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POSGRADO EN GEOCIENCIAS APLICADAS

**Geoquímica, petrogénesis y geocronología del
vulcanismo en el Complejo Volcánico Sierra de
San Miguelito, Campo Volcánico de San Luis
Potosí, México**

Tesis que presenta

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Para obtener el grado de

Doctor(a) en Geociencias Aplicadas

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Constancia de aprobación de la tesis

La tesis "Geoquímica, petrogénesis y geocronología del vulcanismo en el Complejo Volcánico Sierra de San Miguelito, Campo Volcánico de San Luis Potosí, México" presentada para obtener el Grado de Doctor(a) en Geociencias Aplicadas fue elaborada por **Darío Torres Sánchez** y aprobada el **quince de diciembre del dos mil veinte** por los suscritos, designados por el Colegio de Profesores de la División de Geociencias Aplicadas del Instituto Potosino de Investigación Científica y Tecnológica, A.C.

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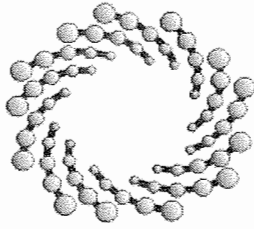
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Dedicatoria

A mi prometida Edith Elizondo

A mis padres Darío Torres González y Sonia Laura Sánchez de la Garza

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Resumen

Geoquímica, petrogénesis y geocronología del vulcanismo en el Complejo Volcánico Sierra de San Miguelito, Campo Volcánico de San Luis Potosí, México

La presente tesis doctoral describe las características geológicas, petrográficas, geoquímicas y geocronológicas de las rocas volcánicas del Complejo Volcánico Sierra de San Miguelito (CVSSM). El CVSSM se constituye principalmente por rocas volcánicas de composiciones máficas, intermedias y félsicas. Las rocas máficas del presentan texturas porfiríticas con matriz vítrea y un ensamblaje mineral principal compuesto por fenocristales de olivino, clinopiroxeno, ortopiroxeno, y plagioclasas, a su vez, estas rocas se caracterizan por contenidos de 38.12-51.48% de SiO₂, 4.59-6.26% MgO y valores de #Mg [#Mg = 100 x (Mg²⁺/Mg²⁺ + Fe²⁺)] que varían entre 41.46 y 55.24. De igual forma, muestran un patrón con enriquecimiento de tierras raras ligeras (REE) y elementos traza menos incompatibles con anomalías negativas de Nb (0.64), Ta, Ba, y P. Las rocas intermedias muestran texturas porfiríticas con matriz vítrea y un ensamblaje mineral principal de fenocristales de clinopiroxeno, ortopiroxeno y plagioclasas. Algo destacable en estas rocas volcánicas, es la interacción entre fenocristales de plagioclasas con texturas de criba junto a plagioclasas con texturas típicas de estos minerales. Muestran contenidos de SiO₂ de 57.46-62.24%, MgO de 2.81-5.71% y valores de #Mg [#Mg = 100 x (Mg²⁺/Mg²⁺ + Fe²⁺)] que varían entre 53.04 y 64.13. Presentan un patrón de tierras raras y elementos traza similares a las rocas máficas, con una presencia similar en anomalías de Nb (0.22-0.33), Ta, Ba, y P. Las rocas félsicas contienen texturas porfiríticas en una matriz vítrea con un ensamblaje mineral principal compuesto por fenocristales de cuarzo, feldespato, plagioclasas y ortopiroxenos. Los contenidos de SiO₂ y Al₂O₃, son de 64.36-82.24%, 10.92-21.26%, respectivamente. Muestran patrones enriquecidos en tierras raras ligeras con una tendencia a la horizontal en tierras raras pesadas, a la vez, es una anomalía negativa pronunciada de Eu, lo cual se asocia a la fraccionación de feldespatos y plagioclasas lo cual coincide con la cantidad de feldespatos y plagioclasas notables

en las rocas félsicas. A su vez, las rocas, muestran anomalías negativas de Nb (0.21-35.38), Ta, Ba y P.

Las características isotópicas de las rocas máficas del CVSSM revelan que se generaron en el manto, lo cual a su vez es demostrable a partir de los valores positivos de $\epsilon_{Nd(t)}$ (+5.2 - +0.9). Por otro lado, tanto las rocas intermedias, como las rocas félsicas muestran una evolución a partir de componentes corticales, lo cual se asocia a valores negativos de $\epsilon_{Nd(t)}$ (-2.5 a -1.8).

A partir de las nuevas dataciones por medio del método de $^{40}Ar/^{39}Ar$ obtenidas para el presente trabajo, junto con la comparación de edades propuestas de trabajos previos, se observó que las rocas del CVSSM se generaron en tres distintos periodos de tiempo. El primer periodo se caracteriza por la formación de lavas félsicas de características efusivas, los cuales dieron lugar a estructuras dómicas. También dentro de este episodio se emplazaron distintos paquetes piroclásticos de distintos espesores, los cuales presentan una composición de igual forma félsica. El segundo periodo se describe como un episodio predominantemente explosivo, en donde, se generaron grandes paquetes piroclásticos de composición félsica junto con la erupción de flujos de lavas aisladas de una composición intermedia. Por último, el tercer periodo se caracteriza por la presencia aislada de flujos de lava masivos de composiciones máficas, principalmente asociados al sistema de fallas. Los modelos cuantitativos generados en el presente trabajo revelan que las rocas máficas evolucionaron por la fusión parcial de una fuente de manto aproximadamente a los 22–21 Ma, mientras que las rocas intermedias se generaron por fraccionamiento de una fuente máfica la cual presentó distintos niveles de asimilación cortical en un periodo de 32–28Ma. Por último, los modelos revelan que las rocas félsicas se generaron a partir de un proceso de fusión parcial de metasedimentos a niveles de corteza media y superior en un periodo de tiempo que abarca de los 34 a 32Ma.

Palabras clave: Geoquímica, Petrogenesis, Rocas volcánicas, Ambiente tectónico, San Luis Potosí, datación $^{40}Ar/^{39}Ar$, Mesa Central, México, isótopos Sr-Nd-Pb.

Abstract

Geochemistry, petrogenesis and geochronological of the volcanism from the Sierra de San Miguelito Volcanic Complex, San Luis Potosí Volcanic Field, Mexico

The present work describes the geological, petrographic, geochemistry and geochronological characteristics from the Sierra de San Miguelito Volcanic Complex (SSMVC) volcanic rocks. The SSMVC mainly of volcanic rocks of mafic, intermediate, and felsic. The mafic rocks display porphyritic textures with vitreous matrix, and a main mineral assemblage of phenocrysts of olivine, clinopyroxene and plagioclases, these rocks are characterized by the contents of 38.12-51.48% SiO₂, 4.59-6.26% MgO and #Mg [#Mg = 100 x (Mg²⁺/Mg²⁺ + Fe²⁺)] values that range 41.46 – 55.24. The mafic rocks show light rare earth elements (LREE) and enrichment incompatible trace elements with small negative anomalies of Nb (0.64), Ta, Ba, and P. In the other hand, the intermediate volcanic rocks from SSMVC consist of phenocrysts of clinopyroxene, orthopyroxene and plagioclases with porphyritic textures with a vitreous matrix, these rocks display contents of SiO₂ of 57.46-62.24%, MgO of 2.81-5.71% and #Mg [#Mg = 100 x (Mg²⁺/Mg²⁺ + Fe²⁺)] values that range 53.04 to 64.13. Intermediate rocks show LREE and trace elements patterns similar to the mafic volcanic rocks negative anomalies of Nb (0.22-0.33), Ta, Ba, and P. The felsic volcanic rocks of the SSMVC are characterized by porphyritic textures with a vitreous matrix, and a main mineral assemblage of quartz, alkali feldspar, plagioclases and orthopyroxene. The contents of SiO₂, Al₂O₃, in these rocks are 64.36-82.24%, 10.92-21.26%, respectively. Felsic rocks display an enriched trend in LREE with negative anomalies of Eu, which are probably associated with the fractionation of feldspar. These rocks display negative anomalies of Nb (0.21–35.38), Ta, Ba, and P.

Isotopic characteristics of mafic volcanic rocks from the SSMVC reveals that these rocks were generated in the mantle, which is demonstrable from their positive values of $\epsilon_{Nd(t)}$ (+5.2 - +0.9). In the other hand, the intermediate rocks and the felsic rocks display involvement of crustal components, as depicted by the negative values of $\epsilon_{Nd(t)}$ (-2.5 to -1.8).

New $^{40}\text{Ar}/^{39}\text{Ar}$ ages presented in this work, and in the compilation of previous ages from the area, three main magmatic episodes were defined as follows: (a) the first episode is mainly characterized by the eruption of felsic lavas of effusive characteristics, which formed domes structures and different pyroclastic deposits of felsic composition that were erupted; (b) the second episode is characterized by explosive eruption of pyroclastic deposits of a felsic composition, as well as, lava flows of intermediate composition in isolated area; (c) the third episode is defined by isolated lava flows of mafic compositions, that are located mainly in the edge of fault systems.

Magmatic models generated during the present work reveals that mafic rocks evolved through a partial melting process of the mantle during 22–21 Ma. The intermediate rocks were generated by the fractionation of mafic sources, which presented different levels of crustal assimilation during 32–28 Ma. Finally, the magmatic models display that felsic rocks were generated by partial melting of metasediments at middle-upper continental crust levels during the period of 34 to 32 Ma.

Keywords: Geochemistry, Petrogenesis, Volcanic rock, Tectonic setting, San Luis Potosí, $^{40}\text{Ar}/^{39}\text{Ar}$ dating, Mesa Central, Mexico, Sr-Nd-Pb isotopes

INTRODUCCIÓN

La provincia del *Basin and Range* (BR), o también conocida como la provincia de Cuencas y Sierras, ha sido de gran importancia en la comunidad geológica ya que sirve como un claro ejemplo de un sistema montañoso en bloques de escala regional, el cual se ha sido formado por deformación extensional. La BR se extiende desde Canadá pasando por el oeste de los EUA y terminando hacia el noroeste de México (Figura 1a), a su vez, esta provincia se ha descrito como una región montañosa Cenozoica con una configuración actual y una geometría interna derivada de las múltiples fases de deformación extensional conducidos por distintos impulsos geodinámicos ([Dickinson, 2002](#)).

La BR es una larga sucesión de estructuras de graben y horst del Cenozoico con una configuración actual y geometría interna derivada de múltiples fases de extensión ([Dickinson, 2002](#)). A su vez, las estructuras tanto estructurales como las rocas volcánicas existentes a través de esta provincia, data desde aproximadamente del Oligoceno temprano hasta el Cuaternario ([Gans, 1981](#); [Eaton, 1982](#); [Dickinson, 2002](#)).

Se ha propuesto que el inicio de las fases extensivas que dieron lugar a la construcción del BR, y la disminución de la elevación de la superficie se atribuye a la reorganización del límite de placas Pacífico-Norteamérica durante el Cenozoico, y más específicamente al establecimiento del sistema transformante de San Andreas ([Cassel et al., 2014](#); [Long, 2019](#)). La disminución en el acoplamiento intraplaca que acompañó a la desaparición de la subducción de la placa Farallón, junto con la correspondiente influencia creciente del cizallamiento dextral en el margen de placa, sigue siendo la explicación más aceptada para el control extensional principal del BR ([Dickinson, 2002](#); [Long, 2019](#)). La provincia del Basin and Range se ha separado en tres secciones principales como: (i) sección norte; (ii) sección central; (iii) sección sur (Figura 1a; [Cosca et al., 2014](#)).

La sección sur del BR se caracteriza principalmente por presentar actividades extensionales syn-transformantes las cuales se han relacionado a la migración rápida de la triple unión de la placa Rivera la cual pasa por Baja California, México hasta su ubicación actual entre la península de Baja California y el terreno mexicano, como también esta sección presenta una gran cantidad de actividad volcánica a través de la provincia, los cuales tuvieron lugar desde el Paleoceno Tardío, hasta el Mioceno Tardío ([House et al., 2001](#); [Spencer et al., 2001](#); [Jones et al., 1992](#); [Dickinson, 2002](#); [Cosca et al., 2014](#)).

Dentro de la sección sur de la BR se encuentra la provincia de la Mesa Central (MC) la cual se localiza en la región centro-norte de México, cubriendo los estados de Chihuahua, Durango, Zacatecas, Aguascalientes, y San Luis Potosí y se ha definido como una meseta elevada la cual provee un registro geológico que abarca desde el Triásico hasta el presente ([Nieto-Samaniego et al., 2007, 2019](#)). La MC se encuentra limitada hacia el N y E por la Sierra Madre Oriental, y para el W por la Sierra Madre Occidental (Figura 1b).

La MC se ha dividido en dos regiones principales: (i) región norte, la cual se caracteriza principalmente por episodios avanzados de erosión con cuencas aluviales-lacustres, y unidades volcánicas dispersas del Oligoceno al Cuaternario; (ii) región sur, se distingue por presentar un amplio volumen de rocas volcánicas del Paleogeno-Neogeno, cortadas por diversos sistemas de fallas normales ([Nieto-Samaniego et al., 1996, 1999, 2007](#)).

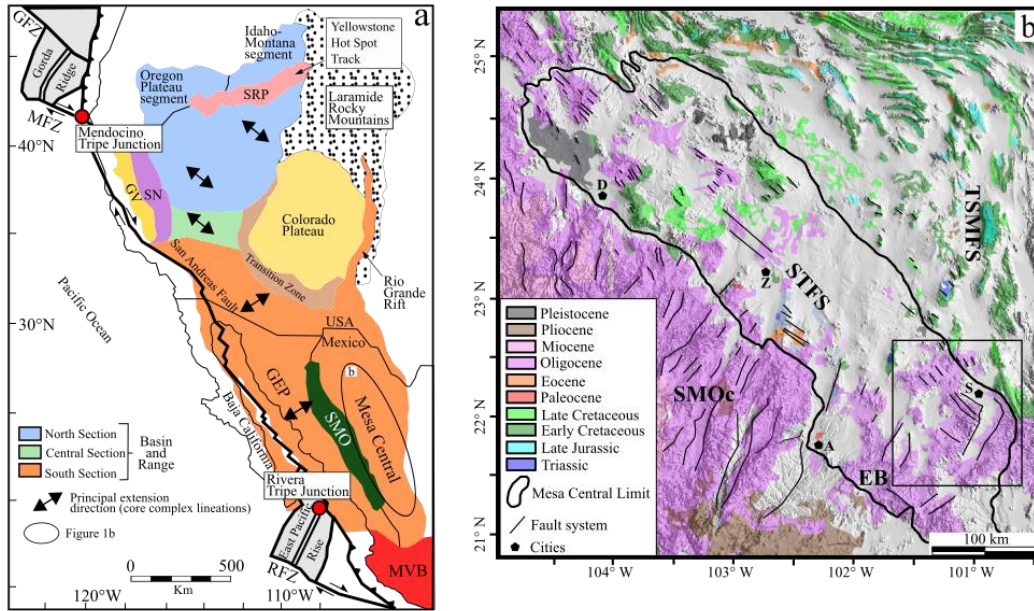


Figura 1. a) Mapa tectónico regional de la situación actual de la sección oeste de Norteamérica y la provincia del Basin and Range con las subdivisiones de las regiones norte, centro y sur del Basin and Range (Modificado de Cosca et al., 2014); b) Mapa geológico simplificado de la Mesa Central. El recuadro negro representa el área de estudio del presente trabajo. (Modificado de Nieto-Samaniego et al., 2007). Abreviaciones: SMOc= Sierra Madre Occidental; STFS= Sistema de Falla San Luis-Tephuanes; TSMFS= Sistema de Fallas Taxco-San Miguel de Allende; EB=Graben El Bajío; S=San Luis Potosí; Z=Zacatecas; A= Aguascalientes; D=Durango

Las estructuras Cenozoicas mayores que conforman a la Mesa Central son:(a) *Falla del Bajío*: la cual presenta dos grupos de fallas con direcciones NW-SE y ENE a NE, ambos grupos proporcionan una forma aserrada al borde norte de El Bajío, en esta zona los desplazamientos mínimos son entre 150 y 250m, en cambio, el segundo segmento presenta una longitud de 80 km aproximadamente y se conforma por una serie de fallas normales de ángulo alto, escalonadas con bloques hundidos al SW;(b) *Sistema de fallas Taxco-San Miguel de Allende*: se constituye por sistemas de falla de dirección N-S. Los principales lineamientos NW-SE que se observan segmentando a este sistema de fallas son los sistemas de fallas San Luis-Tephuanes y la falla el Bajío; (c) *Sistema de fallas San Luis-Tephuanes*: de forma general este sistema se caracteriza por una serie de lineamientos de dirección NW-SE en donde su traza coincide burdamente con el límite de los alforamientos de rocas volcánicas cenozoicas pertenecientes a la Sierra Madre Occidental; (d) *Graben de Aguascalientes*: consiste de un graben asimétrico cuya falla principal se ubica hacia el occidente (Nieto-Samaniego et al., 2007).

Dentro de la región sur de la MC se localiza el Campo Volcánico de San Luis Potosí (CVSLP) el cual incluye secuencias volcánicas de composiciones básicas, intermedias y ácidas las cuales hicieron erupción a partir desde el Eoceno hasta el Cuaternario. Los eventos volcánicos del CVSLP tuvieron lugar durante el Oligoceno Tardío y se caracteriza principalmente por riolitas de alto sílice y largos volúmenes de ignimbritas. A su vez, los eventos volcánicos jóvenes en el CVSLP se caracterizan por bajos volúmenes de lavas de composición básica e intermedia (Tristán-González et al., 2009; Aguillón-Robles et al., 2014; Torres-Sánchez et al., 2019). El CVSLP ha sido dividido en seis complejos principales: (i) Complejo Ahualulco; (ii) Complejo Villa Hidalgo; (iii) Complejo La Repartición; (iv) Complejo Pinos; (v) Complejo Santa María; y el (vi) Complejo de la Sierra de San Miguelito (Torres-Hernández et al., 2006; Rodríguez-Ríos et al., 2007; Tristán-González et al., 2009; Aguillón-Robles et al., 2014; López-Loera, 2014; Torres-Sánchez et al., 2019).

Diversos estudios han sido desarrollados dentro del Complejo Volcánico de la Sierra de San Miguelito (p.ej. Tristán-González et al., 2009; Aguillón-Robles et al., 2014; López-Loera, 2014), sin embargo, aún existen distintas ausencias de información, como de modelos complementarios. Dentro de este apartado se muestra una sobrevista sobre las metas y objetivos que se proponen en el presente estudio.

1.1 Justificación y objetivos de investigación

1.1.1 Justificación

Diversa información ha sido generada para el Complejo Volcánico de la Sierra de San Miguelito (CVSSM), desde cartografía geológica, geología estructural, petrografía, geoquímica de roca total y datación por el método de K/Ar en roca total. Sin embargo, para una propuesta de un modelo petrogenético se requiere: a) contar con un muestreo representativo, tanto de litología como espacialmente del complejo volcánico; b) establecer una composición modal de las diferentes rocas, como sus características texturales y la composición química tanto de la matriz de la roca como de sus minerales; c) contar con una base de datos completa de geoquímica (elementos mayores y traza), relaciones isotópicas (Sr-Nd-Pb) para cada muestra; d) generar información geocronológica para rocas de las distintas litologías de la zona y elaborar un contraste con las edades propuestas; f) deducir el ambiente tectónico asociado al vulcanismo de la región a través de diagramas de discriminación tectónica convencionales, y de diagramas basados en estadística multivariada.

1.1.2 Objetivos

El objetivo principal de este estudio es el proponer un modelo petrogenético cuantitativo para las rocas volcánicas del Complejo Volcánico Sierra de San Miguelito.

Las metas que se alcanzaron para este trabajo son:

- Descripción de la distribución espacial de las diferentes litologías y su relación con los elementos estructurales de la región.
- Caracterización de los distintos tipos de rocas desde una perspectiva petrográfica, geoquímica e isotópica.
- Evaluación cuantitativa de procesos magmáticos que generaron al vulcanismo en el CVSSM.
- Elaboración de un esquema geocronológico de los eventos magmáticos que forman parte del CVSSM.
- Propuesta de un ambiente tectónico para el vulcanismo del CVSSM.

1.2 Hipótesis

Las hipótesis de investigación del presente proyecto son las siguientes:

1. El vulcanismo máfico del Complejo Volcánico de la Sierra de San Miguelito es relacionado a un proceso de fusión parcial del manto.
2. El origen del vulcanismo intermedio del Complejo Volcánico de la Sierra de San Miguelito es relacionado a un proceso de asimilación–cristalización fraccionada con una mezcla de magmas entre rocas félsicas y máficas.
3. El vulcanismo félsico del Complejo Volcánico de la Sierra de San Miguelito se relaciona a un proceso de fusión parcial de distintos niveles de la corteza continental.
4. La actividad magmática tuvo lugar durante el Eoceno Tardío–Mioceno Temprano durante un evento tectónico extensional.

1.2.1 Prueba de hipótesis

Para poder comprobar las hipótesis planteadas anteriormente se ha propuesto las siguientes pruebas:

1. Análisis geoquímico de elementos mayores, traza y tierras raras ayudarán para el entendimiento del ambiente tectónico del vulcanismo.
2. Las nuevas edades Ar/Ar, aportarán al análisis geocronológico para la precisión de las edades de formación.
3. Análisis isotópico se lleva a cabo para el entendimiento de la evolución geodinámica y petrogenesis del Complejo Volcánico de la Sierra de San Miguelito.

1.3 Esquema de tesis

El presente trabajo se basa en tres artículos publicados, uno en participación como co-autor y dos publicados como primer autor.

El capítulo **uno** del presente estudio describe las características geológicas de las provincias geológicas de la Mesa Central, así como, del Campo Volcánico de San Luis Potosí. El capítulo **dos** muestra la descripción geológica y el muestreo del área de estudio. El capítulo **tres** presenta los resultados del análisis petrográfico de las rocas

estudiadas en el presente trabajo. El **cuarto** capítulo presenta un artículo publicado en “*Journal of Spectrometry*” donde el autor de este trabajo participó como co-autor. En este artículo se describe la importancia de la aplicación de modelos de regresión lineal ponderada para la calibración de máquinas de fluorescencia de rayos-X a partir de 59 materiales de referencia.

El capítulo **cinco** presenta el primer artículo publicado en la revista “*Journal of South American Earth Sciences*”, el cual se enfoca en las características petrográficas y geoquímicas de las rocas máficas y félsicas del Complejo Volcánico de la Sierra de San Miguelito, y se presentan los primeros modelos petrogenéticos que ayuden a explicar un posible origen de las rocas máficas y félsicas del CVSSM. El capítulo **seis** presenta el segundo artículo publicado en la revista “*Lithos*”, el cual se enfoca tanto en las conexiones petrogenéticas, como en el contraste de edades entre las rocas básicas, intermedias y ácidas del CVSSM y sus implicaciones en la actividad volcánica Cenozoica de la Mesa Central. El capítulo **siete** presenta las características de química mineral de las distintas unidades del CVSSM, y se propone las temperaturas en las cuales se generaron las distintas rocas volcánicas del CVSSM. El último capítulo (capítulo **ocho**) provee un resumen de los hallazgos principales y sugiere como la investigación puede ser complementada y/o extendida en el área de estudio del presente trabajo.

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CAPITULO 1.-

GEOLOGÍA REGIONAL—REGIÓN SUR DE LA MESA CENTRAL

En el presente capítulo se presenta una descripción de rasgos geológicos destacables de la región sur de la Mesa Central, y se describen las características geológicas del Campo Volcánico de San Luis Potosí. Las siguientes descripciones aportan nueva información acerca de la estratigrafía, geología estructural, así como de la actividad volcánica de la zona de estudio.

1.1 Estratigrafía de la región sur de la Mesa Central

La región sur de la Mesa Central presenta un registro geológico amplio el cual abarca un período de tiempo desde el Mesozoico hasta la época presente (Figura 1.1). Las rocas Mesozoicas en esta región son representadas por dos principales litologías: (i) rocas marinas calcáreas de la secuencia de la Sierra Madre Oriental; (ii) y secuencias del complejo volcano-sedimentario de la Sierra de Guanajuato ([Eguiluz de Antaño et al., 2000](#); [Orozco-Esquivel et al., 2002](#)).

Por otro lado, la evidencia magmática para el Cretácico Tardío–Paleoceno (Figura 1.1) es escasa, principalmente representada por cuerpos intrusivos que se acomodan a lo largo de las principales estructuras tectónicas de la MC, y por cuerpos intrusivos graníticos en la Sierra de Guanajuato para el Paleoceno. Para el Eoceno, el vulcanismo corresponde a cuerpos aislados de lavas andesíticas, depósitos piroclásticos y cuerpos riolíticos reportados en el área de San Luis Potosí y Guanajuato ([Martínez-Reyes, 1992](#); [Orozco-Esquivel et al., 2002](#); [Nieto-Samaniego et al., 2019](#)).

En cambio, para el Oligoceno se ha reportado un evento volcánico voluminoso el cual formó una cubierta volcánica gruesa de lavas de composición andesítica a riolítica, cuales reposan sobre unidades Mesozoicas (Figura 1.1; [Orozco-Esquivel et al., 2002](#)). [Orozco-Esquivel et al \(2002\)](#) agruparon a esta cobertura volcánica del Oligoceno en dos secuencias: (a) *secuencia inferior*, la cual incluye rocas emplazadas antes de la formación del Graben de Villa de Reyes, las cuales presentan una variación composicional de andesitas a riolitas, y en porcentaje bajo, distintos depósitos piroclásticos de composición

riolítica; y, (b) *secuencia superior*, en donde la mayoría de las rocas fueron formadas durante una fase volcánica muy bien definida, la cual inició con el emplazamiento de grandes volúmenes de lavas riolíticas, seguido por un evento ignimbrítico voluminoso de composición igualmente riolítica. Una segunda fase volcánica tomo lugar, la cual produjo de igual forma lavas riolíticas y paquetes ignimbríticos, solo que estos a menor escala que la fase anterior.

Para el Mioceno (Figura 2.1), se ha reportado rocas volcánicas de composición basáltica y andesítica intercaladas con los sedimentos que rellenan cuencas continentales cenozoicas, y afloramientos de rocas volcánicas de la misma composición en numerosas localidades aisladas (McDowell y Keizer, 1977; Luhr et al., 1995; Henry y Aranda-Gómez, 2000; Rodríguez-Ríos et al., 2007; Aguillón-Robles et al., 2009).

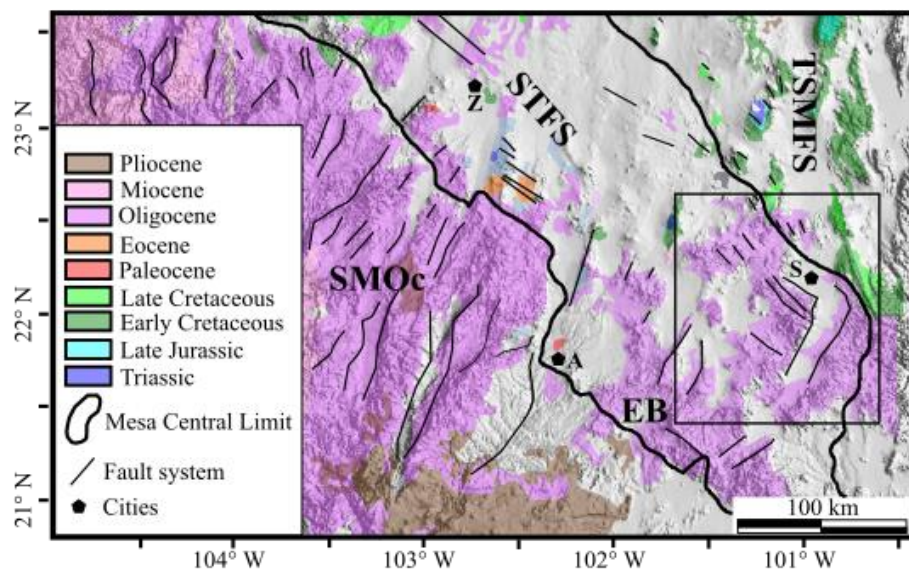


Figura 1.1 Mapa geológico de la sección sur de la Mesa Central (Modificado de Nieto-Samaniego et al., 2007). Abreviaciones: SMOc= Sierra Madre Occidental; STFS=Sistema de Fallas San Luis Tepehuanes; TSMFS=Sistema de Fallas Taxco-San Miguel de Allende; EB= Graben El Bajío; S=San Luis Potosí; Z= Zacatecas; A= Aguascalientes.

Por último, para el Plioceno y Cuaternario (Figura 1.1) se han reconocido rocas volcánicas que afloran principalmente en pequeños conjuntos de aparatos volcánicos como conos cineríticos, flujos de lava y “maares”. Estos cuerpos volcánicos abarcan una composición basáltica alcalina, y corresponden a lavas alcalinas con xenolitos de lertzolita provenientes del manto y de granulita procedentes de la base de la corteza (Labarthe-Hernández et al., 1982; Swanson, 1989; Pier et al., 1989; Schaaf et al., 1994; Aranda-Gómez et al., 2003; Saucedo et al., 2017)

1.2 Geología estructural de la región sur de la Mesa Central

La región sur de la Mesa Central se encuentra afectada por distintos sistemas de fallas, entre los cuales se encuentran el sistema de fallas San Luis–Tepehuanes (SFST), la falla El Bajío (FB) y el graben de Aguascalientes (GA; Nieto-Samaniego et al., 2007).

El SFST se caracteriza por un conjunto de lineamientos de dirección NW-SE (Figura 1.1). La porción oriental del SFST se conforma por diversas estructuras de fallas normales con dirección de rumbo NW-SE y buzamiento SW, que conforman diversos sistemas de horst, grabens, y semi-grabens, los cuales presentaron actividad durante el Oligoceno hasta el Mioceno (Labarthe-Hernández y Jiménez-López, 1992; Silva-Romo, 1996; Xu et al., 2004; Nieto-Samaniego et al., 2007). Por otro lado, la porción occidental del SFST se conforma por estructuras de fallas normales con direcciones NW-SE, y en menor número, fallas con rumbos E-W y NE-SW, la actividad principal de estos sistemas de fallas ocurrieron entre el Eoceno y el Oligoceno (Albinson, 1988; Arada-Gómez et al., 2003; Loza-Aguirre, 2005; Nieto-Samaniego et al., 2007).

La FB se caracteriza por mostrar dos segmentos mayores que se extienden desde Celaya hasta Guanajuato (Figura 1.1). El primer segmento de esta estructura se caracteriza por dos grupos de fallas con direcciones NW-SE y ENE, los cuales brindan a este sector una forma aserrada al borde norte de El Bajío (Trujillo-Candelaria, 1985; Nieto-Samaniego et al., 2005). En cambio, el segundo segmento muestra una extensión ~ de 80km de largo y es representada por una serie de fallas normales de ángulo alto, escalonadas y con bloque hundidos al SW (Quintero-Legorreta, 1992; Nieto-Samaniego et al., 2005).

El GA limita al occidente la región sur de la Mesa Central separándola de la Sierra Madre Occidental. Esta estructura consiste de un graben asimétrico cuya falla principal se ubica al occidente y se extiende aproximadamente por 150 km con dirección N-S (Figura 1.2; [Jiménez-Nava, 1993](#); [Nieto-Samaniego et al., 2007](#)).

1.3 Campo Volcánico San Luis Potosí (CVSLP)

1.3a Estratigrafía del CVSLP

El Campo Volcánico de San Luis Potosí (CVSLP, Figura 1.2) se localiza en la porción sureste de la región sur de la Mesa Central, se definió por primera vez por Labarthe-Hernández et al. (1982) como un campo volcánico conformado por lavas e ignimbritas del Terciario. Actualmente se conoce al CVSLP como un campo volcánico que cubre un área cercana a los 10, 000 km², y se encuentra formado principalmente por secuencias de lavas y flujos piroclásticos que abarcan una edad desde el Eoceno medio hasta el Cuaternario, y un rango composicional que va desde rocas básicas hasta rocas ácidas ([Tristán-González et al., 2009](#)).

Se ha establecido que el CVSLP presentó cinco etapas principales de vulcanismo, donde la primera predominó la emisión de lavas andesíticas durante el Eoceno Medio; la segunda etapa abarcó el Oligoceno Temprano y se compone principalmente por lavas de composición ácida (dacitas y riolitas). La tercera y cuarta etapa se conforman por un vulcanismo félsico y máfico el cual ocurrió durante el Oligoceno Tardío-Mioceno Inferior. La quinta y última etapa se caracteriza por un vulcanismo intraplaca de composición basanítica, la cual abarca principalmente el Cuaternario ([Tristán-González et al., 2009](#); [Aguillón-Robles et al., 2012](#)).

En base a la semejanza estratigráfica, geoquímica y geocronológica [Tristán-González et al. \(2009\)](#) dividieron al CVSLP en diversos complejos volcánicos (Figura 2.3), los cuales se han denominado como:

- a) **Ahualulco**: se caracteriza por presentar diversas estructuras volcánicas de composición andesítica, domos riolíticos del Eoceno Medio-Oligoceno Temprano, y flujos piroclásticos asociados a estos. Dentro de este complejo

volcánico también se pueden apreciar derrames de lavas basaníticas del Cuaternario.

- b) **Pinos:** el cual se conforma por derrames de lavas y domos poligenéticos del Oligoceno temprano de un rango composicional principalmente dacítica y riolítica, y derrames piroclásticos asociados a estos.
- c) **Villa Hidalgo:** se caracteriza por mostrar domos y derrames de lava de composición dacítica-andesítica y flujos piroclásticos asociados a estos emplazados durante el Oligoceno. A su vez, dentro de este campo se observan lavas máficas del Cuaternario, las cuales contienen xenolitos tanto de manto como de corteza.
- d) **La Repartición:** se comprende por depósitos de flujos piroclásticos, un conjunto de lavas dacíticas, andesíticas y basálticas. También se presentan diques tufocíticos y acumulaciones de material piroclástico y epiclástico.
- e) **Sierra de San Miguelito:** está formado principalmente por domos de composición riolítica intercalados con un paquete grueso de depósitos piroclásticos del Oligoceno; también, se presenta vulcanismo puntual de composición básica. Algo característico en este complejo es que las secuencias se encuentran afectadas por una gran cantidad de fallas normales de ángulo alto, las cuales se acomodaron en patrón de dominó, formando así semi-fosas tectónicas estrechas.
- f) **Santa María:** se caracteriza por presentar cadenas de domos de composición traquítica hasta riolítica, junto con depósitos piroclásticos asociados a estos. El gran paquete volcánico de este complejo fue afectado por una serie de fallas normales NW-SE. Durante el último evento efusivo de este complejo se inyectaron puntualmente coladas de lavas basálticas a lo largo de fallas normales, principalmente sobre la zona de fallas marginales al SE del Graben de Villa de Reyes.

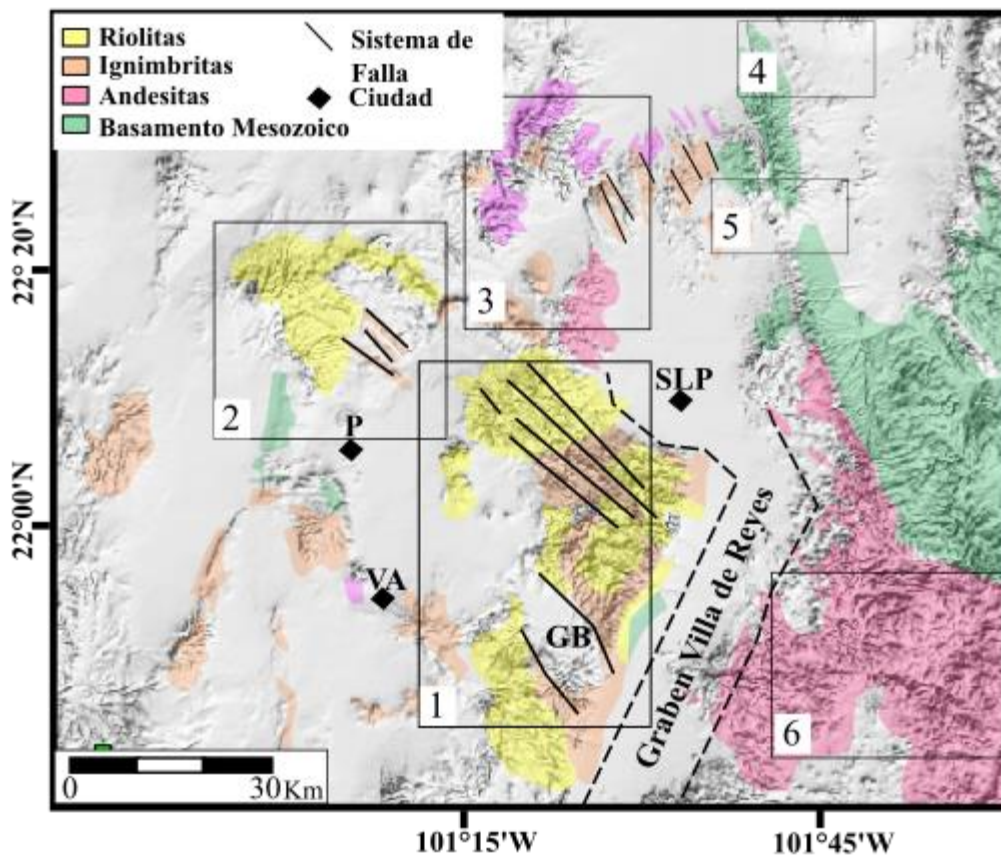


Figura 1.2 Mapa geológico del CVSLP (Tomado y modificado de Tristán-González et al., 2009). Abreviaciones: SLP=San Luis Potosí, VA=Villa de Arista; P=Pinos; GB=Graben Bledos. 1.-Complejo Volcánico Sierra de San Miguelito; 2.- Complejo Volcánico Pinos; 3.- Complejo Volcánico Ahualulco; 4.- Complejo Volcánico la Repartición; 5.- Complejo Volcánico Villa Hidalgo; 6.- Complejo Volcánico Santa María

1.3b Geología estructural del CVSLP

Con respecto a la geología estructural del CVSLP se han caracterizado dos regiones estructurales distintas: (i) el Graben de Villa de Reyes (GVR) localizado entre el Complejo Volcánico de la Sierra de San Miguelito y el Complejo Volcánico Santa María; y (ii) el Graben de Bledos (GB) el cual se localiza hacia el sur del Complejo Volcánico de la Sierra de San Miguelito (Figura 1.2).

El GVR se ha definido como una fosa tectónica de edad Oligocénica, la cual se encuentra rellena por depósitos volcánicos y clásticos de edades que abarcan desde el Oligoceno temprano hasta la época actual (Figura 1.2). Esta estructura presenta una longitud de ~ 200 km y un ancho variable de 10-20 km, y este se limita por un sistema de fallas de

dirección NE. El fondo de la fosa se conforma por domos exógenos del Oligoceno y en algunos sitios llegan a aflorar como pequeñas cumbres, como se observa en las inmediaciones del poblado de Villa de Reyes (Labarthe-Hernández et al., 1982; Tristán-González, 1986; López-Loera et al., 2013). A su vez, el GVR forma parte de una serie de estructuras tectónicas presentes en la región sur de la Mesa Central las cuales corresponden a la etapa de máxima extensión, y esta estructura se encuentra cruzado por un sistema de fallas de orientación NW-SW las cuales se arreglan en un patrón en dominó (Labarthe-Hernández et al., 1982; Tristán-González, 1986; Nieto-Samaniego et al., 1997; Labarthe-Hernández y De la Huerta-Cobos, 1998; Aranda-Gómez et al., 2000; Torres-Aguilera, 2005; Rodríguez-Ríos, 2009; López-Loera et al., 2013).

Por otra parte, el GB consiste en un valle de ~ 18 km de largo y ~8 km de ancho. Esta estructura se ha considerado como un semi-fosa con dos fallas notables en su margen NE, con rumbos NW y buzamientos al SW. Por otra parte, la edad de las fallas se ha definido entre los 29 Ma y 27 Ma (Labarthe-Hernández y Jiménez-López, 1992). A su vez, esta semi-fosa se encuentra rellena por depósitos volcánicos de composiciones riolíticas y basálticas como de depósitos conglomeráticos y aluviales (Labarthe-Hernández y De La Huera-Cobos, 1998).

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CAPITULO 2.- TRABAJO DE CAMPO Y MUESTREO

2.1 Localización del área de estudio

El Complejo Volcánico de la Sierra de San Miguelito (CVSSM) se localiza en la sección central de México, hacia el borde occidental del estado de San Luis Potosí, México (Figura 2.1), donde forma una extensa sierra en los bordes sur, oeste y noroeste de la ciudad de San Luis Potosí. Desde una perspectiva fisiográfica, el CVSSM forma parte de la provincia de la Mesa Central.

Las vías de acceso hacia el CVSSM son principalmente la carretera estatal No. 80 San Luis Potosí–Guadalajara, con la desviación a la comunidad de Escalerillas y Presa del Peaje, como también el anillo periférico libramiento sur y este con desviación hacia la Cañada del Lobo, Capulines, Iglesia del Desierto, Presa de San José, Villa Magna y Horizontes (Figura 2.1). De igual forma se puede acceder hacia el CVSSM por la Carretera Estatal No. 49 con desviación hacia el poblado Suspiro Picacho y también algunos caminos de terracería que llegan a comunicar pequeñas comunidades con poblados.

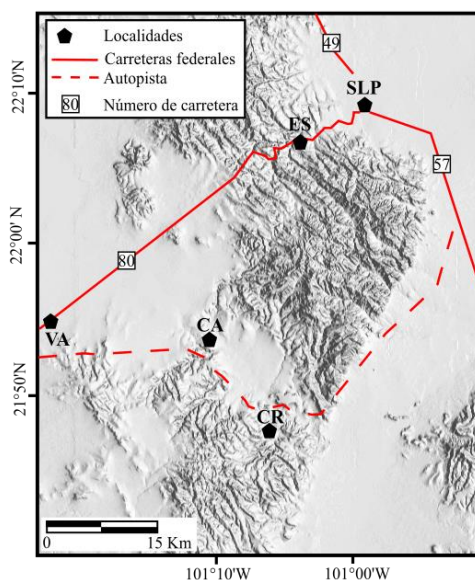


Figura 2.1.- Mapa de ubicación del área de estudio. Abreviaciones: CA= Cabras; CR= Carranco; ES= Escalerillas; SLP= San Luis Potosí; VA= Villa Aldama.

2.2 Geología estructural del CVSSM

El CVSSM se considera como un área relativamente alta de la región sur de la provincia de la Mesa Central. Los grandes ejes y el alineamiento de domos volcánicos comúnmente indican las orientaciones de largas fallas normales (Xu et al., 2008).

Las estructura geológica principal que limita al CVSSM es el Graben de Villa de Reyes (GVR; Figura 2.2), el cual se caracteriza por ser una fosa tectónica del Oligoceno rellena por depósitos volcánicos y clásticos de edad del Oligoceno temprano hasta depósitos recientes (López-Loera et al., 2013; Xu et al., 2013). La longitud aproximada del GVR es de 200 km con un ancho variable de 10-20 km, a su vez, el GVR forma parte a las distintas series de estructuras tectónicas de la provincia de la Mesa Central las cuales corresponde a la etapa de máxima extensión del área (Labarthe-Hernández et al., 1982; Tristán-González, 1986; Nieto-Samaniego et al., 1996; Henry y Aranda-Gómez, 1992; López-Loera et al., 2013).

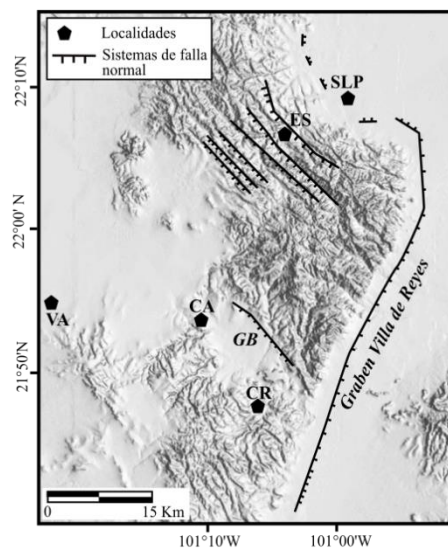


Figura 2.2.- Mapa de los sistemas de fallas principales del CVSSM. Abreviaciones: CA= Cabras; CR= Carranco; ES= Escalerillas; SLP= San Luis Potosí; VA= Villa Aldama; GB= Graben de Bledos.

El CVSSM se compone por numerosas fallas normales con rumbos que oscilan entre los 300° y 340°. La mayoría de estas fallas presentan direcciones de buzamiento hacia el SW y varían desde los 45° a los 75° de buzamiento (Xu et al., 2004). Se han clasificado a estas fallas como un sistema *domino* debido a que muestran dirección de buzamiento de falla uniforme, dirección de buzamiento de bloques uniformes y ángulos de buzamientos similares a los bloques (Figura 2.2; Labarthe-Hernández y Jiménez-López, 1992; Xu et al., 2004, 2008, 2013).

2.3 Estratigrafía local del CVSSM

La estratigrafía del Complejo Volcánico de la Sierra de San Miguelito cubre desde el Oligoceno hasta el Mioceno temprano. El CVSSM muestra siete unidades litológicas principales (Figura 2.3) las cuales son la unidad San Miguelito, San José, El Zapote, Cantera, La Placa, Panalillo, y Cabras.


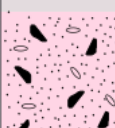


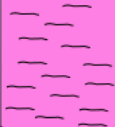

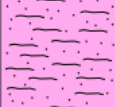
Mioceno	Unidad Cabras		21.5 ± 0.5 a 22.1 ± 0.5 Ma ⁵⁻⁶	Flujos de lava masivos porfídicos, de coloración oscuro-claro y de composición máfica, Contacto discontinuo con la unidad Panalillo
	Unidad Panalillo		25.4 ± 0.6 a 31.9 ± 0.7 Ma ^{1,5}	Ignimbrita de bajo grado de soldado, con pomex y una alta cantidad de fragmentos líticos. Contacto discontinuo con la Unidad Placa.
Oligoceno Tardío	Unidad Placa		26.9 ± 0.4 a 29.0 ± 0.7 Ma ⁵⁻⁶	Flujo de lava masiva, de color oscuro, porfírica de composición máfica. Contacto discontinuo con la Unidad Cantera.
	Unidad Cantera		29.0 ± 1.5 Ma ⁵	Depósito de ignimbrita con un grado de soldado alto, rico en cristales, pobres en líticos con estructuras de fiames y de composición riolítica Contacto discontinuo con la Unidad Zapote.
Oligoceno Temprano	Unidad Zapote		27.4 ± 0.4 a 31.2 ± 0.7 Ma ⁴⁻⁵	Flujo de lava masivo, pobre en cristales de color rosado y de composición riolítica. Contacto discontinuo con la Unidad San José
	Unidad San José		29.0 ± 1.5 Ma ⁵	Depósito de ignimbrita con un grado de soldado medio y de composición riolítica Contacto discontinuo con la Unidad San Miguelito.
	Unidad San Miguelito		21.1 ± 0.3 a 32.7 ± 1.0 Ma ¹⁻⁶	Flujos de lava masivos, ricos en cristales, de color rosado bajo de composición riolítica.

Figura 2.3.- Columna litológica simplificada del Complejo Volcánico de la Sierra de San Miguelito. Edades son tomadas de los siguientes autores: ¹Labarthe-Hernández et al. (1982); ²Tuta et al. (1988); ³Aguillón-Robles et al. (1994); ⁴Aguillón-Robles et al. (2009); ⁵Tristán-González et al. (2009); ⁶Aguillón-Robles et al. (2014).

La unidad más antigua que se puede encontrar en el CVSSM es la unidad San Miguelito (Figura 2.4a) la cual se caracteriza por ser el principal componente de los domos exógenos que se observan dentro del complejo; además, el grosor de la unidad San Miguelito varía entre los 130 m hasta llegar a espesores amplios de 800m, y se observa que es aún más alta en el centro de los domos (Figura 2.4; [Labarthe-Hernández y Jiménez-López, 1992](#)). La mineralogía que se puede observar en rocas de mano consiste principalmente en fenocristales de cuarzo y sanidino embebidos en una matriz desvitrificada.

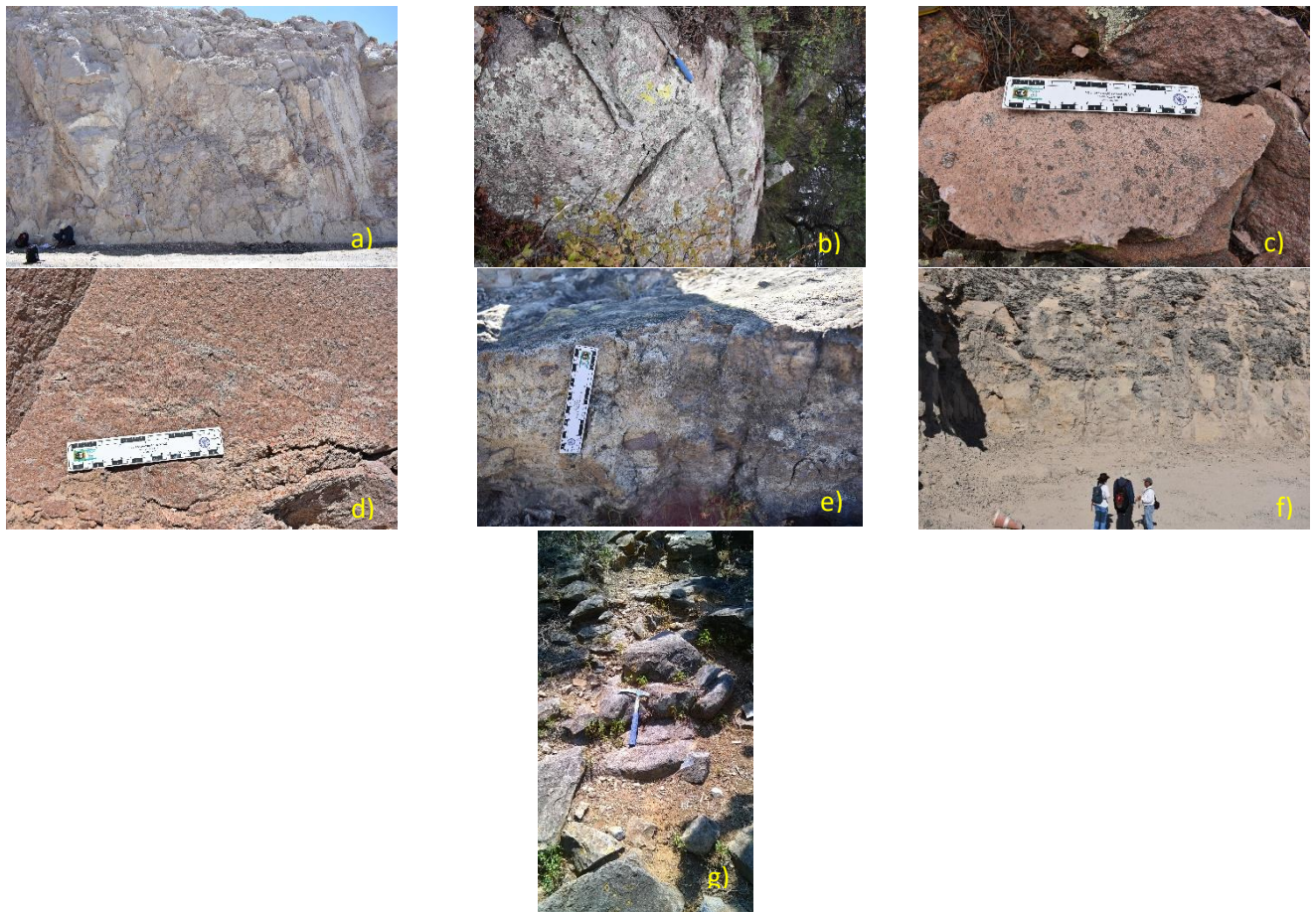


Figura 2.4.- Fotografías de campo representativas de cada unidad litológica del Complejo Volcánico Sierra de San Miguelito. a) Afloramiento masivo de la unidad San Miguelito; b) Afloramiento de bloque masivo de la unidad El Zapote; c) Muestra de mano de la unidad San José, donde se aprecian ; d) Afloramiento de bloque de la unidad Cantera, donde se observan estructuras de *flames*; e) Afloramiento de bloque de roca de la unidad Panalillo donde se observan alta cantidad de líticos; f) Afloramiento masivo basáltico de la unidad La Placa; e) Afloramiento de bloques basálticos de la unidad Cabras.

Estudios previos han reportado edades de roca total por el método de K/Ar de roca total para la unidad San Miguelito (Tabla 2.1; [Labarthe-Hernández y Jiménez-López, 1992](#); [Aguillón-Robles et al., 1994, 2014](#); [Tristán-González et al., 2009](#)). La datación por este método ha variado entre los 33 y 30 Ma para la generación de esta unidad. En algunas localidades esta unidad se observa en contacto junto a una unidad riolítica denominada como la unidad El Zapote, como a su vez, con distintas unidades piroclásticas como lo son la unidad Cantera, Panalillo y San José.

La unidad El Zapote, presenta ciertas características similares a la unidad San Miguelito (lavas masivas de composición riolítica con mineralogía de cuarzo y feldespatos), sin embargo, esta unidad presenta distintos espesores; además se concentra en una sola región del Complejo Volcánico de la Sierra de San Miguelito. A su vez, la unidad El Zapote descansa directamente en el tope de la unidad San Miguelito, y a diferencia de esta, la unidad El Zapote presenta texturas de grano fino con tendencia fluida, y pobre en fenocristales. (Figura 2.4b; [Labarthe-Hernández y Jiménez-López, 1992](#)). Ha sido datada de igual forma por el método de K/Ar por trabajos previos en una edad de 31 a 27 Ma (Tabla 2.1; [Labarthe-Hernández y Jiménez-López, 1992](#); [Aguillón-Robles et al., 2014](#); [Tristán-González et al., 2009](#)).

Entre la unidad San Miguelito y la unidad El Zapote se encuentra una unidad piroclástica la cual se le ha denominado como San José (Figura 2.4c). Esta unidad se caracteriza por localizarse en un sector del CVSSM, como también por espesores de 3 a 15m, consiste principalmente de depósitos de ceniza que se encuentran parcialmente consolidados ([Labarthe-Hernández y Jiménez-López, 1992](#); [Tristán-González et al., 2009](#); [Gaytán-Martínez et al., 2017](#)).

La unidad Cantera subyace a la unidad San Miguelito, El Zapote y San José, como a su vez, cubre la mayoría del CVSSM. La unidad Cantera se determina por depósitos piroclásticos altamente compactado caracterizados por pómez y ceniza de colores grisáceas a rosas, abundantes estructuras de fiames y fragmentos líticos de composiciones de arenisca y riolita, inmersos en una matriz vítrea (Figura 2.4d; [Labarthe -Hernández y Jiménez-López, 1992](#); [Tristán-González et al., 2009](#); [Caballero -Miranda et al., 2009](#); [Gaytán-Martínez et al., 2017](#)). Diversos autores han datado esta unidad por el método de K/Ar, mostrando una edad de formación de 29 Ma ([Labarthe-Hernández y Jiménez-López, 1992](#); [Tristán-González et al., 2009](#)).

La unidad La Placa consiste de flujos de lava masivas de composiciones básicas (Figura 2.4f) las cuales se exponen en afloramientos aislados, mostrando un rango de espesor de 10 a 15m, y se caracteriza por fenocristales de olivino escasos en una matriz vítrea. Estudios previos han datado esta unidad por el método de K/Ar y han reportado edades de 29 a 26 Ma (Tabla 2.1; [Tristán-González et al., 2009](#); [Aguillón-Robles et al., 2014](#)).

La unidad de composición ácida más joven dentro del CVSSM, es la unidad denominada ignimbrita Panalillo. Se compone principalmente de depósitos de piroclásticos de pómez y ceniza, ricos en líticos, y presenta bajos grados de soldamiento en comparación a la unidad ignimbritica Cantera (Figura 2.4e). Estudios previos han asignado esta unidad en 29 a 26 Ma por estar ubicada entre dos unidades fechadas por el método de K/Ar (Tabla 2.1; [Labarthe-Hernández y Jiménez-López, 1992](#); [Torres-Hernández et al., 2006](#); [Tristán-González et al., 2009](#)).

La unidad más reciente en el CVSSM es la unidad Cabras, esta unidad, igual que la unidad La Placa, se presenta en afloramientos aislados dentro de la región del complejo, como a su vez, se caracteriza por flujos de lava gruesos (~25 m de espesor) con texturas afaníticas-porfídicas con fenocristales de olivino escasos y composiciones básicas (Figura 2.4g). [Tristán-González et al. \(2009\)](#) y [Aguillón-Robles et al. \(2014\)](#) reportan edades de roca total por el método de K/Ar de 22 a 21 Ma.

Tabla 2.1.- Edades de literatura de las unidades litológicas del Complejo Volcánico de la Sierra de San Miguelito. Abreviaciones: SM=San Miguelito; RT=Roca total; FLD=Feldespatos; BT=Biotita.

Unidad	Edad	Error	Método	Tipo de muestra	Referencia
Cabras	21.5	0.5	K/Ar	RT	Aguillon-Robles etal (2014)
Cabras	21.5	0.5	K/Ar	RT	Tristan-González etal. (2009)
Cabras	21.6	0.5	K/Ar	RT	Aguillon-Robles etal (2014)
Cabras	22.1	0.5	K/Ar	RT	Aguillon-Robles etal (2014)
Panalillo	25.4	0.6	K/Ar	RT	Tristan-González etal. (2009)
Panalillo	25.4	0.6	K/Ar	RT	Tristan-González etal. (2009)
Panalillo	25.7	0.4	K/Ar	FLD	Tristan-González etal. (2009)
Panalillo	27.6	0.6	K/Ar	FLD	Labarthe-Hernández etal. (1982)
Panalillo	28.3	0.9	K/Ar	RT	Tristan-González etal. (2009)
Panalillo	28.9	0.5	K/Ar	RT	Tristan-González etal. (2009)
Panalillo	29	0.9	K/Ar	RT	Tristan-González etal. (2009)
Panalillo	29	0.9	K/Ar	RT	Tristan-González etal. (2009)
Panalillo	29.9	0.6	K/Ar	FLD	Labarthe-Hernández etal. (1982)
Panalillo	30.3	0.7	K/Ar	FLD	Labarthe-Hernández etal. (1982)
Panalillo	31.9	0.7	K/Ar	FLD	Labarthe-Hernández etal. (1982)
Placa	26.9	0.4	K/Ar	RT	Tristan-González etal. (2009)
Placa	28	0.6	K/Ar	RT	Tristan-González etal. (2009)
Placa	29	0.7	K/Ar	RT	Aguillon-Robles etal (2014)
Zapote	27.4	0.4	K/Ar	FLD	Tristan-González etal. (2009)
Zapote	28.7	0.7	K/Ar	RT	Aguillon-Robles etal. (2009)
Zapote	31.2	0.7	K/Ar	RT	Tristan-González etal. (2009)
SM	21.1	0.3	K/Ar	RT	Tristan-González etal. (2009)
SM	28.7	0.8	K/Ar	RT	Tristan-González etal. (2009)
SM	29.1	0.7	K/Ar	RT	Tristan-González etal. (2009)

Tabla 2.1.- *Continuación*

Unidad	Edad	Error	Método	Tipo de muestra	Referencia
SM	29.2	0.8	K/Ar	BT	Aguillon-Robles etal. (1994)
SM	30.3	0.9	K/Ar	RT	Tristan-González etal. (2009)
SM	30.4	0.5	K/Ar	FLD	Tristan-González etal. (2009)
SM	30.7	0.7	K/Ar	RT	Tristan-González etal. (2009)
SM	31	0.6	K/Ar	BT	Labarthe-Hernández etal. (1982)
SM	31.1	0.7	K/Ar	FLD	Tristan-González etal. (2009)
SM	31.1	0.3	K/Ar	RT	Tuta etal. (1988)
SM	31.3	0.7	K/Ar	RT	Tristan-González etal. (2009)
SM	31.6	0.9	K/Ar	RT	Tristan-González etal. (2009)
SM	32	0.7	K/Ar	RT	Tristan-González etal. (2009)
SM	32.3	1.6	K/Ar	FLD	Labarthe-Hernández etal. (1982)

2.4 Muestreo

Para el presente trabajo un total de 41 muestras (Figura 2.5) representativas de las principales unidades litológicas del CVSSM fueron seleccionadas. Lugares en donde se presentan distintos grados de alteración, como a su vez, afloramientos en donde se observa deformación de la roca fueron evitados durante el muestreo.

Los límites estratigráficos y litológicos de las rocas volcánicas fueron dibujados, en gran parte en características geomorfológicas probadas en el campo y después refinados por los análisis petrológicos, geoquímicos y geocronológicos mostrados en las siguientes secciones del presente trabajo.

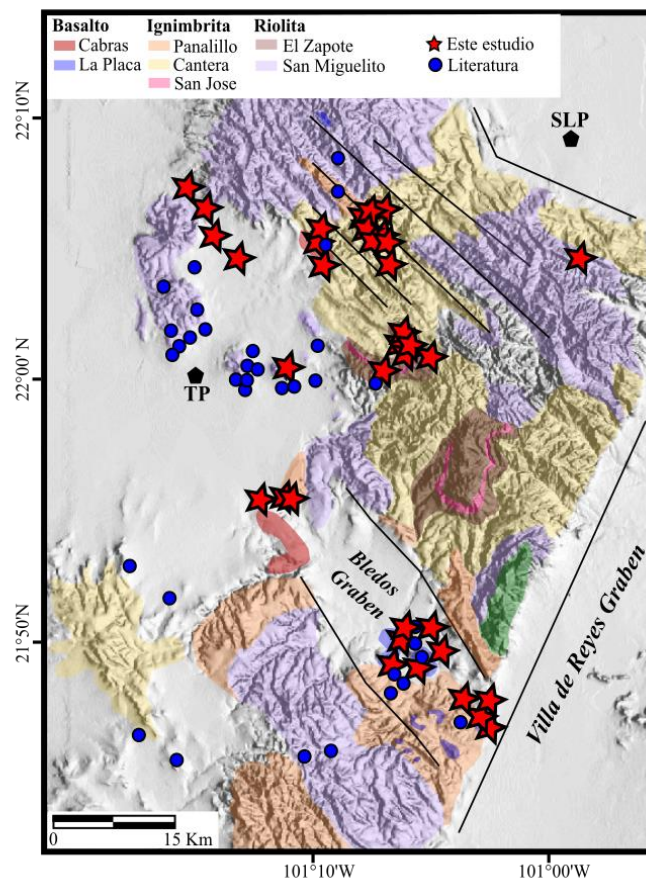


Figura 2.5.- Mapa del CVSSM representado las muestras recolectadas en este estudio (estrellas rojas) y muestras compiladas de literatura (círculos azules).

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CAPITULO 3.-

ANÁLISIS PETROGRÁFICO

3.1 Metodología del análisis petrográfico

Para la elaboración del análisis petrográfico, se elaboraron 20 láminas delgadas de las rocas volcánicas del Complejo Volcánico Sierra de San Miguelito (CVSSM), las cuales se llevaron a cabo en el Laboratorio de Preparación de la Facultad de Ciencias de la Tierra, UANL.

Para una revisión detallada de la metodología de elaboración de las secciones delgadas, se puede revisar la sección A.1 del apartado de Anexos de este trabajo. Las composiciones modales son reportadas en la Tabla A.1 en la sección de Anexos.

3.2 Características petrográficas de las unidades litológicas del CVSSM

A partir de las observaciones de campo y petrográficas, las rocas del CVSSM se pueden dividir en tres grupos principales. Para este trabajo se han dividido las rocas volcánicas del CVSSM en: a) rocas básicas; b) rocas intermedias; y c) rocas ácidas. En las siguientes secciones se presentarán las características petrográficas principales de estos tres grupos.

3.2.1 Rocas básicas

Las rocas básicas del CVSSM muestran texturas porfiríticas con una limitada presencia de vesículas y una matriz vítrea (Tabla A.1).

El ensamblaje mineral principal que presentan las rocas básicas del CVSSM (Figura 3.1) consiste de fenocristales euhedrales a subhedrales de plagioclasas (10-68%), generalmente mostrando maclas simples (maclas Carlsband) y un zonamiento normal; fenocristales subhedrales de clinopiroxeno que muestran una abundancia aproximada del 6-20%; fenocristales subhedrales de ortopiroxeno (4-7%); como también una pequeña cantidad de microcristales subhedrales a xenoformos de olivino (<6%).

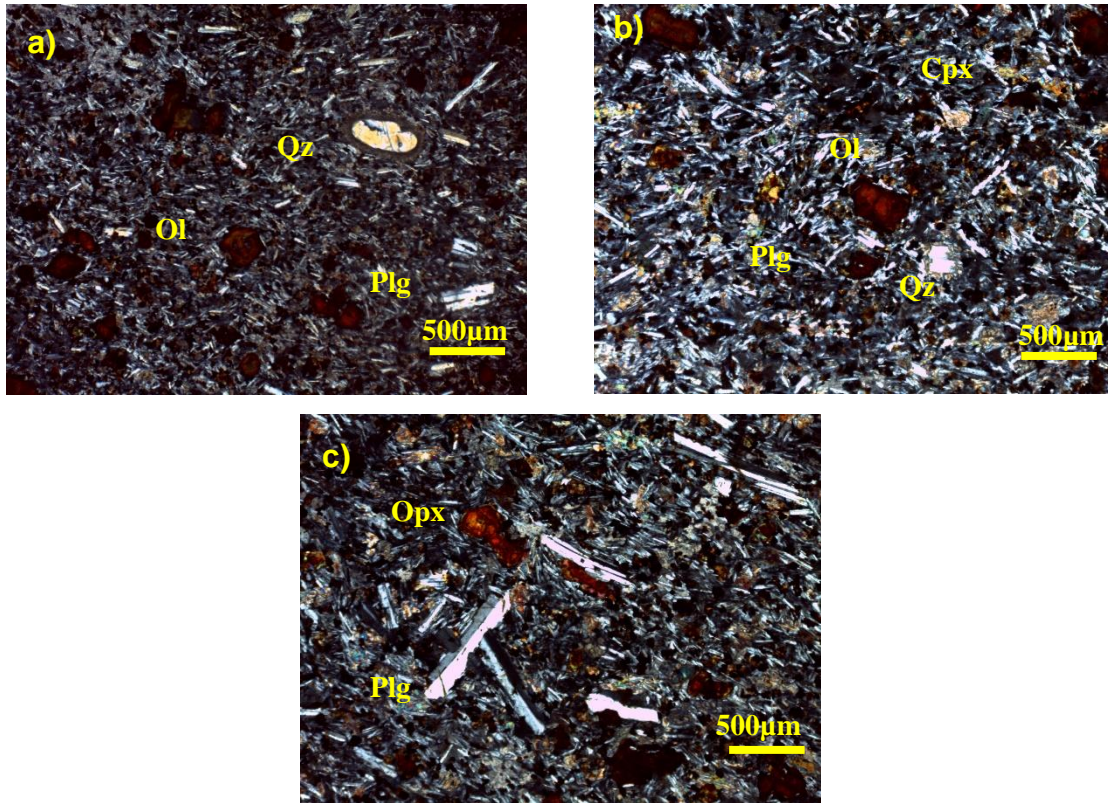


Figura 3.1.- Microfotografías en nicols cruzados (NX) con un aumento de 2.5x de muestras representativas de la unidad máfica Cabras. a) Fenocristales de plagioclasas, olivino y cuarzo con aros de reacción y microcristales de plagioclasa en la matriz; b) Fenocristales de plagioclasas, olivino y cuarzo con aros de reacción y microcristales de plagioclasa y piroxenos en la matriz; c) Fenocristales de plagioclasas con microcristales de plagioclasa en la matriz. Plg=plagioclasa; Qz= cuarzo; Cpx= clinopiroxeno; Opx= ortopiroxeno; Ol=olivino.

3.2.2 Rocas intermedias

Las rocas intermedias del CVSSM muestran texturas porfiríticas sin presencia de vesículas, y una matriz vítrea (Tabla A.1).

El ensamblaje mineralógico que muestran las rocas intermedias es similar al de las rocas básicas, con la única diferencia en las proporciones que muestran en ciertos minerales. Las rocas intermedias del CVSSM se componen por: fenocristales euhedrales a subhedrales de plagioclasas (10-68%); fenocristales subhedrales de clinopiroxeno que muestran una abundancia aproximada del 6-20%; fenocristales subhedrales de ortopiroxeno (4-7%); como también cantidad limitada de microcristales subhedrales a xenofomos de olivino (<6%).

Además, las rocas intermedias del CVSSM muestran las siguientes características peculiares: (i) diversas texturas de desequilibrio que incluyen plagioclasas con texturas normales y de criba en la misma muestra; (ii) cristales redondeados y embebidos con aros de reacción de cuarzo; (iii) plagioclasas con zonación mineral compleja, tales como zonación oscilatoria e inversa, o bien, cristales zonados de forma normal e inversa en la misma muestra (Figura 3.2).

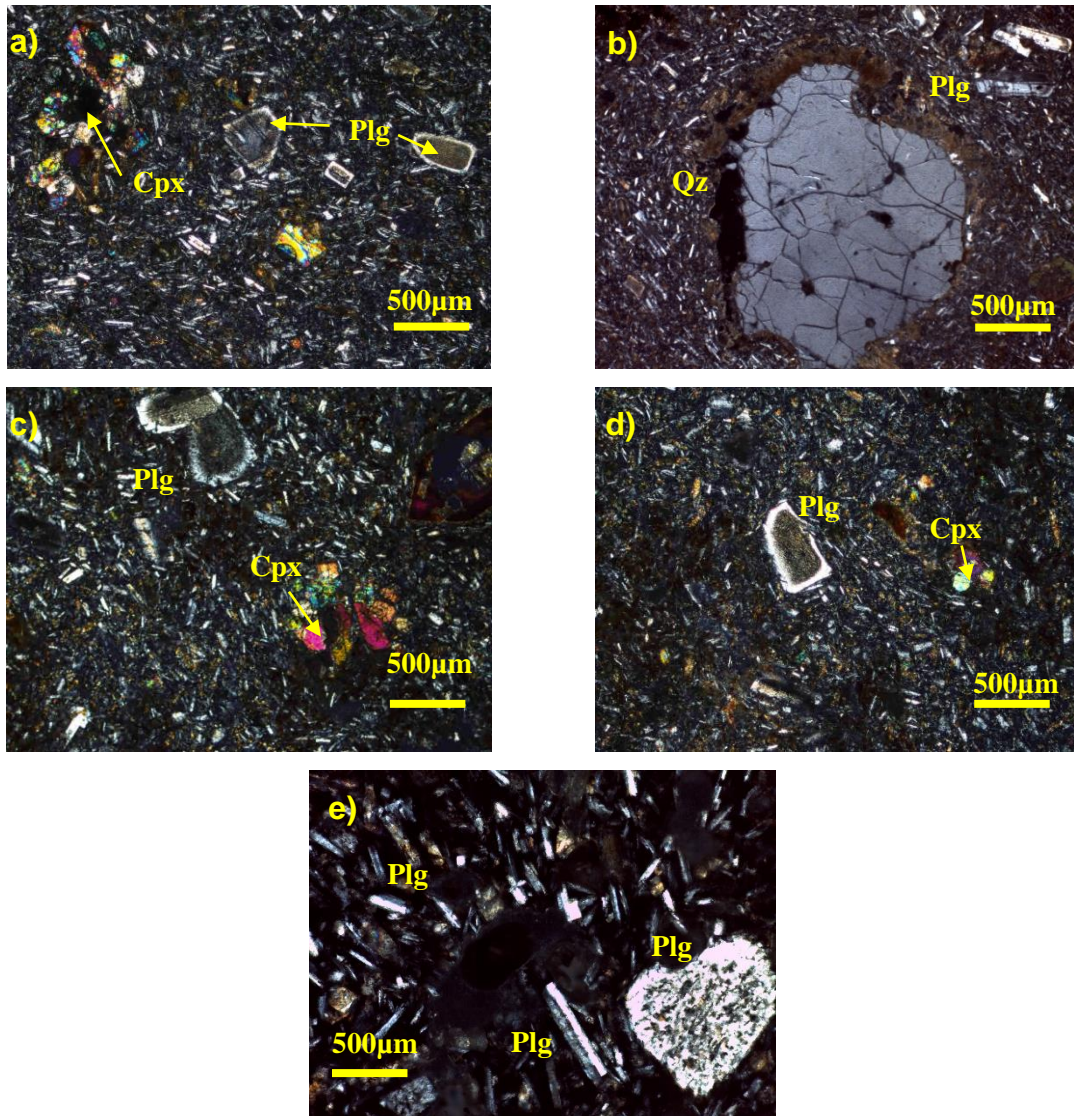


Figura 3.2.- Microfotografías en niclos cruzados (NX) con un aumento de 2.5x de muestras representativas de la unidad intermedia Placa. a) Fenocristales de plagioclasas con texturas de criba y clinopiroxenos y microcristales de plagioclasa en la matriz; b) Fenocristal de cuarzo con aro de reacción inmerso en una matriz vítrea con microcristales de plagioclasas; c) Fenocristales de plagioclasas con texturas de criba y fenocristales de clinopiroxeno inversos en una matriz vítrea con microcristales de plagioclasas; d) Fenocristales de plagioclasa con textura de criba y clinopiroxenos; e) Fenocristales de plagioclasa con textura de criba en una matriz vítrea con microcristales. Plg=plagioclasa; Qz= cuarzo; Cpx= clinopiroxeno; Opx= ortopiroxeno; Ol=olivino.

3.2.3 Rocas ácidas

Las rocas ácidas muestran texturas porfíricas, con excepción de algunas muestras que muestran texturas porfiriticas fluidales (Tabla A.1; Figura. 3.3).

El ensamblaje mineralógico principal de las rocas ácidas del CVSSM consiste de fenocristales xenoformos de cuarzo (27-41%); fenocristales euhedrales a subhedrales de feldespatos potásicos (11-15%); fenocristales euhedrales a subhedrales de plagioclasas (~44%) con diámetros de 0.5-2 mm; ortopiroxenos subhedrales (1-4%) y fenocristales de clinopiroxenos (1-4%) de un diámetro aproximado de 0.2-0.7 mm.

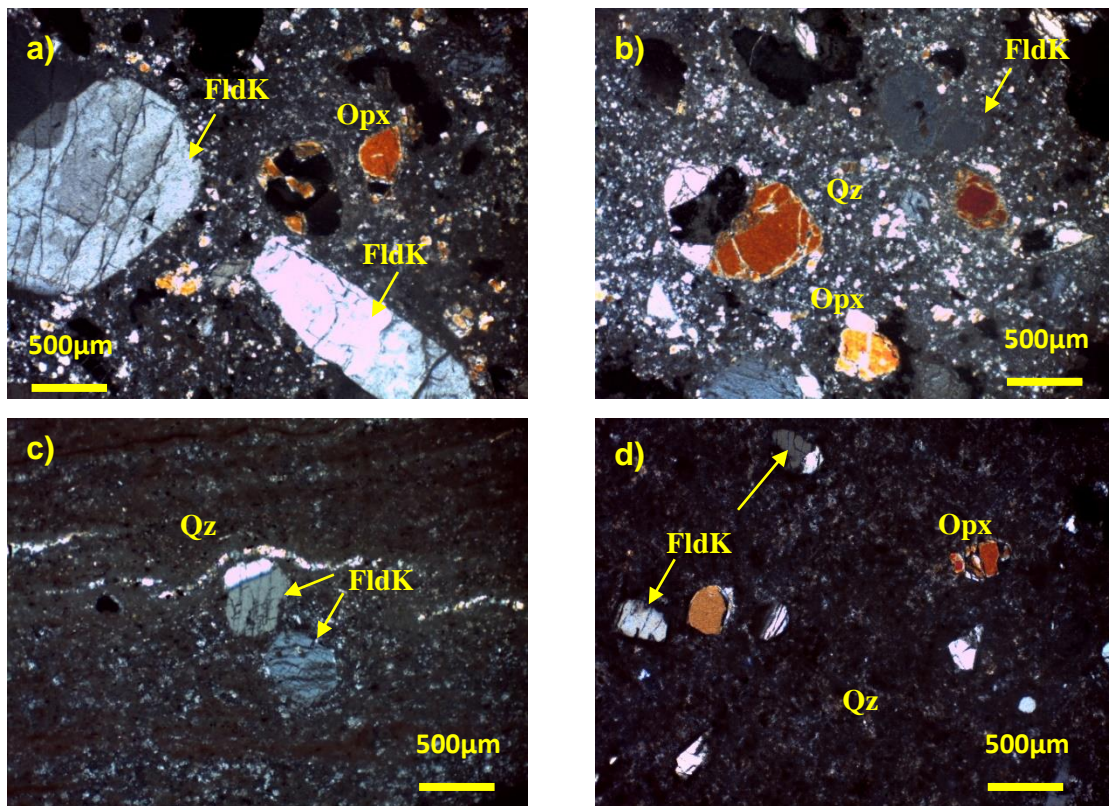


Figura 3.3.- Microfotografías en nicos cruzados (NX) con un aumento de 2.5x de muestras representativas de las unidades félsicas. a) Fenocristales de feldespatos alcalinos de la unidad San Miguelito; b) Fenocristales de feldespato potásico y ortopiroxenos con microcristales de cuarzo en la matriz de la unidad San Miguelito; c) Fenocristales de feldespato potásico con microcristales de cuarzo en la matriz de la unidad El Zapote; d) Fenocristales de feldespato potásico, cuarzo y ortopiroxeno de la unidad El Zapote. Qz= cuarzo; FldK= feldespato potásico; Opx= ortopiroxeno.

De igual forma las rocas ácidas de carácter piroclástico consisten en depósitos de flujo semi-consolidados y consolidados. Estas muestras de roca presentan texturas porfiríticas y un ensamblaje mineral principal de fenocristales de suhedrales de cuarzo (15-38%), feldespatos potásicos (15-22%) euhedrales a subhedrales, plagioclasas (2-43%), ortopiroxenos subhedrales y xenomorfos (2-5%), cristales de clinopiroxeno (2-7%) y en algunas muestras se pueden llegar a observar microcristales de olivinos en poca abundancia (Table A1; Figura 4.4).

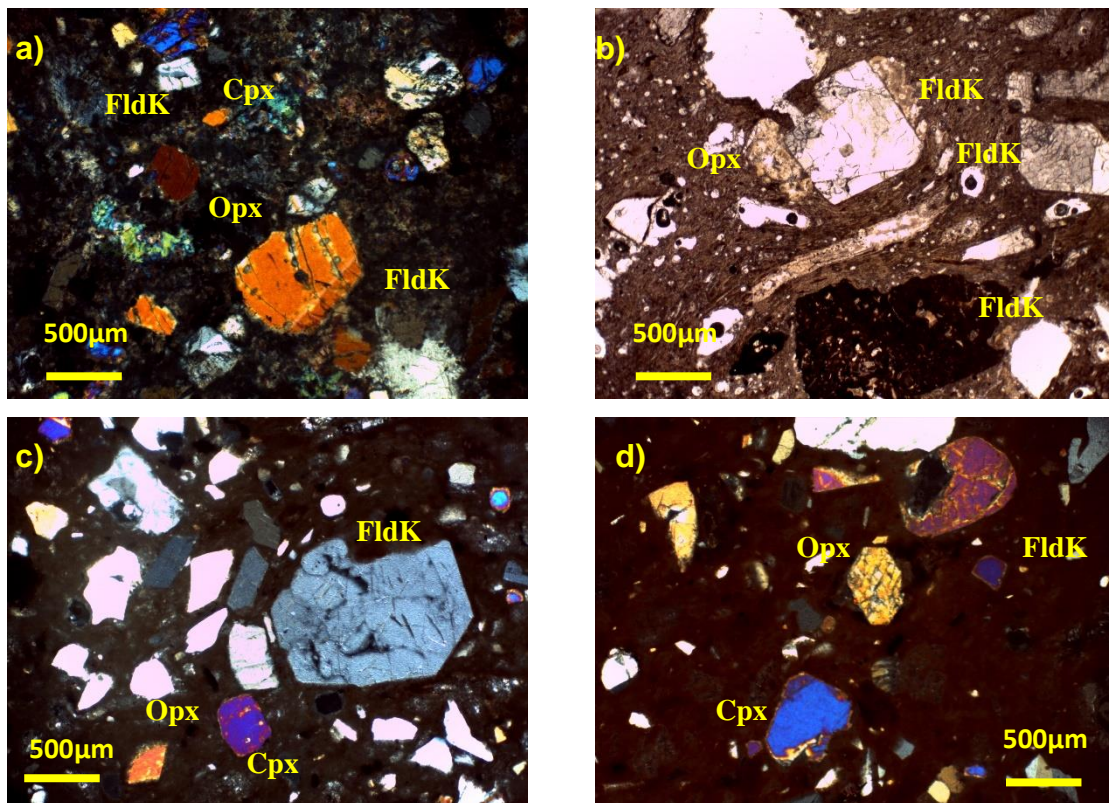


Figura 3.4 Microfotografías en nicols cruzados (NX) y nicols paralelo con un aumento de 2.5x de muestras representativas de las unidades ácidas de carácter piroclástico. a) Fenocristales de clinopiroxeno, ortopiroxeno y feldespatos potásicos de la unidad Cantera; b) Fenocristales de feldespatos potásicos y ortopiroxeno de la unidad Cantera; c) Fenocristales de feldespatos potásicos, ortopiroxeno y clinopiroxeno de la unidad Panalillo; d) Fenocristales de feldespato potásico, ortopiroxeno y clinopiroxeno de la unidad Panalillo. FldK= feldespato potásico; Opx= ortopiroxeno.

CAPITULO 4.-

CALIBRACIÓN DE FLUORESCENCIA DE RAYOS-X PARA LA MEDICIÓN DE ELEMENTOS MAYORES

4.1 Presentación del trabajo de investigación publicado





El presente artículo de investigación ha sido publicado en *Journal of Spectrometry* en el cual el autor de esta tesis participó activamente. Dentro de este trabajo me encuentro como cuarto co-autor, como a su vez, el primer autor de este trabajo es el Dr. Surendra Pal Verma Jaiswal. Los demás co-autores del presente artículo de investigación son: Dr. Sanjeet Kumar Verma, Dra. M. Abdelaly Rivera-Gómez, Dra. Lorena Díaz-González, Ing. Alejandra Amezcua-Valdez, M.C. Beatriz Adriana Rivera-Escoto, Dr. Mauricio Rosales-Rivera, Dr. John S. Armstrong-Altrin, Dr. Héctor López-Loera, Dr. Fernando Velasco Tapia y el Dr. Kailasa Pandarinath.

Este trabajo de investigación brinda una amplia contribución dentro de la calibración de máquinas de fluorescencia de rayos-X (FRX). Dentro de este trabajo se aplicó tanto regresiones lineales ordinarias (RLO) y modelos de regresión lineal ponderada (RLP) para la calibración y comparación de una máquina de FRX, a través de 59 materiales de referencia geoquímica (MRG). Se obtuvieron archivos a partir de una recopilación actualizada de datos químicos y su procesamiento a partir de pruebas conocidas de discordancia e importancia.

La contribución principal es lo demostrado entre la comparación de los modelos de RLO y RLP, en donde se mostró claramente que este último, con valores generalmente más altos de r , es preferible para calibraciones rutinarias de los procedimientos analíticos de máquinas de FRX.

Research Article

Statistically Coherent Calibration of X-Ray Fluorescence Spectrometry for Major Elements in Rocks and Minerals

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We applied both the ordinary linear regression (OLR) and the new uncertainty weighted linear regression (UWLR) models for the calibration and comparison of a XRF machine through 59 geochemical reference materials (GRMs) and a procedure blank sample. The mean concentration and uncertainty data for the GRMs used for the calibrations (Supplementary Materials) (available here) were achieved from an up-to-date compilation of chemical data and their processing from well-known discordancy and significance tests. The drift-corrected XRF intensity and its uncertainty were determined from mostly duplicate pressed powder pellets. The comparison of the OLR (linear correlation coefficient $r \sim 0.9523$ – 0.9964 and 0.9771 – 0.9999 , respectively, for before and after matrix correction) and UWLR models ($r \sim 0.9772$ – 0.9976 and 0.9970 – 0.9999 , respectively) clearly showed that the latter with generally higher values of r is preferable for routine calibrations of analytical procedures. Both calibrations were successfully applied to rock matrices, and the results were generally consistent with those obtained in other laboratories although the UWLR model showed mostly narrower confidence limits of the mean (slope and intercept) or lower uncertainties than the OLR. Similar sensitivity (~ 2.69 – 46.17 $\text{kc}\cdot\text{s}^{-1}\cdot\%^{-1}$ for the OLR and ~ 2.78 – 59.69 $\text{kc}\cdot\text{s}^{-1}\cdot\%^{-1}$ for the UWLR) also indicated that the UWLR could advantageously replace the OLR model. Another novel aspect is that the total uncertainty can be reported for individual chemical data. If the analytical instruments were routinely calibrated from the UWLR model, this action would make the science of geochemistry more quantitative than at present.

1. Introduction

All modern analytical instruments require some kind of calibration of the instrumental response (y -variable) as a function of the concentration (x -variable) [1–3]. This calibration is generally achieved through an ordinary least-squares linear regression (OLR) model. However, such a procedure is not strictly valid because all requirements for the statistical validity of the OLR model are not fulfilled. Usually, the assumptions “independent concentration variable x is error-free or less than one-tenth of the error in the dependent response variable y ” and “error in y is homoscedastic” (i.e., equal errors for all y values) are not satisfied and, therefore, more sophisticated and statistically coherent regression procedures, such as weighted least-squares linear regression (WLR) models, should be used [4–18].

X-ray fluorescence (XRF) spectrometry is among the most popular analytical techniques for the determination of all major and some trace elements in rocks [4, 19–27]. Natural geochemical reference materials (GRMs) are commonly used for XRF calibrations and posterior characterization of those and other GRMs as well as of similar rock and mineral matrices [4, 19, 28–30]. As for most other analytical instruments, XRF spectrometers are also calibrated under the statistically incoherent OLR model.

To apply the WLR and compare it with the OLR, both central tendency (e.g., mean) and dispersion (e.g., confidence limits of the mean) estimates on both x -axis (concentration, generally expressed in the unit of % m/m, i.e., mass/mass unit expressed in percent) and y -axis (response, in this case XRF intensity, generally reported in the unit of $\text{kc}\cdot\text{s}^{-1}$, i.e., kilo counts per second) variables are required. More precise (and accurate) estimates of the central tendency will also be useful for both types of regressions. Therefore, precise concentrations of GRMs with the respective lowest possible “confidence limits of the mean” (referred hereafter as the “uncertainty” of the measured variable) [2, 17, 18] are required to apply the regression procedure. Sometimes, we had to use also the term “error” (instead of the uncertainty) because the use of the error is widespread in the literature.

We report the following five aspects: (a) evaluation of 59 GRMs to achieve the least possible uncertainties in the mean concentrations of all major elements (SiO_2 to P_2O_5); (b) the comparison of regression models (OLR and WLR) applied to net drift-corrected XRF intensities before the correction of matrix effects; (c) the second (or final) comparison of both models after achieving the matrix correction as well as for the estimation of sensitivities of the regression models; (d) application of the entire procedure to four GRMs treated as “unknown” samples and their comparison with the previous literature compilations; and (e) development of a computer program to achieve the abovementioned objectives. Thus, the regression equations (intercept and its uncertainty, slope and its uncertainty, and linear correlation coefficient values) for each constituent from SiO_2 to P_2O_5 and their application to similarly complex rock matrices are presented in this work.

2. Evaluation of Major Element Data for GRMs

A total of 59 GRMs (listed in alphabetical order in Table S1; this and four other tables are provided in Supplementary Materials), along with a procedure blank, were used in this study. The procedure blank was a pellet prepared in duplicate with only pure $\text{N,N}'$ -Ethylene bis(stearamide) beads without any sample (Section 3). The individual data reported in earlier compilations [31–47] were first compiled in new databases.

The statistical parameters obtained in these early compilations could not be directly used for instrumental calibrations due to the following reasons: (i) the statistical methods used to achieve the statistical estimates were outdated (see [17, 18, 48, 49] for possible reasons), and the inferred statistical values were of low quality (high values of dispersion); (ii) there are still determinations reported during about 30 or more years (postcompilation years) that were not obviously available to those compilers; (iii) the precision of more recent determinations is likely to have improved due to the availability of online computers on most modern instruments; (iv) newer more reliable statistical techniques are now available for improving both precision and accuracy of the statistical inferences, e.g., the use of discordancy tests with the highest power and lowest swamping and masking effects [18, 48, 50–52]; and (v) importantly, new computer programs have been developed by our group [52–54], available at <http://tlaloc.ier.unam.mx> for download or online processing of data (after previous registration onto our server), which can be advantageously used for efficient processing of experimental databases.

The same kinds of objections are applicable even today for the originator’s websites, such as <https://gbank.gsj.jp/geostandards/welcome.html> for Japanese GRMs or https://crustal.usgs.gov/geochemical_reference_standards for United States GRMs. The statistical information at these websites is based on early compilations (around 30 or more years ago). Furthermore, we were unable to use the recent work [55] because this paper reported significantly larger uncertainty values as compared to those achievable from our new validated statistical procedure [51–54]; besides, updated statistical information on the mean and its uncertainty was not available in [55] for many GRMs used in our work.

The initial databases were complemented by individual data from a large number of posterior publications (~480; Table S1), whose complete listing is available at our server <http://tlaloc.ier.unam.mx> under the heading of “Quality Control.” These major element data were classified according to the analytical method groupings [56]. Data from each method group were considered as a univariate statistical sample. Appropriate discordancy and significance tests were applied from thoroughly automatized software UDASys2 [52] and UDASys3 (unpublished), which, in their “recommended procedure,” apply the most powerful five (two new and three conventional) recursive tests with prior application of respective single-outlier tests having nil swamping and low masking effects [48, 57–60]. Although the application of discordancy tests is identical for both UDASys2 and UDASys3, the difference lies in that the latter applies the

significance (ANOVA, F and t) tests in order to provide the final results automatically.

The resulting statistical information after the application of well-known discordancy tests at the strict 99% confidence level (mean and uncertainty values rounded according to the flexible rules [18]) is listed in Table S2. These GRM compositional data showed by far the lowest 99% uncertainty (Table S2), much lower than any existing compilation [31–47, 55]. We may also stress once again that this was achieved through an objective combination of discordancy tests having the highest performance and lowest swamping and masking effects [17, 18, 48, 53], i.e., from the methodology having the lowest type I and type II errors and the highest power.

Therefore, the population mean of these GRMs is now known within the narrowest possible 99% confidence limits of the mean to best represent the concentration (x) axis in the instrumental calibrations as suggested [2, 5, 7, 10, 17, 18, 53]. These data (in units of % m/m; Table S2) will also be useful for those who wish to achieve instrumental calibrations or simply use them for quality control of their results for rock and mineral matrices.

3. XRF Instrumentation and Intensity Measurements

A wavelength dispersive X-ray fluorescence (WDXRF) spectrometer Rigaku ZSX Primus II model (rhodium X-ray tube; 4 kW maximum power) was used for this work. We made the effort to best represent the response (y) axis (x -ray intensity in the units of kilo counts per second, $\text{kc}\cdot\text{s}^{-1}$) for the calibrations. For each GRM, duplicate (41 samples) or even triplicate (8 samples) pressed powder pellets were prepared. First, an appropriate amount of each GRM was dried overnight in an oven at about 105°C . For each pellet, accurately weighed 3.5 g of moisture-free GRM was thoroughly mixed with accurately weighed 3.0 g pure $\text{N,N}'$ -ethylene bis(stearamide) beads, $<840\ \mu\text{m}$ as wax (Sigma-Aldrich), and stored in a desiccator. Pressed powder pellets were prepared at $20\ \text{tons}\cdot\text{inch}^{-2}$ pressure (about 310 MPa). However, for 10 GRMs, sufficient material was not available; therefore, only a single pellet could be prepared but the measured intensity uncertainty (u_{99}) at the 99% confidence level was increased by a factor of 2 to take into account the sample preparation variance. Similarly, accurately weighed 6.5 g of pure $\text{N,N}'$ -ethylene bis(stearamide) beads, $<840\ \mu\text{m}$ as wax, was pressed to prepare a procedure blank sample. This was done in duplicate.

For the intensity measurements, the optimum instrumental conditions were first established through preliminary experiments prior to the routine measurements (Table S2). Each pellet was run at least 8 to 10 times in a random sequence, along with two drift monitors prepared from two volcanic rocks (basalt and rhyolite) from the San Luis Potosí Volcanic Field, San Luis Potosí (central Mexico).

The peak and background measuring conditions and time periods are also listed in Table S3. Appropriate mean drift corrections from two monitors were applied to all intensity measurements. Both monitors were run randomly

8 to 10 times each day. First, the expected monitor intensity was established as an average value of the first two days when the intensities were fairly stable and reproducible. Then, the average drift correction factors were calculated for each chemical element from the two monitors run in the XRF instrument periodically before and after a set of GRMs used for the calibration. These correction factors were then applied to the bracketed GRMs for the entire period of calibration, including the first two days and analysis of “unknown” samples.

Now, although the X-ray counts may obey a Poisson distribution, we are dealing with average values of count rates, which are likely to follow a normal distribution because of the central limit theorem. A normal distribution of measured intensities was also assured for each pellet from the application of discordancy tests as explained above for GRM concentrations. The intensity results for all pellets from a given GRM were then combined, the tests applied again to the combined data, and new mean and 99% uncertainty values were calculated for X-ray intensity of each GRM. This was done to take into account the variance of the sample preparation method, which was significantly higher than the instrumental variance of intensity measurements for individual pellets. The drift-corrected intensity values and their 99% uncertainties ($\text{kc}\cdot\text{s}^{-1}$) for all GRMs, along with the concentration data and their 99% uncertainties (% m/m), are listed in Table S2.

4. Regression Models

Two different regression models (OLR and UWLR) were used and compared in this work. The OLR model most frequently used for instrumental calibrations (x -axis concentration and y -axis response; GRM concentration and X-ray intensity, respectively, in XRF spectrometry) requires the following assumptions to be fulfilled [4, 7, 10, 12–18]: (i) all errors are in the y -axis; (ii) x -axis is either error-free or has at most 10% error of the y -axis errors; (iii) errors in both axes are normally distributed; and (iv) errors in the y -axis are homoscedastic. Some or all of these assumptions are violated in most instrumental calibrations through the OLR model.

Thus, from the literature on the GRMs, it has been demonstrated that the concentration axis is not error-free (see non-zero uncertainties for all GRM concentrations in Table S2) [31–47, 51–53]. One can also clearly see that the errors in the intensity axis are not homoscedastic (see unequal, i.e., heteroscedastic uncertainties for any element in different GRMs in Table S2). For a heteroscedastic linear regression system, even if each error or noise term is still Gaussian, the OLR model is no longer the maximum likelihood estimate and consequently, it is no longer efficient [10]. The main advantage that the WLR has over the OLR is the ability to handle regression situations in which the data points are of varying quality as is the case with most instruments including the XRF spectrometers.

However, the major disadvantage of the WLR is that the approach is based on the assumption that the weights are known exactly. They can be estimated using several different equations or algorithms, but when the weights are produced

from small numbers of replicated observations, the regression parameters can be unpredictably affected [10]. In the example of the XRF calibration that we are presenting, the numbers of observations were relatively large for both the x and y axes (concentration and X-ray intensity parameters). Besides, instead of the sample variance, we used the uncertainty values (that take into account the number of observations in the formula for uncertainty or confidence limits of the mean calculations) [2, 18] for estimating the weight factors. The problem of the sensitivity to outliers in the regression equations [10] was also appropriately handled by discordancy tests programmed in the UDASys and BiDASys software [53, 54, 61].

Therefore, although frequently used, the OLR model is not statistically correct or coherent. The statistically coherent WLR, especially the uncertainty-based WLR (UWLR [17]) model, should be used. The confidence level, such as 95% or 99% (significance level of 5% or 1%, respectively, or α of 0.05 and 0.01, respectively), can be explicitly expressed in the confidence limits of the mean or uncertainty used in the UWLR model as well as to estimate the weight factors [17]. We will deal with the 99% uncertainty to have the type I error small (about 1%). Unfortunately, software of most analytical instruments, including XRF spectrometers, allows only the OLR calibration. Therefore, any sophisticated regression model, such as the UWLR, will have to be applied outside the instrumental software. Thus, the probability concept (99% confidence level) can be explicitly used in the UWLR model for weight factors based on the inverse of the squared 99% uncertainty of the mean.

We now present a synthesis of the regression equations for instrumental calibrations [2, 10, 17, 18, 61].

4.1. Ordinary Least-Squares Linear Regression (OLR) Model.

Let us assume that we have a series of n reference materials or standard calibrators having individual mean concentrations x_i with respective uncertainties u_{x_i} , where i varies from 1 to n . In order to calibrate an instrument, each of these n calibrators were run several times, obtaining individual mean responses y_i with respective uncertainties u_{y_i} , where i varies from 1 to n . Thus, we have n bivariate concentration-response data pairs or calibrators (x_i, y_i) with the respective uncertainties (u_{x_i}, u_{y_i}).

We can apply the OLR model to these data for obtaining a calibration equation. The OLR fits a least-squares linear equation to the n pairs (x_i, y_i) but does not take into account the respective uncertainties (u_{x_i}, u_{y_i}).

The general regression equation for the OLR is as follows (the subscript O is for the OLR model):

$$y_O(\pm u_{y_O}) = b_O(\pm u_{b_O}) + [m_O(\pm u_{m_O}) \times x], \quad (1)$$

where m is the slope, u_m is the resulting uncertainty in the slope, b is the intercept, u_b is the resulting uncertainty in the intercept, x is the independent variable, y_O is the dependent variable from the OLR model, and u_{y_O} is the resulting uncertainty in y . The following equations allow the calculations of these parameters:

$$m_O = \frac{\sum_{i=1}^n \{(x_i - \bar{x}) \times (y_i - \bar{y})\}}{\sum_{i=1}^n (x_i - \bar{x})^2}, \quad (2)$$

where \bar{x} and \bar{y} are, respectively, the mean values of the x and y variables:

$$u_{m_O} = \left\{ \sqrt{\frac{\sum_{i=1}^n (y_i - \hat{y}_i)^2}{(n-2) \sum_{i=1}^n (x_i - \bar{x})^2}} \right\} \times t_{(n-2)}^{CL}, \quad (3)$$

where \hat{y}_i is the value of y_O for x_i in equation (1) and t is the Student's t test value for $(n-2)$ degrees of freedom, and the superscript CL is the confidence level, generally 95% or 99%:

$$b_O = \bar{y} - (m_O \times \bar{x}),$$

$$u_{b_O} = \left\{ \sqrt{\frac{\sum_{i=1}^n (y_i - \hat{y}_i)^2 \times \sum_{i=1}^n x_i^2}{n(n-2) \sum_{i=1}^n (x_i - \bar{x})^2}} \right\} \times t_{(n-2)}^{CL}. \quad (4)$$

It is a general practice in most instrumental calibrations to ignore all uncertainties in equation (1) and use an OLR equation without any error (or uncertainty) as follows:

$$y_O = b_O + [m_O \times x]. \quad (5)$$

The resulting standard deviation values of repeat measurements of unknown samples are reported as the final errors. However, these are only partial errors because the errors in the calibration equation (1) are not taken into account. In this work, we will use equation (1) to report total errors (in fact, 99% uncertainties) for the OLR model.

4.2. Uncertainty Weighted Least-Squares Linear Regression (UWLR) Model.

For the UWLR model, the n pairs (x_i, y_i) of calibrators as well as the respective uncertainties (u_{x_i}, u_{y_i}) are taken into account in order to achieve the best least-squares linear fit.

The uncertainties u_{x_i} in the x -axis are first propagated to the y -axis, combined with the u_{y_i} , and the total uncertainty u_i values on the y -axis are used for the weighting factors [2, 10, 17, 18, 61]:

$$u_i = \sqrt{(m_O \times u_{x_i})^2 + (u_{y_i})^2}. \quad (6)$$

The weights are calculated from u_i as follows:

$$w_i = \frac{n \times (u_i)^{-2}}{\sum_{i=1}^n (u_i)^{-2}}, \quad (7)$$

where w_i values have the following property:

$$\sum_{i=1}^n w_i = n. \quad (8)$$

Thus, the UWLR fits a linear equation to the n pairs (x_i, y_i) with the respective weighting factors w_i as follows (the subscript UW is for the UWLR model):

$$y_{UW}(\pm u_{y_{UW}}) = b_{UW}(\pm u_{b_{UW}}) + [m_{UW}(\pm u_{m_{UW}}) \times x]. \quad (9)$$

Note that this regression line will pass closer to the data with lesser uncertainty u_i . The intercept and slope variables and their uncertainties are calculated from the following equations:

$$m_{\text{UW}} = \frac{\{\sum_{i=1}^n (w_i \times x_i \times y_i)\} - \{n \times \bar{x}_{\text{UW}} \times \bar{y}_{\text{UW}}\}}{\{\sum_{i=1}^n (w_i \times x_i^2)\} - \{n \times (\bar{x}_{\text{UW}})^2\}}, \quad (10)$$

where \bar{x}_{UW} and \bar{y}_{UW} are, respectively, the weighted mean values of the x and y variables:

$$u_{m_{\text{UW}}} = \left\{ \sqrt{\frac{\sum_{i=1}^n (y_i - \hat{y}_{i\text{UW}})^2}{(n-2)\sum_{i=1}^n (x_i - \bar{x}_{\text{UW}})^2}} \right\} \times t_{(n-2)}^{\text{CL}}, \quad (11)$$

where $\hat{y}_{i\text{UW}}$ is the value of y_{UW} for x_i in equation (9):

$$b_{\text{UW}} = \bar{y}_{\text{UW}} - (m_{\text{UW}} \times \bar{x}_{\text{UW}}),$$

$$u_{b_{\text{UW}}} = \left\{ \sqrt{\frac{\sum_{i=1}^n (y_i - \hat{y}_{i\text{UW}})^2 \times \sum_{i=1}^n x_i^2}{n(n-2)\sum_{i=1}^n (x_i - \bar{x}_{\text{UW}})^2}} \right\} \times t_{(n-2)}^{\text{CL}}. \quad (12)$$

The best regression equation for a calibration curve should have the following characteristics (without distinguishing the subscripts O and UW): (i) intercept b small approaching to zero; (ii) slope m large; and (iii) both u_b and u_m small. Further, the quality of the regression, whether a calibration curve or any other bivariate relationship, is also expressed as the linear regression coefficient (r ; r_{O} and r_{UW} , respectively, for the OLR and UWLR), which is ideally +1.00000 for a calibration curve [5, 18, 61].

5. Application of Regression Models for XRF Calibration

5.1. Original Drift-Corrected Net Intensities and GRM Concentrations: The First Set of Two Regression Equations for Each Element. The evaluations for both regression types on the drift-corrected net intensity-concentration (Int-Conc) relationships (Table S2) for all major elements from SiO_2 to P_2O_5 were performed (Table S4), for which the new online software BiDASys was used [61] at <http://tlaloc.ier.unam.mx>. BiDASys allows the application of the conventional OLR as well as the newly proposed UWLR model [17] and provides the output of all regression parameters in an Excel® file. Contrary to the common practice, we will refrain from showing the numerous x - y (variable x is drift-corrected net intensity "Int" and variable y is the GRM concentration "Conc") plots. This is because Table S4 statistically quantifies the visual interpretation of such diagrams. The quality parameters (standard errors se_i and se_m , uncertainty u_b and u_m , and linear correlation coefficient r and its squared value R^2 parameters) are reported in Table S4. Because we are using these several different quality parameters, the concern against the use of solely R^2 parameter [62] is not important for comparison purposes.

We will explain the implications of the statistical results for the first element SiO_2 ; the statistics for other elements

(Table S4) can be similarly understood. The OLR regression equation from the first row of statistical information in Table S4 is as follows (after the element SiO_2 , subscript O is for the OLR and p is for provisional concentration; note many decimal places are used for the regression variables in such equations, because these values are not final results, and we should not introduce rounding errors during the calculation stage):

$$C_{\text{SiO}_{2\text{op}}} (\pm u_{C_{\text{SiO}_{2\text{op}}}}) = 11.47071 (\pm 4.90411) + \left[0.14325 (\pm 0.01605) \right] \times I_{\text{SiO}_2} (\pm u_{I_{\text{SiO}_2}}). \quad (13)$$

Similarly, the UWLR equation from the second row of statistical information in Table S4 is as follows:

$$C_{\text{SiO}_{2\text{UWp}}} (\pm u_{C_{\text{SiO}_{2\text{UWp}}}}) = -0.01316 (\pm 2.83181) + \left[0.18539 (\pm 0.00927) \right] \times I_{\text{SiO}_2} (\pm u_{I_{\text{SiO}_2}}). \quad (14)$$

The implications of these regression equations can be understood from the comparison of the uncertainties of the intercept and slope, which are lower for the UWLR (equation (14)) than for the OLR (equation (13)). This means that the uncertainty of the calculated concentration will be lower for the UWLR than for the OLR. Correspondingly, the r value for the UWLR (0.99004, $n = 60$; $R^2 = 0.98017$) is much higher than that for the OLR (0.95229, $n = 60$; $R^2 = 0.90687$; Table S4). Similar trend in the r (and R^2) values was obtained for all other elements except MnO (Table S4).

5.2. Matrix-Effect-Corrected Intensities and GRM Concentrations: The Second Set of Two Regression Equations for Each Element. Matrix correction is certainly required because the abovementioned least-squares linear regression fits are far from "perfect" ($r \neq +1.00000$; in fact, $r < 1$; $n = 60$; $r = 0.95229$ – 0.99638 for the OLR and $r = 0.97715$ – 0.99760 for the UWLR; Table S4). There is a vast literature on the subject of matrix effects in XRF and their correction procedures [63–75]. In this study, the Lachance-Traill algorithm [73] was used for the matrix effect correction [63, 71]. This was done outside the XRF instrument software. In a review of the existing algorithms, Rousseau [63] showed that the Lachance-Traill algorithm could be considered as one of the most appropriate procedures for the matrix effect correction because other algorithms have limited application range or lack of accuracy. Thus, for each element from SiO_2 to P_2O_5 , a system of overdetermined equations was solved and the resulting alpha coefficients were used to correct all intensities for matrix effects.

From the alpha coefficients, matrix-corrected intensities and improved concentration values for the GRMs and their uncertainties were calculated iteratively under the condition that the convergence parameter (absolute relative difference

of the GRM calculated and input concentrations) for each compositional constituent (SiO_2 to P_2O_5) be minimized.

New regression equations for achieving the corrected concentrations were established from the relationship of the calculated GRM concentrations (ConcCalc) and the original GRM concentrations (Conc) given in Table S2, for which the online BiDASys software [61] was used at <http://tlaloc.ier.unam.mx>. These equations can be formulated from the regression coefficient values given in Table S4 (see ConcCalc-Conc rows corresponding to the OLR and UWLR). Again, we will highlight their significance for SiO_2 only.

The OLR regression equation from the third row of statistical information in Table S4 is as follows:

$$C_{\text{SiO}_2\text{O}} \left(\pm u_{C_{\text{SiO}_2\text{O}}} \right) = 1.87016 (\pm 4.2048) + \left[0.96759 (\pm 0.07369) \right] \times C_{\text{SiO}_2\text{c}} \left(\pm u_{C_{\text{SiO}_2\text{c}}} \right), \quad (15)$$

where the subscripts O and c stand for the OLR model and calculated concentration (ConcCalc), respectively.

Similarly, the UWLR equation from the fourth row of statistical information in Table S4 is as follows:

$$C_{\text{SiO}_2\text{UW}} \left(\pm u_{C_{\text{SiO}_2\text{UW}}} \right) = -0.02185 (\pm 1.47356) + \left[1.00325 (\pm 0.02701) \right] \times C_{\text{SiO}_2\text{UWc}} \left(\pm u_{C_{\text{SiO}_2\text{UWc}}} \right), \quad (16)$$

where the subscripts UW and c stand for the UWLR model and calculated concentration (ConcCalc), respectively.

Equations (15) and (16) show that the concentration values from the UWLR would be more reliable (lesser uncertainty values in both intercept and slope) than the OLR model. The r value is higher for the UWLR (0.99704, $n = 60$; $R^2 = 0.99408$; Table S4) than the OLR (0.97710, $n = 60$; $R^2 = 0.95472$).

After the matrix correction, in fact most regression equations are better because all r and R^2 values are higher for both OLR and UWLR than without the correction (Table S4; Figure 1 for r only). For the OLR, the matrix correction increased the r values ($n = 60$) from 0.95229–0.99638 ($R^2 = 0.90687 - 0.99277$) to 0.97710–0.99992 ($R^2 = 0.95472 - 0.99994$). Similarly, for the UWLR, this increase was from 0.97715–0.99760 ($R^2 = 0.95472 - 0.99521$) to 0.99704–0.99993 ($R^2 = 0.99408 - 0.99986$). Thus, after matrix correction, all r values increased for both OLR and UWLR. For the UWLR, the r values approached the ideal value of 1.00000 (Figure 1). One has to keep in mind that when the r values are closer to the maximum possible value of 1 (the “ideal” fit), the improvement expressed by the actual (absolute) value of r will apparently be small. However, as long as the r value increases for the UWLR as compared to the OLR (Figure 1; Table S4), we can objectively infer that the UWLR is a better regression model than the OLR.

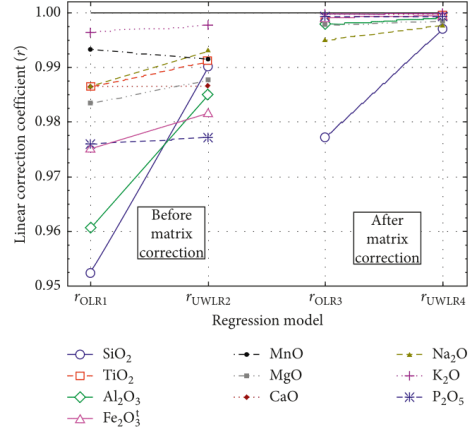


FIGURE 1: Linear correlation coefficient (r) values for the ordinary least-squares linear regression (OLR) and uncertainty-based weighted least-squares linear regression (UWLR) models for the XRF calibration of major elements (SiO_2 to P_2O_5) in rocks and minerals. OLR1: OLR model 1 for Int-Conc before matrix correction; UWLR2: UWLR model 2 for Int-Conc before matrix correction; OLR3: OLR model 3 for ConcCalc-Conc after matrix correction; and UWLR4: UWLR model 4 for ConcCalc-Conc after matrix correction. Symbols are shown as inset. The horizontal line at the r value of 1 represents the “ideal” or “perfect” linear fit.

Before the matrix correction, the intercepts of the Int-Conc regression lines were closer to zero for the UWLR (range ~ -0.013 to $+0.011$) than for the OLR (range ~ -2.098 to $+11.47$) model (Table S4; Figure 2). The same is true for the intercept values (ConcCalc-Conc relationship) after the matrix correction (~ -0.025 to $+0.021$ for the UWLR and ~ -0.110 to $+1.87$ for the OLR).

Finally, the uncertainties on both intercept and slope parameters were mostly lower for the UWLR than the OLR (Table S4). We highlight these differences (lower uncertainties for the UWLR) from dimensionless (free of the measurement units) parameters δu_b and δu_m defined as follows:

$$\delta u_b = \frac{u_{b\text{O}} - u_{b\text{UW}}}{u_{b\text{UW}}} \times 100, \quad (17)$$

$$\delta u_m = \frac{u_{m\text{O}} - u_{m\text{UW}}}{u_{m\text{UW}}} \times 100.$$

Plots of these two parameters are presented in Figure 3. If $u_{b\text{O}} > u_{b\text{UW}}$, the δu_b will be positive, otherwise it will be negative. The same is true for δu_m . For the comparison of two models OLR and UWLR before the matrix correction, the uncertainty for the UWLR were lower than the OLR for 7 elements (positive δu_b and δu_m), whereas for after the matrix correction, it was so for 8 elements (out of 10; Figure 3). The exceptions were for 3 elements MnO, CaO, and P_2O_5 (negative δu_b and δu_m) for the uncertainties before matrix

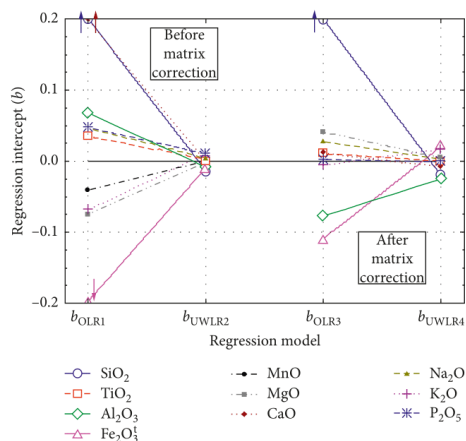


FIGURE 2: Intercept (b) values for the ordinary least-squares linear regression (OLR) and uncertainty-based weighted least-squares linear regression (UWLR) models for the XRF calibration of major elements (SiO_2 to P_2O_5) in rocks and minerals. Symbols are shown as inset. For abbreviations, see Figure 1. Note some intercept values plotted outside the graph; this is indicated by arrows next to the data point. The horizontal line at the intercept value of zero represents the “ideal” intercept.

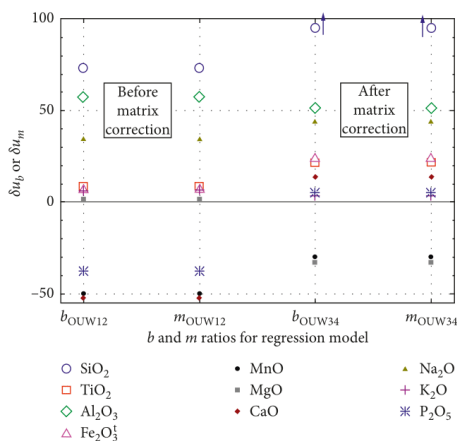


FIGURE 3: New parameters δu_b (for the intercept) and δu_m (for the slope) for the evaluation of intercept (b) and slope (m) of two regression models (OLR: ordinary least-squares linear regression and UWLR: uncertainty-based weighted least-squares linear regression) before (OUW12) and after (OUW34) the matrix correction. The horizontal solid line at the y value of zero represents the line with no difference in the uncertainties of the two models. The arrows indicate that these data plotted above the scale are used for the diagram.

correction and for 2 elements MnO and MgO for those after the matrix correction (Figure 3). Even for the exceptions of the elements MnO and MgO, the UWLR values should be usable (Table S4), i.e., it is not actually necessary to resort to the OLR model for these two exceptions (2 out of 10 cases). Thus, we can use the UWLR model for all purposes.

6. Sensitivities of Major Elements

We calculated the sensitivities as the slope of the Conc-IntCorr (GRM concentrations of Table S2 and matrix-corrected intensities of Table S5; see Supplementary Materials at <http://taloc.ier.unam.mx>) from the regression curve (line) for all 10 elements and for both models (Table 1). Because the r values are significantly high (all >0.961 , $n = 60$; Table 1) and the residuals are randomly distributed (graphs not shown), the straight line is the most likely, statistically valid fit for the concentration-matrix-corrected intensity data [5, 17, 18]. Therefore, the slope of the regression line represents an average sensitivity value for a given element under the chosen working conditions (Table S3).

The intercept values were closer to zero (zero being the theoretically ideal intercept) for the UWLR regression (~ -0.113 to $+0.104$; Table 1) as compared to the OLR (~ -47.8 to $+12.3$; Table 1). The sensitivity values represented by the slopes of the regression lines (Table 1) were generally similar for both models (~ 2.69 – $46.17 \text{ kc}\cdot\text{s}^{-1}\cdot\%^{-1}$ for the OLR and ~ 2.78 – $59.69 \text{ kc}\cdot\text{s}^{-1}\cdot\%^{-1}$ for the UWLR). The sensitivity actually depends on the measuring conditions (Table S3), which were the same for both models.

For the matrix-corrected intensity-concentration (IntCorr-Conc) regressions, the parameters are listed in Table 2. All intercepts for the UWLR model, without exception, were closer to zero as compared to the OLR model. This confirms the superiority of the UWLR model.

7. Application to Rock Matrices

The calibrations achieved in this work (Table S4) were applied to the analysis of four GRMs (attapulgitic or Fuller’s earth clay ATT1; bentonite clay CSB1; granite GH; and tonalite TLM1) taken as “unknown” samples. These GRMs, having similarly complex matrices as the calibration samples, were not included in the calibrations because their mean values were available only from early description or compilations (for ATT1 and CSB1 [76]; for GH [77]; and for TLM1 [78]). We were unsuccessful in complementing these “old” concentration values with newer ones for these GRMs. Therefore, these GRMs were used as unknown samples. They were analysed in exactly the same manner as the calibration samples.

All calculations for the unknown samples were done outside the instrumental software. The drift-corrected net intensities and the corresponding uncertainties were processed from the first set of two regression equations (Int-Conc OLR and UWLR models; Table S4) to obtain provisional concentration and uncertainty values. The

TABLE 1: Instrumental sensitivities (x - y : concentration-matrix-corrected intensity (Conc-IntCorr) regression model; mean and 99% uncertainty) for major elements.

Element	Regression		Number of data pairs (calibrators)	Regression equation parameters						Quality of regression equation	
	Variables x - y	Model		Intercept (b)	se_b	u_b	Slope (m)	se_m	u_m	r	R^2
SiO ₂	Conc-IntCorr	OLR	60	-47.8441	13.1332	34.9772	6.3592	0.2401	0.6395	0.96105	0.92362
	Conc-IntCorr	UWLR	60	0.1035	5.3870	14.3471	5.3476	0.0985	0.2623	0.99157	0.98321
TiO ₂	Conc-IntCorr	OLR	60	-0.2298	0.3000	0.7991	13.2255	0.2889	0.7694	0.98644	0.97307
	Conc-IntCorr	UWLR	60	-0.0188	0.2637	0.7024	12.3181	0.2539	0.6763	0.99113	0.98235
Al ₂ O ₃	Conc-IntCorr	OLR	60	8.1398	4.1673	11.0987	6.1191	0.2269	0.6043	0.96237	0.92615
	Conc-IntCorr	UWLR	60	0.0594	3.0552	8.1369	7.0715	0.1663	0.4430	0.98557	0.97135
Fe ₂ O ₃ ^t	Conc-IntCorr	OLR	60	12.2628	2.4739	6.5886	4.3943	0.1337	0.3561	0.97419	0.94904
	Conc-IntCorr	UWLR	60	0.0715	2.7723	7.3835	5.4403	0.1498	0.3991	0.98112	0.96259
MnO	Conc-IntCorr	OLR	60	2.0367	0.2963	0.7891	46.1686	0.6958	1.8531	0.99348	0.98700
	Conc-IntCorr	UWLR	60	0.0147	0.7535	2.0069	59.6875	1.7695	4.7127	0.99160	0.98328
MgO	Conc-IntCorr	OLR	60	1.1050	1.0231	2.7246	3.6928	0.0901	0.2400	0.98317	0.96662
	Conc-IntCorr	UWLR	60	0.0108	0.9641	2.5676	3.9863	0.0849	0.2262	0.98745	0.97505
CaO	Conc-IntCorr	OLR	60	-14.3055	3.5661	9.4974	15.4215	0.3371	0.8977	0.98643	0.97304
	Conc-IntCorr	UWLR	60	-0.0121	5.6252	14.9814	11.5666	0.5317	1.4160	0.98657	0.97332
Na ₂ O	Conc-IntCorr	OLR	60	0.0166	0.1553	0.4137	2.6888	0.0571	0.1521	0.98716	0.97449
	Conc-IntCorr	UWLR	60	-0.0111	0.1154	0.3073	2.7843	0.0424	0.1130	0.99344	0.98692
K ₂ O	Conc-IntCorr	OLR	60	1.2200	0.5896	1.5702	13.8296	0.1431	0.3811	0.99691	0.99383
	Conc-IntCorr	UWLR	60	-0.1130	0.5491	1.4623	14.4829	0.1333	0.3549	0.99793	0.99587
P ₂ O ₅	Conc-IntCorr	OLR	60	-1.7376	0.5433	1.4469	39.1055	1.2932	3.4442	0.96972	0.94035
	Conc-IntCorr	UWLR	60	0.0626	0.7039	1.8747	29.4822	1.6756	4.4624	0.96926	0.93946

b , intercept; se , standard error; u , uncertainty at 99%; m , slope; r , linear correlation coefficient; R^2 , squared linear correlation coefficient.

provisional concentrations were then used to obtain matrix corrections for each sample. The method was iteratively applied with the newer concentrations to obtain the final calculated concentration values (Table 3). These calculated concentration values were used to compute the final mean concentrations (\bar{x}) and 99% uncertainties of the mean (u_{99}) for each sample from the second sets of regression equations (ConcCalc-Conc, OLR and UWLR models; Table S4). The loss on ignition (LOI) was required to optimise the final results.

The results are listed in Table 3 and compared with the literature compilations [75–77]. On the other hand, because 99% uncertainties were not reported in the original compilations, they were computed for the comparison from the standard deviation, number of determinations, and appropriate two-sided t values at 99% confidence level [2, 18].

Firstly, although the mean concentration values determined by the OLR and UWLR models showed a general agreement, the 99% uncertainty values (u_{99} ; Table 3) were generally lower for the UWLR models, which clearly

TABLE 2: Regression (x - y): matrix-corrected intensity-concentration (IntCorr-Conc) parameters.

Element	Regression		Number of data pairs (calibrators)	Regression equation parameters						Quality of regression equation	
	Variables x - y	Model		Intercept (b)	se_b	u_b	Slope (m)	se_m	u_m	r	R^2
SiO ₂	IntCorr-Conc	OLR	60	10.8541	1.6762	4.4641	0.1452	0.0055	0.0146	0.96105	0.92362
	IntCorr-Conc	UWLR	60	-0.0148	0.9765	2.6006	0.1848	0.0032	0.0085	0.99157	0.98321
TiO ₂	IntCorr-Conc	OLR	60	0.0344	0.0220	0.0587	0.0736	0.0016	0.0043	0.98644	0.97307
	IntCorr-Conc	UWLR	60	0.0020	0.0204	0.0542	0.0801	0.0015	0.0040	0.99114	0.98235
Al ₂ O ₃	IntCorr-Conc	OLR	60	-0.1675	0.6763	1.8011	0.1514	0.0056	0.0150	0.96237	0.92615
	IntCorr-Conc	UWLR	60	-0.0070	0.4419	1.1769	0.1397	0.0037	0.0098	0.98557	0.97135
Fe ₂ O ₃ ^t	IntCorr-Conc	OLR	60	-2.0576	0.5960	1.5873	0.2160	0.0066	0.0175	0.97419	0.94904
	IntCorr-Conc	UWLR	60	-0.0092	0.5458	1.4536	0.1772	0.0060	0.0160	0.98112	0.96259
MnO	IntCorr-Conc	OLR	60	-0.0412	0.0067	0.0178	0.0214	0.0003	0.0009	0.99348	0.98700
	IntCorr-Conc	UWLR	60	0.0001	0.0134	0.0356	0.0165	0.0007	0.0017	0.99160	0.98328
MgO	IntCorr-Conc	OLR	60	-0.0977	0.2748	0.7319	0.2618	0.0064	0.0170	0.98317	0.96662
	IntCorr-Conc	UWLR	60	-0.0014	0.2692	0.7168	0.2376	0.0063	0.0167	0.98745	0.97505
CaO	IntCorr-Conc	OLR	60	1.0561	0.2173	0.5789	0.0631	0.0014	0.0037	0.98643	0.97304
	IntCorr-Conc	UWLR	60	0.0064	0.4616	1.2293	0.0853	0.0029	0.0078	0.98657	0.97331
Na ₂ O	IntCorr-Conc	OLR	60	0.0447	0.0567	0.1511	0.3624	0.0077	0.0205	0.98716	0.97449
	IntCorr-Conc	UWLR	60	0.0054	0.0421	0.1122	0.3558	0.0057	0.0152	0.99345	0.98693
K ₂ O	IntCorr-Conc	OLR	60	-0.0720	0.0430	0.1146	0.0719	0.0007	0.0020	0.99691	0.99383
	IntCorr-Conc	UWLR	60	0.0082	0.0391	0.1042	0.0687	0.0007	0.0018	0.99793	0.99587
P ₂ O ₅	IntCorr-Conc	OLR	60	0.0537	0.0128	0.0341	0.0241	0.0008	0.0021	0.96972	0.94035
	IntCorr-Conc	UWLR	60	0.0010	0.0202	0.0538	0.0320	0.0013	0.0033	0.96929	0.93952

b , intercept; se , standard error; u , uncertainty at 99%; m , slope; r , linear correlation coefficient; R^2 , squared linear correlation coefficient.

indicates that this model should be used routinely, instead of the conventional OLR model. Secondly, there is also a general agreement among all mean values, especially for granite GH and tonalite TLM1. The two clay samples (ATT1 and CSB1) showed some differences with the preliminary values obtained by the originators of these GRMs [75]. These values for comparison were obtained in only one laboratory. The errors (uncertainties) reported in the literature were underestimated, because they did not include those resulting from the calibrations. Furthermore, the accuracy data of the originator's laboratory were not reported [75], such as the

results for established GRMs and their comparison to other laboratories.

8. Computer Program XRFCalcUnknown

An online computer program JSpectrom_XRFCalcUnknown will be available at our server <https://tloc.ier.unam.mx> for use for unknown samples, which will guide other users to achieve the UWLR calibration outside of the instrumental software and its routine application to unknown samples. This program incorporates the iteration process to achieve reliable

TABLE 3: Application of the XRF calibrations for the determination of major elements (mean concentration in % m/m, mass/mass unit expressed in percentage, habitually called wt.% and 99% uncertainty u_{99} or simply u) in four rock samples (data reported as rounded value following flexible criteria proposed by [18]) and their comparison with the literature data.

Rock	Method (ref.)	SiO ₂	TiO ₂	Al ₂ O ₃	Fe ₂ O ₃ ¹	MnO	MgO	CaO	Na ₂ O	K ₂ O	P ₂ O ₅										
		\bar{x}	u_{99}	\bar{x}	u_{99}	\bar{x}	u_{99}	\bar{x}	u_{99}	\bar{x}	u_{99}										
ATT1	OLR	61.7	6.1	0.4961	0.0173	10.00	0.49	4.246	0.292	0.02394	0.00188	5.373	0.377	2.139	0.086	0.083	0.094	0.7777	0.0378	0.8074	0.0134
	UWLR	62.30	2.23	0.4951	0.0143	9.666	0.321	4.402	0.235	0.02416	0.00269	6.17	0.52	2.144	0.076	0.058	0.065	0.7781	0.0366	0.7828	0.0125
	[76]	59.6	0.8	0.49	0.015	9.50	0.29	3.31	0.06	0.025	0.018	9.14	0.06	1.87	0.03	0.1	0.03	0.86	0.06	0.76	0.01
CSBI	OLR	52.7	5.5	0.1448	0.0157	24.60	0.71	3.493	0.289	0.01408	0.00188	1.523	0.270	1.117	0.085	2.126	0.119	0.4169	0.0374	0.0383	0.0060
	UWLR	52.76	2.04	0.1343	0.0130	24.40	0.48	3.564	0.233	0.01398	0.00268	1.604	0.398	1.105	0.075	2.123	0.083	0.4337	0.0361	0.0357	0.0057
	[76]	55.3	0.8	0.16	0.03	21.75	0.18	3.88	0.03	0.020	0.003	1.94	0.08	1.28	0.06	2.07	0.20	0.54	0.03	—	—
GH	OLR	73.3	6.5	0.0968	0.0157	13.98	0.53	1.336	0.285	0.05084	0.00190	0.074	0.262	0.843	0.084	3.618	0.151	5.302	0.059	0.0119	0.0059
	UWLR	73.62	2.38	0.0861	0.0129	13.714	0.349	1.466	0.229	0.05064	0.00271	0.038	0.390	0.828	0.074	3.625	0.105	5.216	0.057	0.0091	0.0057
	[77]	75.8	0.39	0.08	0.07	12.50	0.16	1.34	0.2	0.05	0.009	0.03	0.18	0.69	0.13	3.85	0.11	4.76	0.05	0.08	0.03
TLM1	OLR	57.7	5.7	0.8497	0.0200	17.38	0.58	7.637	0.306	0.11614	0.00199	3.972	0.319	6.807	0.100	2.675	0.130	1.7585	0.0403	0.1378	0.0063
	UWLR	57.67	2.11	0.8402	0.0165	17.046	0.386	7.682	0.247	0.11592	0.00281	4.281	0.447	6.821	0.088	2.667	0.090	1.742	0.0390	0.1336	0.0060
	[78]	58.85	0.35	0.820	0.013	17.48	0.25	7.67	0.15	0.105	0.013	3.3	0.08	6.67	0.08	2.98	0.07	1.67	0.08	0.15	0.013

concentrations as demonstrated in this work. Example input data files and a ReadMe document are provided to facilitate the application of JSpectrum_XRFCalcUnknown. One important aspect of the program is that for a sample to be identified as an “unknown” sample, the value of LOI (loss on ignition in percent) should be input in the first sheet of the measured intensity file.

A novel aspect of the present work is that total 99% uncertainty can be calculated for individual datum in a given sample (treated as unknown; Table 3). This innovation if put into practice can entirely change the geochemical literature, and in fact make geochemistry a more quantitative science. Further, if an appropriate GRM is analysed as unknown and the analytical data (both mean and total uncertainty) are reported along with the field samples, the data accuracy can be statistically judged from such reports.

9. Conclusions

The XRF spectrometer calibrated under both the OLR and UWLR models clearly showed that the UWLR provides more reliable results (lower uncertainty estimates) than the OLR model commonly practiced for most XRF instruments. The sensitivity and LOD values presented for both models also supported the use of the UWLR model. The UWLR model should therefore be used routinely in such calibrations. The use of a large number of well-characterized GRMs is also recommended for this purpose as illustrated in the present work. The application of our procedure was well documented for the analysis of similarly complex rock matrices. The reporting of total uncertainty values for individual datum is highly recommended for all future geochemical research. This work for the XRF shows that such a practice is easy to achieve in any other analytical calibration procedures. As the major conclusion, we can confirm that the statistically coherent WLR model was shown to perform better than the frequently used conventional statistically incoherent OLR model.

Data Availability

The list of all compiled references (Table S1) will be available at <http://tlaloc.ier.unam.mx>. These references are not included with the manuscript because they are too many (~480). Similarly, as stated, the online program JSpectrum_XRFCalcUnknown will also be added onto this web portal <http://tlaloc.ier.unam.mx>. This program needs to be available online for future use; it cannot be submitted to the journal.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

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Supplementary Materials

Five tables (Tables S1–S5) are provided. (*Supplementary Materials*)

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CAPITULO 5.-

IMPLICACIONES PETROGENÉTICAS Y TECTÓNICAS DE ROCAS VOLCÁNICAS DEL OLIGOCENO-MIOCENO DEL COMPLEJO VOLCÁNICO DE LA SIERRA DE SAN MIGUELITO, MÉXICO CENTRAL

5.1 Presentación del trabajo de investigación publicado

El presente artículo de investigación ha sido publicado en *Journal of South American Earth Sciences*, en el cual el autor de esta tesis participó activamente. Me encuentro como primer autor y el Dr. Sanjeet K. Verma, Dr. Surendra P. Verma, Dr. Fernando Velasco Tapia y el Dr. José Ramón Torres Hernández son los co-autores del presente artículo de investigación.

El presente trabajo publicado ha contribuido con nuevas descripciones de campo del Complejo Volcánico de la Sierra de San Miguelito (CVSSM), junto con la recolección de nuevas muestras de roca en diferentes localidades dentro del CVSSM, tomando en cuenta trabajos previos de la zona. A su vez, se reportan nuevos datos de concentraciones de elementos mayores y elementos traza para las unidades volcánicas del CVSSM. Una parte en destacar en este estudio, es la innovación en el reporte de las concentraciones de elementos mayores, la cual consiste en estimar el 99% de incertidumbre a partir de la calibración basada en el modelo de regresión lineal de mínimos cuadrados ponderado, como también, en la preparación y medición de *pellets* duplicados, los cuales se midieron 6 veces en una secuencia aleatoria.

De igual forma, en el presente trabajo se reporta los primeros modelos geoquímicos cuantitativos para las rocas volcánicas del CVSSM, tanto para procesos de fusión parcial cortical en las rocas volcánicas de composición ácida, como también, modelos cuantitativos de procesos de mezcla de magmas entre las rocas volcánicas de composición básica y ácida. Los presentes modelos sirvieron para brindar una mejor comprensión en la petrogenesis y la implicación tectónica de las rocas volcánicas del CVSSM.

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Petrogenetic and tectonic implications of Oligocene–Miocene volcanic rocks from the Sierra de San Miguelito complex, central Mexico

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ABSTRACT

Paleogene–Neogene volcanic complexes are commonly found in the southern part of the Mesa Central province (MC), Mexico. The Sierra de San Miguelito Complex (SSMC) is located in the southeastern part of the MC and consists of two main phases of Oligocene and Miocene units as follows: (1) basaltic group of porphyritic texture of basalt/trachybasalt compositions; and (2) rhyolite-ignimbrite group of porphyritic texture of trachydacite/trachyte compositions. Chondrite-normalized rare-earth element patterns for the basaltic group display enrichment in light rare earth elements (LREE) and large ion lithophile elements (LILE), whereas the rhyolite-ignimbrite group shows enrichment in LREE and high field strength elements (HFSE). The Eu anomalies within the SSMC become progressively more negative from the basaltic group to the rhyolite-ignimbrite group ($\text{Eu}/\text{Eu}^* = 1.13$ to 0.02) suggest upper crust (crustal contamination) participates during magma generation. Trace element modelling reveals that the basaltic group evolved through a mixing process between mafic and felsic end-members. In contrast, chondrite-normalized and multi-element models reveal that the rhyolite-ignimbrite group evolved by partial melting of the upper continental crust. New multi-dimensional discrimination diagrams along with the Nb–Ta anomalies reveal that the SSMC volcanic rocks were generated in an extensional environment.

1. Introduction

The Mesa Central province (MC) is an elevated plateau located in the central-northern Mexico and is bounded to the N and E by the Sierra Madre Oriental, and toward the W by the Sierra Madre Occidental (Fig. 1a; Nieto-Samaniego et al., 2007). The major fault systems active during the Cenozoic and surrounding as well as within the MC are as follows (Fig. 1b): (a) the El Bajío system at the south; (b) the Taxco-San Miguel de Allende at the west; and (c) the San Luis-Tepehuanes that extends along the MC province. The San Luis-Tepehuanes fault system characterized by a NW-SE lineaments separates the MC into two main regions: (i) northern region, which is characterized by advance stage of erosion with alluvial–lacustrine basins and shows low magmatic evidence from the Oligocene and Quaternary ages; and (ii) southern region

that is mainly covered by Paleogene-Neogene volcanic rocks with normal faults (Nieto-Samaniego et al., 1996; 1999, 2007).

At the southern region of the MC, the main volcanic activity is concentrated (Fig. 1b; see the box marked as c) and amplified in Fig. 1c). Previous studies at the southern region of the MC showed that partial melting and fractional crystallization processes of the lower crust in an extensional setting generated felsic magmas in the region (Orozco-Esquivel et al., 2002; Aguilón-Robles et al., 2014; Sieck et al., 2019). Based on the petrographic evidence, Rodríguez-Ríos and Torres-Aguilera (2009) proposed that mafic volcanism exposed in this area is derived from partial melting of the upper mantle and mixing with the overlying crust. In addition, Tristán-González et al. (2009) and Aguilón-Robles et al. (2014) inferred that associated mafic and intermediate magmas were derived in a post-subduction setting from partial melting of the

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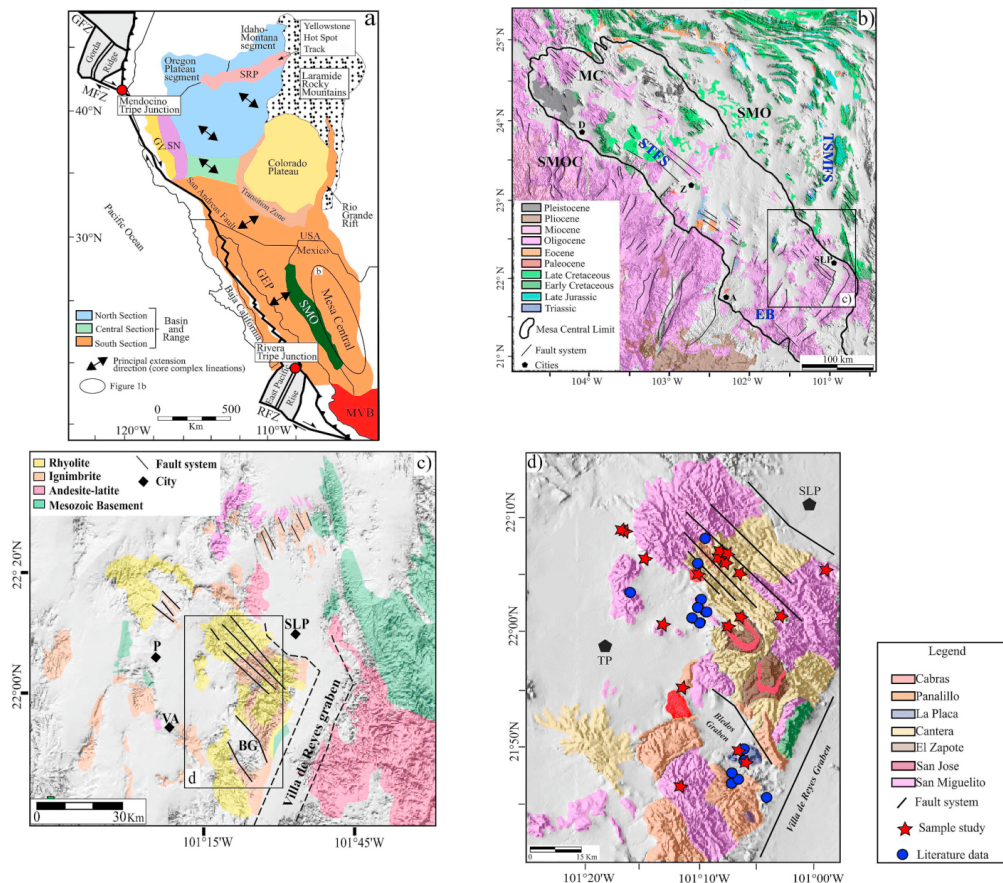


Fig. 1. (a) Present-day regional tectonic map of western North America and the Basin and Range province with subdivisions of the northern, central and southern Basin and Range from Jones et al. (1992) and Sonder and Jones (1999). Principal extensional directions established from core complex lineation in each region are simplified and modified from Dickinson (2002) and Cosca et al. (2014); (b) Simplified geological map of the Mesa Central province (modified after Nieto-Samaniego et al., 2007); (c) Geological map of the San Luis Potosí volcanic field (SLPVF; modified after Tristán-González, 1986); (d) Geological map of the Sierra de San Miguelito complex (SSMC), showing the space distribution of the geological units. Abbreviations: GEP – Gulf Extensional Province; SN – Sierra Nevada; RFZ – Rivera Fracture Zone; GV – Great Valley; SRP – Snake River Plain; GFZ – Gorda Fracture Zone; MFZ – Mendocino Fracture Zone; MVB – Mexican Volcanic Belt; STFS – San Luis-Tepahuanes Fault System; TSMFS – Taxco-San Miguel de Allende Fault System; EB – El Bajío Fault System; MC – Mesa Central province; SMO – Sierra Madre Oriental province; SMOC – Sierra Madre Occidental province; the cities are D – Durango; Z – Zacatecas; A – Aguascalientes; TP – Tepetate; SLP – San Luis Potosí; USA – United States of America.

upper mantle. Nevertheless, no detailed geochemical studies have been performed on volcanic rocks from this area. Therefore, in this work, we present a detailed study of field-relationships and geochemistry of volcanic rocks from the Sierra de San Miguelito Complex (SSMC; Fig. 1d) and discuss a quantitative geochemical model to provide a better understanding of the petrogenesis and tectonic implications of the magmatic activity within the southern region of the MC.

2. Geological setting

The San Luis Potosí Volcanic Field (SLPVF; Fig. 1c) is located in the southern region of the MC and includes felsic – mafic volcanism, ranging in age from Eocene to Quaternary. The main volcanic event of the

SLPVF occurred during late Oligocene and is characterized by high-silica rhyolites and large volume ignimbrites. The youngest volcanic event at the SLPVF was characterized by less voluminous mafic to intermediate lavas (Aguillón-Robles et al., 2014). The SLPVF has been divided into six main volcanic complexes as follows: (i) Ahuualulco; (ii) Villa Hidalgo; (iii) La Repartición; (iv) Pinos; (v) Santa María; and (vi) Sierra de San Miguelito (Torres-Hernández et al., 2006; Rodríguez-Ríos et al., 2007; Tristán-González et al., 2009; Aguillón-Robles et al., 2014; López-Loera, 2014).

The Sierra de San Miguelito complex (SSMC; Fig. 1d) consists mainly of Oligocene – Early Miocene rocks of silicic lavas associated to silicic domes, ignimbrite packages, and isolated basaltic volcanism. The volcanic activity of the SSMC was emplaced through fault systems that

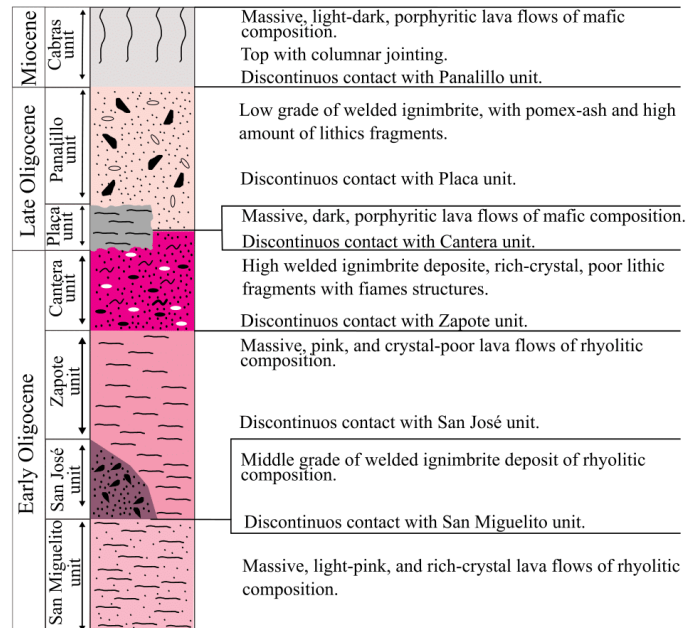


Fig. 2. Schematic stratigraphic column for the volcanism in the SSMC.

exhibit a “domino style geometry” with strike direction of 300°–340° dip to the SW and tilted to NW systematically with dip angles between 12° and 31° (Xu et al., 2004, 2013). The stratigraphy of the SSMC covers the period of Oligocene to Early Miocene and most of the lithology in this complex corresponds to Oligocene. These lithological units could be distributed in three groups as follows: (a) basaltic group (La Placa and Cabras unit); (b) rhyolite group (San Miguelito and El Zapote unit); (c) ignimbrite group (San José, Cantera and Panalillo unit) (Fig. 2). The early Miocene epoch is characterized by the Cabras unit (Figs. 1d and 2).

2.1. Basaltic group

The basaltic group is constituted by the La Placa and Cabras units (Fig. 2). Basaltic group displays outcrop in isolated regions of the study area (Fig. 1d). Previous studies (Tristán-González et al., 2009; Aguillón-Robles et al., 2014) have dated the volcanism of the La Placa unit and reported whole-rock K–Ar age of 29 to 26 Ma. The La Placa unit is characterized by massive lava flows that are intercalated with the ignimbrite group (Panalillo and Cantera units). The La Placa unit shows a thickness range from 10 to 15 m and is mainly characterized by scarce phenocrysts and a vitreous matrix. On the other hand, Tristán-González et al. (2009) and Aguillón-Robles et al. (2014) proposed whole-rock K–Ar age of 22 to 21 Ma for the Cabras unit. This unit finds in isolated outcrops of lava flows having a thickness of 25 m and is characterized by aphanitic-porphyritic texture, scarce phenocrysts and rare presence of vesicles in a vitreous matrix (Labarthe-Hernández and Jiménez-López, 1992; Tristán-González et al., 2009; Gaytán-Martínez et al., 2017).

2.2. Rhyolite group

The rhyolite group consists of the San Miguelito and El Zapote units (Fig. 2). Previous studies (Labarthe-Hernández and Jiménez-López,

1992; Aguillón-Robles et al., 1994, 2014; Tristán-González et al., 2009) reported whole-rock K–Ar dates for the San Miguelito and El Zapote units ranging from 33 to 21 Ma (Tristán-González et al., 2009). These units consist of lavas flows associated to exogenous domes, whose last eruptive phase formed a thick covering at the top of the domes, known as breccia and vitreous fragments. The thickness of these units are variable (130–800 m), although it is higher at the center of domes. The rhyolite group includes mainly quartz and sanidine phenocrysts embedded in a devitrified and fluidal matrix (Tristán-González, 1986; Labarthe-Hernández and Jiménez-López, 1992; Aguillón-Robles et al., 1994; Tristán-González et al., 2009).

2.3. Ignimbrite group

The ignimbrite group is formed by the San José, Cantera and Panalillo units (Fig. 2). Based on stratigraphic aspects Tristán-González et al. (2009) proposed an age of 29 Ma for the San José unit. This unit consists of ash and pyroclastic flows, having a middle-grade of welding and a thickness of ~3–15 m. The San José unit appears in contact with the rhyolite group (San Miguelito and El Zapote unit; Labarthe-Hernández and Jiménez-López, 1992; Tristán-González et al., 2009). In contrast, the Cantera unit was dated at about 29 Ma by the K–Ar method (Labarthe-Hernández and Jiménez-López, 1992; Tristán-González et al., 2009; Gaytán-Martínez et al., 2017). The Cantera unit is characterized by pyroclastic flows of pumice and ash with fiamme structures and lithics fragments. It is present in the major part of the study area and overlays the San Miguelito unit. The Cantera unit shows high-grade of welding and presents an average thickness of 2–30 m.

The Panalillo unit displays variable ages of 29 to 26 Ma (Labarthe-Hernández and Jiménez-López, 1992; Torres-Hernández et al., 2006; Tristán-González et al., 2009), is constituted by pumice–ash pyroclastic flows with a lesser grade of welding, and shows an average

Table 1
Petrographic information of the Sierra de San Miguelito complex volcanic rocks.

Sample	Group	Unit	Lat (N)°	Long(W)°	Texture	Phenocrysts						Groundmass		Disequilibrium type	
						Ol	Plg	Opx	Cpx	Qz	FK	Op	Qz + R	Pl-N-S	
SLP-01	Basaltic	Cabras	21°55'17"	101°11'50"	P	4	10	4	6			26	50		*
SLP-02	Basaltic	La Placa	21°46'45"	101°03'11"	P	6	68		20			5	1		
SLP-03	Basaltic	La Placa	21°49'47"	101°04'42"	P	3	19	7	3			1	67	*	*
SLP-06	Rhyolite-Ignimbrite	San Miguelito	22°05'23"	101°13'06"	P					3	1	41	15	30	10
SLP1712	Rhyolite-Ignimbrite	San Miguelito	22°00'03"	101°10'51"	P					3	1	46	10	36	4
SLP-07	Rhyolite-Ignimbrite	El Zapote	22°04'12"	101°04'45"	P					44	4	4	26	15	7
SLP1715	Rhyolite-Ignimbrite	Panallillo	21°46'47"	101°02'13"	P	2	10	18	35	15	15	1	4		
SLP-04	Rhyolite-Ignimbrite	Panallillo	21°48'45"	101°04'11"	P	2	2	9	11	32	14	4	26		
SLP1719	Rhyolite-Ignimbrite	Panallillo	21°49'02"	101°06'08"	P					4		16	22	9	49
SLP1723	Rhyolite-Ignimbrite	Panallillo	22°06'06"	101°06'35"	P					43	1	38	15	3	
SLP1724	Rhyolite-Ignimbrite	Cantera	22°05'26"	101°06'34"	P					3	6	24	20	6	41
SLP-05	Rhyolite-Ignimbrite	San José	22°01'20"	101°04'44"	P					4	2	16	21	5	52

Modal data are presented in percentage (%). Texture: P = porphyritic. Minerals: Ol: olivine; Plg: plagioclase; Opx: orthopyroxene; Cpx: clinopyroxene; Qz: quartz; FK: potassium feldspar; Amp: amphibole; Op: opaque. Disequilibrium type: Qz + R = quartz with reaction rim, Pl-N-S = Plagioclase with normal and sieved texture, * = Presence of Disequilibrium type.

thickness of 20–30 m. This unit shows a high amount of lithics fragments and overlies the Cantera unit, with minor intercalation with the San Miguelito unit in some part of the study area (Tristán-González, 1986; Labarthe-Hernández and Jiménez-López, 1992; Tristán-González et al., 2009; Gaytán-Martínez et al., 2017).

3. Sampling and petrography

For this study, we collected representative samples of the main basaltic and rhyolite-ignimbrite groups from the SSMC. The sampling coordinates are presented in Tables 1 and 2. Modal compositions were determined by point counting on thin section using a Leica petrographic microscope and a PELCON Automatic point counter. In order to obtain a representative mode, approximately 1000 points per sample were counted. Modal composition data are reported in Table 1.

3.1. Basaltic group

This group is characterized by porphyritic textures with limited presence of vesicles and a vitreous matrix (Table 1). The main mineral assemblage (Fig. 3a and b) from this unit consists of phenocrysts of euhedral to subhedral plagioclase (10–68%) generally showing Carlsbad twins and normal zoning, subhedral phenocrysts of clinopyroxene (6–20%), subhedral phenocrysts of orthopyroxene (4–7%), and a lesser amount of subhedral to xenomorphic microcrystals of olivine (< 6%). Furthermore, the basaltic group samples display the following characteristics: (i) diverse disequilibrium textures that include normal and sieved plagioclases in the same sample; (ii) rounded and embayed crystals with reaction rims; and (iii) plagioclase with complex mineral zoning, such as reverse and oscillatory zoning or normally and reversely zoned crystals in the same sample (Fig. 3a and b).

3.2. Rhyolite group

The rhyolite group consists of massive flows, which show porphyritic textures, except in some places, where a fluidal texture can be observed. The principal minerals (Fig. 3c and d) consist of phenocrysts of xenomorphic quartz (27–41%), euhedral to subhedral potassium feldspar (sanidine; 11–15%), euhedral to subhedral plagioclase (44%) with 0.5–2.0 mm of diameter, subhedral orthopyroxene (1–4%) and clinopyroxene (1–4%) of 0.2–0.7 mm of diameter. (Table 1; Fig. 3c and d).

3.3. Ignimbrite group

Fresh ignimbrite samples were collected from the SSMC. This group consists of welded and semi-welded pyroclastic flow deposits and represents porphyritic textures. The main minerals (Fig. 3e–g) are subhedral phenocrysts of quartz (15–38%), potassium feldspar (sanidine; 15–22%), plagioclase (2–43%), orthopyroxene (2–5%), and clinopyroxene (2–7%); olivine is present in less abundance in some samples. All subhedral phenocrysts are embedded in a vitreous matrix (Table 1; Fig. 3e–g).

4. Geochemical analytical procedures

4.1. Major and trace element analysis

Major element for 23 samples were determined by a wavelength dispersive X-ray fluorescence (WD-XRF) spectrometer Rigaku ZSX Primus II at Institute of Scientific and Technological Research of San Luis Potosí, IPICYT (San Luis Potosí, Mexico). The analytical procedures, accuracy, and precision were given by Verma et al. (2018, 2019). The computer program MECUX is available for online use at <http://tlaloc.ier.unam.mx>. The innovation in reporting major elements consists in estimating total 99% uncertainty from the calibration based on uncertainty-weighted least-squares linear regression model and preparation and measurement of duplicate pellets, each of which run 6 times, in a random sequence, in the XRF instrument. Thus, there is 99% confidence that the true value will be within the interval (mean \pm 99% uncertainty) implied from the reported results.

Trace element concentrations were determined in ALS Chemex Laboratories (Vancouver, Canada) using the "ME-MS81" method (for more details visit: www.alsglobal.com/geochemistry). Trace element concentrations were obtained by Inductively Coupled Plasma Mass Spectrometry (ICP-MS) with an analytical precision based on geochemical reference materials SY-2 and OREAS146.

Besides new geochemical data for 23 samples, additional literature data for 31 samples from the SSMC were compiled from Leroy et al. (2002; 11 samples); Orozco-Esquivel et al. (2002; 8 samples); Aguilón-Robles et al. (2014; 4 samples) Torres-Hernández et al. (2014; 8 samples). All computations (anhydrous and iron-oxidation ratio adjustments, norm compositions, and rock classifications) were automatically done from the IgRoCS software (Verma and Rivera-Gómez, 2013), with iron-oxidation ratio adjustments proposed by Middlemost (1989). Rock classification was based on the total alkali – silica diagram (TAS–Le Bas et al., 1986; Fig. 4).

Table 2
Major element geochemistry and CIPW norms of volcanic rocks from the Sierra de San Miguelito Volcanic Complex, San Luis Potosí, Mexico. n = number of analyses of each sample; [Mg# = $100 \times (\text{Mg}^{2+} / (\text{Mg}^{2+} + \text{Fe}^{2+}))$]. CIPW norms are computed on an anhydrous, 100% adjusted basis and $\text{Fe}_2\text{O}_3/\text{FeO}$ ratio after *Middlemost (1989)*, using a program proposed by *Verma and Růvera-Gómez (2013)*.

Sample	SLP-01	SLP-02	SLP-03	SLP-04	SLP-05	SLP-06	SLP-07
Rock unit	Cabras	La Placa	La Placa	La Placa	La Placa	Cantera	Panaillo
Rock (TAS)	Foidite, melaphelinitite	Foidite, melaphelinitite	Potassic trachybasalt	Alkali basalt	Rhyolite	Rhyolite	Rhyolite
Long. (°W)	101°11'50"	101°03'11"	101°05'12"	101°04'42"	101°06'34"	101°02'11"	101°05'43"
Lat. (°N)	21°55'17"	21°46'45"	21°49'47"	21°49'47"	22°05'26"	21°46'46"	21°49'48"
n	12	12	12	12	12	12	12
SiO ₂	39.11 ± 1.50	37.38 ± 1.44	44.76 ± 1.5	46.34 ± 1.54	71.01 ± 1.94	72.73 ± 1.91	71.6 ± 1.94
TiO ₂	3.9991 ± 0.039	1.7453 ± 0.0225	1.7345 ± 0.021	2.1155 ± 0.0256	0.1137 ± 0.011	0.196 ± 0.011	0.187 ± 0.013
Al ₂ O ₃	15.76 ± 0.314	18.556 ± 0.305	17.139 ± 0.297	15.616 ± 0.302	11.245 ± 0.304	11.388 ± 0.301	12.726 ± 0.319
Fe ₂ O ₃	14.898 ± 0.131	14.657 ± 0.12	10.977 ± 0.105	11.721 ± 0.111	1.608 ± 0.078	1.613 ± 0.078	1.611 ± 0.079
MnO	0.38551 ± 0.003	0.34943 ± 0.003	0.34366 ± 0.003	0.34362 ± 0.003	0.01435 ± 0.002	0.01786 ± 0.002	0.02757 ± 0.0023
MgO	4.513 ± 0.255	6.025 ± 0.251	5.177 ± 0.249	4.488 ± 0.249	0.841 ± 0.263	0.845 ± 0.236	0.722 ± 0.236
CaO	14.355 ± 0.175	15.963 ± 0.159	12.486 ± 0.143	11.74 ± 0.148	2.134 ± 0.119	2.217 ± 0.119	2.052 ± 0.119
Na ₂ O	2.082 ± 0.077	2.333 ± 0.073	2.642 ± 0.075	2.142 ± 0.074	0.482 ± 0.069	0.613 ± 0.069	0.45 ± 0.069
K ₂ O	1.984 ± 0.048	1.403 ± 0.045	2.406 ± 0.048	2.252 ± 0.048	5.366 ± 0.065	4.012 ± 0.056	4.842 ± 0.061
P ₂ O ₅	2.4266 ± 0.043	0.8962 ± 0.013	0.6307 ± 0.010	0.7638 ± 0.013	0.0442 ± 0.007	0.0592 ± 0.007	0.0501 ± 0.0067
LOI	0.496	0.692	1.703	2.481	7.146	0.824	5.818
Sum	100.005	99.999	99.998	100.005	100.004	100.001	100.003
Q					44.63	50.71	47.33
Or	11.93		14.59	13.81	34.19	25.34	30.42
Ab	0.44		5.67	16.65	4.40	40.43	4.05
An	28.28		28.53	27.28	11.10	11.34	10.47
Ne	9.48		9.36	1.15			
C					0.94	2.29	3.33
Di-Mg	11.58		15.01	11.91			
Di-Fe	11.76		10.28	11.22			
Hy-Mg					2.26	2.25	0.85
Hy-Fe					1.36	1.21	1.38
Mt	3.35		3.47	2.68	0.78	0.78	0.77
Il	7.73		3.38	4.16	0.23	0.40	0.21
Ap	5.72		1.50	1.83	0.11	0.15	0.12
Mg#	41.46		54.26	47.23	60.04	28.99	56.28
FeO/MgO	2.97		1.91	2.35	1.72	6.33	2.01

Sample	SLP17-08	SLP17-09	SLP-04	SLP17-17	SLP17-25	SLP-05	SLP17-04
Rock unit	Panaillo	Panaillo	Panaillo	Panaillo	Panaillo	San José	San Miguelito
Rock (TAS)	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Rhyolite
Long. (°W)	101°09'24"	101°09'2"	101°04'11"	101°05'11"	101°05'45"	101°04'44"	101°12'33"
Lat. (°N)	22°05'10"	22°05'07"	21°48'45"	21°48'46"	22°05'14"	22°01'20"	22°00'48"
n	12	12	12	12	12	12	12
SiO ₂	70.98 ± 1.76	72.69 ± 1.89	78.88 ± 2.01	69.4 ± 1.81	71.98 ± 1.94	68.68 ± 1.63	74.32 ± 2.16
TiO ₂	0.2157 ± 0.0113	0.2019 ± 0.0114	0.0796 ± 0.0111	0.2344 ± 0.0115	0.1484 ± 0.0112	0.1627 ± 0.0112	0.0595 ± 0.0111
Al ₂ O ₃	14.692 ± 0.298	13.978 ± 0.309	10.895 ± 0.295	15.567 ± 0.304	10.129 ± 0.297	16.971 ± 0.291	14.004 ± 0.348

(continued on next page)

Table 2 (continued)

Sample	SLP17-08	SLP17-09	SLP-04	SLP17-17	SLP17-25	SLP17-05	CR17-04
Rock unit	Panahillo	Panahillo	Panahillo	Panahillo	Panahillo	San José	San Miguelito
Rock (TAS)	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Rhyolite
Long. (W)	101°09'24"	101°09'22"	101°04'11"	101°05'11"	101°05'45"	101°04'44"	101°12'33"
Lat. (N)	22°05'10"	22°05'07"	21°48'45"	21°48'46"	22°05'14"	22°01'20"	22°00'48"
n	12	12	12	12	12	12	12
Fe ₂ O ₃	2.627 ± 0.079	2.401 ± 0.079	1.096 ± 0.078	2.484 ± 0.079	1.619 ± 0.078	2.15 ± 0.078	1.407 ± 0.079
MnO	0.02747 ± 0.00233	0.03816 ± 0.00235	0.03232 ± 0.0023	0.11047 ± 0.0023	0.0173 ± 0.0023	0.04593 ± 0.0023	0.01724 ± 0.0023
MgO	0.076 ± 0.236	0.045 ± 0.236	0.155 ± 0.236	0.737 ± 0.236	0.783 ± 0.236	0.096 ± 0.236	0.019 ± 0.236
CaO	0.606 ± 0.118	0.516 ± 0.118	0.924 ± 0.118	2.081 ± 0.119	2.397 ± 0.119	1.075 ± 0.118	0.328 ± 0.118
Na ₂ O	3.189 ± 0.077	2.946 ± 0.079	2.416 ± 0.077	3.378 ± 0.078	0.42 ± 0.069	2.822 ± 0.073	3.467 ± 0.106
K ₂ O	6.61 ± 0.064	6.458 ± 0.066	4.836 ± 0.058	4.948 ± 0.058	5.291 ± 0.063	6.647 ± 0.061	5.554 ± 0.073
P ₂ O ₅	0.0786 ± 0.0067	0.0669 ± 0.0067	0.0574 ± 0.0067	0.1068 ± 0.0068	0.0509 ± 0.0067	0.1133 ± 0.0067	0.0342 ± 0.0067
LOI	0.9	0.654	0.627	0.954	7.165	1.239	0.791
Sum	100.001	99.994	99.998	100.000	100.000	100.001	100.000
Q	25.30	29.25	44.17	24.92	46.25	24.25	32.01
Or	39.49	38.48	28.78	29.57	33.72	39.83	33.12
Ab	27.28	25.14	20.59	28.91	3.83	24.21	29.60
An	2.52	2.14	4.24	9.74	10.92	4.66	1.42
Ne	1.39	1.37	0.14	1.14	0.82	3.50	1.79
Di-Mg					0.44		
Di-Fe	0.19	0.11	0.39	1.86	1.72	0.24	0.05
Hy-Mg	2.04	1.87	0.90	2.03	1.08	1.74	1.19
Hy-Fe	1.19	1.09	0.50	1.13	0.79	0.98	0.64
Mt	0.41	0.39	0.15	0.45	0.30	0.31	0.11
Il	0.18	0.16	0.13	0.25	0.13	0.27	0.08
Ap	7.67	5.11	28.89	46.01	58.14	11.37	3.73
Mg#							
FeO/MgO	31.10	48.01	6.36	3.03	1.86	20.15	66.63
Sample	SLP17-03	SLP17-04	SLP17-05	SLP-07	SLP17-30	SLP17-31	SM17-03
Rock unit	San Miguelito	San Miguelito	San Miguelito	El Zapote	Zapote	Zapote	San Miguelito
Rock (TAS)	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Rhyolite
Long. (W)	101°07'00"	101°07'36"	101°07'31"	101°04'45"	101°06'43"	100°56'08"	101°14'07"
Lat. (N)	22°07'02"	22°07'04"	22°07'07"	22°04'12"	22°00'40"	22°05'28"	22°06'23"
n	12	12	12	12	12	12	12
SiO ₂	69.43 ± 1.84	70.85 ± 1.88	68.12 ± 1.64	68.57 ± 1.78	72.82 ± 1.96	72.36 ± 1.77	69.95 ± 2.02
TiO ₂	0.1305 ± 0.0112	0.1299 ± 0.0112	0.1458 ± 0.0112	0.1436 ± 0.0112	0.1302 ± 0.0112	0.1352 ± 0.0112	0.1334 ± 0.0112
Al ₂ O ₃	17.153 ± 0.329	15.643 ± 0.317	16.264 ± 0.289	18.084 ± 0.322	15.101 ± 0.326	14.863 ± 0.295	17.027 ± 0.357
Fe ₂ O ₃	1.668 ± 0.079	1.645 ± 0.078	2.052 ± 0.079	1.771 ± 0.078	1.693 ± 0.079	1.627 ± 0.078	1.456 ± 0.078
MnO	0.03938 ± 0.0023	0.03955 ± 0.0024	0.0575 ± 0.0023	0.04941 ± 0.00235	0.03128 ± 0.00235	0.03287 ± 0.0023	0.02497 ± 0.00235
MgO	0.362 ± 0.236	0.258 ± 0.236	0.194 ± 0.236	0.136 ± 0.236	0.02178 ± 0.236	0.046 ± 0.236	0.313 ± 0.236
CaO	0.751 ± 0.118	0.971 ± 0.118	1.828 ± 0.118	0.76 ± 0.118	0.692 ± 0.118	1.018 ± 0.118	0.699 ± 0.118
Na ₂ O	2.743 ± 0.079	2.393 ± 0.078	2.764 ± 0.073	3.229 ± 0.079	2.938 ± 0.082	3.346 ± 0.081	2.745 ± 0.082
K ₂ O	6.357 ± 0.069	6.519 ± 0.071	7.082 ± 0.064	6.165 ± 0.067	5.812 ± 0.066	5.857 ± 0.068	6.547 ± 0.073

(continued on next page)

Table 2 (continued)

Sample	SLP17-03	SLP17-04	SLP17-05	SIP-06	SIP-07	SLP17-30	SLP17-31	SM17-03
Rock unit	San Miguelito	San Miguelito	San Miguelito	San Miguelito	El Zapote	Zapote	Zapote	San Miguelito
Rock (TAS)	Rhyolite	Rhyolite	Rhyolite	Trachydacite	Rhyolite	Rhyolite	Rhyolite	Rhyolite
Long. (W)	101°07'00"	101°07'36"	101°07'31"	101°13'06"	101°04'45"	101°06'43"	100°58'08"	101°14'07"
Lat. (N)	22°07'02"	22°07'04"	22°07'07"	22°05'23"	22°04'12"	22°00'40"	22°05'28"	22°06'23"
n	12	12	12	12	12	12	12	12
P ₂ O ₅	0.0531 ± 0.0067	0.0507 ± 0.0066	0.0952 ± 0.0067	0.1104 ±	0.0669 ± 0.0067	0.0608 ± 0.0067	0.0748 ± 0.0067	0.0664 ± 0.0068
LOI	1.316	1.496	1.395	0.822	1.047	0.663	0.64	1.036
Sum	100.002	99.995	99.997	100.000	100.003	100.002	99.999	99.997
Q	26.92	29.54	20.55	13.94	24.20	31.72	28.07	26.95
Or	38.11	39.16	42.51	44.33	36.86	34.62	34.88	39.14
Ab	23.55	20.58	23.75	26.63	27.64	25.06	28.53	23.49
An	3.43	4.56	8.58	4.77	3.37	3.06	4.60	3.07
Nc								
C	4.59	3.06	0.97	6.22	4.93	2.89	1.36	4.36
Di+Mg								
Di+Fe								
Hy-Mg	0.91	0.65	0.49	1.13	0.34	0.18	0.12	0.79
Hy+Fe	1.35	1.33	1.70	1.48	1.40	1.33	1.28	1.12
Mt	0.76	0.75	0.94	0.87	0.81	0.77	0.74	0.66
Il	0.25	0.25	0.28	0.37	0.28	0.25	0.26	0.26
Ap	0.12	0.12	0.22	0.26	0.16	0.14	0.17	0.16
Mg#	38.40	31.06	21.36	40.30	18.07	10.75	7.51	38.17
FeO/MgO	4.15	5.74	9.52	3.83	11.72	21.46	31.83	4.19

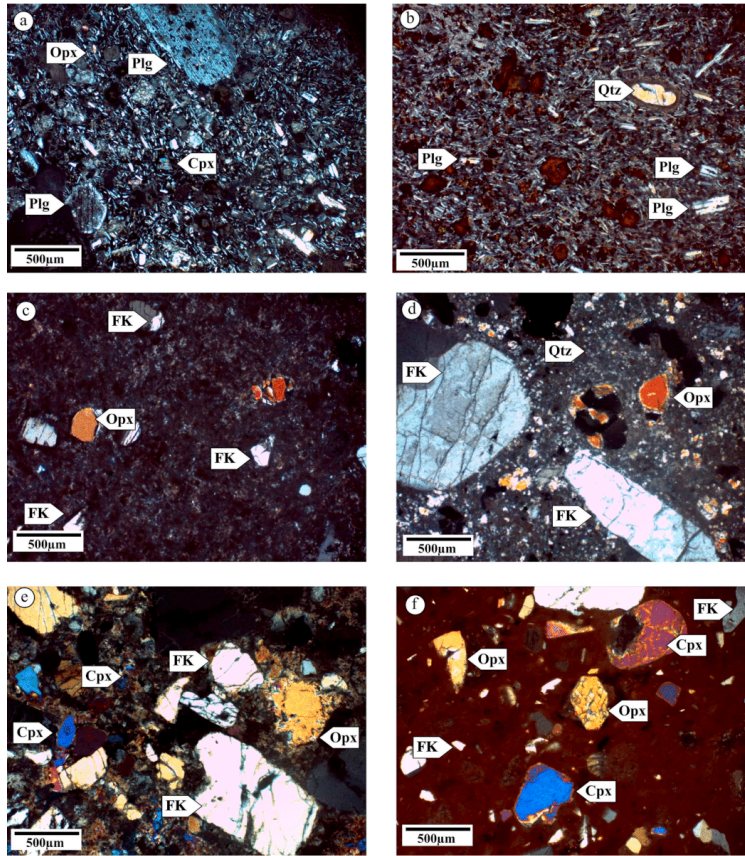


Fig. 3. Photomicrographs of the SSMC volcanic rocks. (a) Phenocrysts of plagioclase and quartz with reaction rims in the Cabras unit; b) Phenocrysts of potassium feldspar, clinopyroxene and orthopyroxene in the Panalillo unit; c) Sieved textures in plagioclase in the Placa unit; d) Phenocrysts of potassium feldspar, orthopyroxene and clinopyroxene in the Cantera unit; e) Phenocrysts of potassium feldspar, orthopyroxene, clinopyroxene in the El Zapote unit; f) Phenocrysts of potassium feldspar, orthopyroxene, clinopyroxene in the San José unit; g) Potassium feldspar phenocrysts and orthopyroxene crystals of porphyritic texture in the San Migueluelo unit. Abbreviations of mineral names are reported in Table 1.

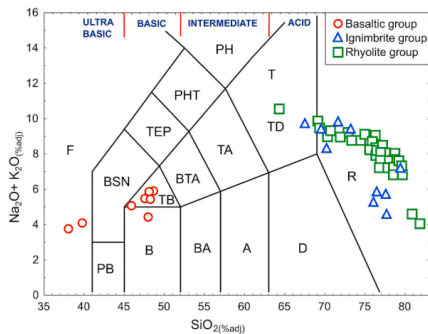


Fig. 4. Total Alkali–Silica classification diagram (TAS) for the volcanic rocks from the SSMC (Le Bas et al., 1986) prepared from the adjusted data by using the IgRoCS computer program (Verma and Rivera-Gómez, 2013).

5. Geochemical results

5.1. Major elements

The basaltic group samples show a foidite to trachybasalt composition (Fig. 4), covering the range of $(\text{SiO}_2)_{\text{adj}} = 38.12\text{--}48.01\%$ m/m, $(\text{Al}_2\text{O}_3)_{\text{adj}}$ values of 16.04–18.92% m/m, $(\text{MgO})_{\text{adj}}$ values are 4.59–6.14% m/m, $(\text{TiO}_2)_{\text{adj}} = 1.78\text{--}4.07\%$ m/m and Mg# (calculated as $100 \times \text{Mg}^{2+}/(\text{Mg}^{2+} + \text{Fe}^{2+})$), ranges from 41.46 to 54.26. Most of the samples of this group are nepheline normative ($n = 4$; Table 2).

In contrast, the rhyolite-ignimbrite group samples display a rhyolitic composition, with the exception of one sample that shows trachydacite composition. The rhyolite-ignimbrite group samples contain 64.36–79.43% m/m $(\text{SiO}_2)_{\text{adj}}$, relatively high $(\text{Al}_2\text{O}_3)_{\text{adj}}$ ranging from 10.92 to 21.26% m/m. $(\text{MgO})_{\text{adj}}$ values are 0.01–0.91% m/m, $(\text{TiO}_2)_{\text{adj}} = 0.06\text{--}0.24\%$ m/m and Mg# ranges from 3.73 to 60.07; most of the samples are hypersthene normative ($n = 19$; Table 2).

5.2. Trace elements

Trace element compositions of the basaltic group samples include Sc = 21–28 ppm ($\mu\text{g}\cdot\text{g}^{-1}$), Cr = 100–240 ppm, Ni = 7–92 ppm, and

Table 3 (continued)

Sample	SLP17-25	SLP-05	CR17-04	SLP17-03	SLP17-04	SLP17-05	SLP-06	SLP-07	SLP17-30	SLP17-31	SM17-03
Rock unit	Panahillo	San José	San Miguelito	San Miguelito	San Miguelito	San Miguelito	San Miguelito	El Zapote	Zapote	Zapote	San Miguelito
Rock (TAS)	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Trachydacite	Rhyolite	Rhyolite	Rhyolite	Rhyolite
Long. (W)	101°05'45"	101°04'44"	101°12'33"	101°07'00"	101°07'36"	101°07'31"	101°13'06"	101°04'45"	101°06'43"	100°58'08"	101°14'07"
Lat. (N)	22°05'14"	22°01'20"	22°00'48"	22°07'02"	22°07'04"	22°07'07"	22°05'23"	22°04'12"	22°00'40"	22°05'28"	22°06'23"
Co	2			2		1		1	1		1
Cr	9.05	10	10	18.1	17.35	20.4	11.9	14.8	13.25	13.5	13.05
Cs	11.75	5.18	5	2	2	2	2	3	2	2	2
Cu	3	5	2	2	2	2	2	2	2	2	2
Ga	15.9	23.1	42.8	24.6	23.2	23.1	22.2	23	21.6	23.1	22.7
Hf	4.6	8.9	4.8	4.8	4.4	4	4.8	5.8	5.6	5.4	5.6
Nb	13.3	23	100.5	26.1	24.5	25.1	20.8	23.3	21.3	25.2	21.6
Ni	1	2	2	1	2	2	2	1	2	2	2
Pb	22	35	62	37	36	33	32	40	33	38	33
Rb	183	336	397	407	390	403	319	368	330	369	315
Sc	5	7	10	10	10	10	7	8	8	9	8
Sr	648	19.9	5.6	16	158.5	117.5	32.8	14.5	18.1	12.9	30.2
Ta	1.2	2.1	9.7	2.5	2.4	2.5	1.9	2.2	2.4	2.4	2
Th	21.9	34	18.25	33.8	31.2	31	25.3	29.2	26.7	37.9	30.8
U	6.15	6.31	6.65	6.78	6.58	5.74	4.23	4.48	5.33	6.22	5.17
V	17	26					8				8
Y	37.6	69.7	16.9	68.7	56.3	57.8	39	14.4	26	70.1	51.3
Zn	62	64	130	71	96	75	70	76	69	65	68
Zr	132	161	135	120	108	92	121	147	144	143	152
La	55.9	76	6.6	61.7	47.2	43.5	37.9	22	44.9	73.5	54.5
Ce	88	107	16.6	132	113.5	107	86.5	63.4	90.5	127	107
Pr	14.5	20.6	1.25	17.85	13.1	12.5	10.05	3.51	13.2	20.4	16.3
Nd	53	77.4	4	67.6	49.6	46.7	37.1	10.1	49.3	76.3	62.2
Sm	10.2	15.8	1.18	15.3	11.15	10.2	8.39	1.82	10.4	16.75	13.9
Eu	0.36	0.33	0.05	0.13	0.11	0.07	0.43	0.22	0.37	0.18	0.38
Gd	7.86	13.2	1.4	12.55	9.15	8.92	7.7	1.8	7.6	13.8	11.55
Tb	1.21	2.03	0.35	1.99	1.57	1.61	1.23	0.44	1.15	2.21	1.79
Dy	7.22	12.6	2.64	13.05	10.2	10.6	7.67	2.86	6.66	13.75	10.7
Ho	1.4	2.49	0.61	2.59	2.06	2.22	1.5	0.64	2.56	2	2
Er	3.82	6.67	2.03	7.24	6.19	6.43	4.1	2.09	7.22	5.23	5.23
Tm	0.61	1.06	0.35	1.15	1	1.06	0.61	0.39	0.56	1.12	0.8
Yb	3.61	6.38	2.59	7.16	6.17	6.61	3.9	2.69	3.62	6.92	4.8
Lu	0.55	0.94	0.36	1.01	0.91	0.99	0.56	0.41	0.51	1.02	0.71

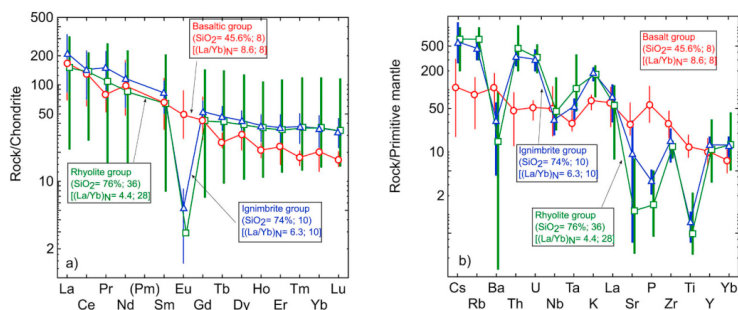


Fig. 5. (a) Chondrite-normalized rare-earth element (REE) plot for the SSMC samples (values taken from McDonough and Sun, 1995; mean value of silica and mean value of $(La/Yb)_N$ ratios along with the number of samples are also included for each rock groups). (b) Primitive mantle-normalized multi-element plot for the SSMC samples (values taken from McDonough and Sun, 1995; mean value of silica and mean value of $(La/Yb)_N$ ratios along with the number of samples are also included for each rock groups).

variable concentration of LILE ($Ba = 288\text{--}966$ ppm; $Sr = 371\text{--}519$ ppm; Table 3). The chondrite-normalized REE diagram (Fig. 5a) shows an enrichment in light REE (LREEs) and heavy REEs (HREEs; $(La/Yb)_{CN} = 3.9\text{--}7.3$), as well as small Eu anomalies $Eu(Eu/Eu^* = 0.70\text{--}1.15)$, the subscript $_{CN}$ refers to the chondrite normalized value, chondrite values are from McDonough and Sun (1995; Fig. 5a). Total REE concentration of the basaltic group varies from 101.09 to 176.63 ppm. The primitive mantle-normalized multi-element diagram (Fig. 5b) for the basaltic group samples display slightly flat patterns in incompatible elements with Nb ($Nb/Nb^* = 0.20\text{--}0.78$), Ta ($Ta/Ta^* = 0.21\text{--}0.90$) and P anomalies (Fig. 5b).

Trace element concentrations of the rhyolite-ignimbrite group samples are characterized by $Sc = 2.0\text{--}10$ ppm, $Cr = 1.2\text{--}10$ ppm, $Ni = 0.60\text{--}14$ ppm, and enriched and variable concentrations of LILE (e.g. $Ba = 6.8\text{--}410$ ppm, $Sr = 5.6\text{--}648$ ppm; Table 3). The chondrite-normalized REE diagram (Fig. 5a) displays an enrichment in LREEs and highly variable heavy REEs (HREEs) ($(La/Yb)_{CN} = 1.7\text{--}10.5$), as well as higher Eu anomalies as compared to the basaltic group. The strong negative Eu ($Eu/Eu^* = 0.02\text{--}0.36$) anomalies are probably consistent with plagioclase fractionation, either in the solid residuum left by partial melting (Rollinson, 1993). The primitive mantle-normalized multi-element diagram for the rhyolite-ignimbrite group shows enriched patterns in incompatible elements with Nb ($Nb/Nb^* = 0.26\text{--}1.06$), Ti ($Ti/Ti^* = 0.04\text{--}0.28$), Ba and P anomalies (Fig. 5b).

6. Discussion

6.1. Alteration effects

The loss-on-ignition (LOI) values for the basaltic group display a range from 0.496 to 2.48%. The LOI of the rhyolitic-ignimbrite group

shows the range from 0.627 to 1.49%, except four samples that display high values $> 5\%$ m/m. Therefore, the effect of post-emplacment alteration on elemental mobility could be considered for only some samples. As probably an alteration-independent index for geochemical diversity, Zr concentration implements a significant correlation with other elements and can be used to test their mobility (Pearce et al., 1992; Wang et al., 2016). Similarly, in this work, elements including $(TiO_2)_{adj}$, Ba, Rb, V, Nb, and Sr, for volcanic rocks plotted against Zr to evaluate their mobility during alteration (graph not shown) displayed a statistically significant correlation. Therefore, these elements could be considered relatively immobile during alteration and are useful in understanding of petrogenetic interpretation.

6.2. Petrogenetic processes

Petrography analysis has revealed disequilibrium evidence in the basaltic group rocks, whereas (Eu/Eu^*) and (Th/Yb) bivariate plots show a limited contribution of the continental crust in their petrogenesis (Fig. 6a–b). In order to explore the effects of magma mixing between the felsic group and mafic group magmas, a mixing model were developed between two samples L2 and SLP01. Sample L2 has the most felsic nature with $SiO_{2(adj)} = 79.41$ %m/m, $MgO_{(adj)} = 0.04$ %m/m (values reported by Leroy et al., 2002), sampled in the study area, and sample SLP01 has more mafic characteristics i.e., $SiO_{2(adj)} = 39.81$ %m/m, $MgO_{(adj)} = 4.59$ %m/m for the SSMC (Fig. 7a and b). It is clear from Fig. 7 that the basaltic group lavas plot close to the representative mixing curve, but also show shift of some points from the modelled mixing curve, probably as a result of the prior fractional crystallization process.

The $(Gd/Yb)_{PM}$ versus Eu/Eu^* and Nb/Yb versus Th/Yb ratios of the rhyolite-ignimbrite group rocks from the SSMC may be comparable to

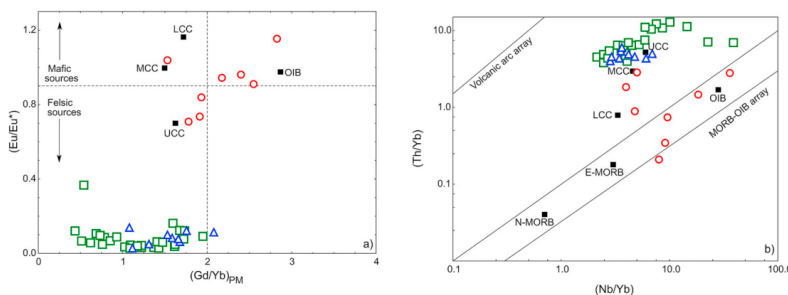


Fig. 6. Geochemical variation diagrams for volcanic rocks from the SSMC. (a) (Eu/Eu^*) – $(Gd/Yb)_{PM}$ diagram; (b) (Nb/Yb) – (Th/Yb) diagram. Abbreviations: N – MORB – normal Mid – Ocean Ridge basalt; E – MORB – enriched Mid – Ocean Ridge basalt; OIB – Ocean Island basalt; LCC – lower continental crust; MCC – middle continental crust; UCC – upper continental crust.

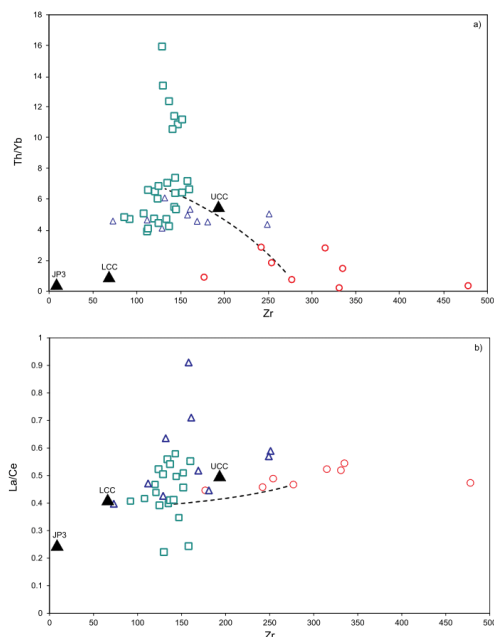


Fig. 7. (a) Zr–Th/Yb and (b) Zr–La/Ce variation diagrams for the basaltic group and rhyolite-ignimbrite group rocks from the SSMC showing mixing model. JP3: hornblende pyroxenite sample represent an enriched mantle (taken from Heinrich and Besch, 1992); LCC: lower continental crust (value taken from Rudnick and Gao, 2003); UCC: upper continental crust (value taken from Rudnick and Gao, 2003).

the continental crust (Fig. 6a–b), especially when the uncertainty in the average crustal compositions could be taken into account.

A quantitative treatment was carried out to represent a partial melting process for the rhyolite-ignimbrite group of the SSMC by applying the modal batch partial melting equations reported by Zou (2007). Trace and rare-earth element compositions from the average upper and lower continental crust reported by Rudnick and Gao (2003) were used as the crustal source (C_c). The mineralogical compositions of crustal reservoirs were as follows: for lower crust (LCC): $0.30 \text{ Pl} + 0.30 \text{ Cpx} + 0.20 \text{ Gt} + 0.10 \text{ Amp} + 0.05 \text{ Opx} + 0.05 \text{ Op}$ and upper crust (UCC): $0.50 \text{ Pl} + 0.10 \text{ Fsp} + 0.15 \text{ Bt} + 0.05 \text{ Amp} + 0.30 \text{ Qtz} + 0.01 \text{ Ap}$ (see more detail in Velasco-Tapia, 2017). The partial melting models for the rhyolite-ignimbrite group were carried out for melting degrees of 20–50% ($F = 0.2\text{--}0.5$). The results obtained were used to elaborate chondrite-normalized REE plots (Fig. 8a–b). The rhyolite-ignimbrite group rocks showed patterns with comparable features with the upper continental crust and their derived liquids. Thus, the crustal melting process may have taken place at different depths with conditions of the upper continental crust.

6.3. Tectonic setting

6.3.1. Evaluation of Nb and Ta anomalies

The evaluation of the Nb and Ta anomalies has been described statistically by Verma (2015, 2019) for basic, intermediate, and acid magmas from the Mexican Volcanic Belt, numerous continental and island arcs, continental rifts or extensional areas, including continental collision setting around the world. In this work, we have calculated

these values for the basic and acid rocks from the SSMC rocks and compared them with the compilation of Verma (2015) in Table 4. The calculated value for Nb and Ta anomaly values for basic (0.604 ± 0.343 and 0.53 ± 0.33), and acid (0.676 ± 0.218 and 1.189 ± 0.445) units could be considered similar to the continental rift or extension areas of the Eastern Anatolia (Turkey) and the Rio Grande rift, New Mexico (USA) and different from the continental arcs (Table 4). Therefore, the volcanic rocks from SSMC may have been derived from the rifting or extensional environment.

6.3.2. Subduction or continental crust sensitive parameters

Subduction–continental crust signal parameters were proposed and used to identify the subduction or continental crust contribution in a tectonic environment (Verma, 2009, 2015, 2019). Higher values of these parameters, especially those having LILE in the numerator and HFSE in the denominator, reflect contribution from the subduction process. Alternatively or concurrently, high values of these parameters may also represent a significantly higher influence from the continental crust. This is especially applied for evolved intermediate and acid rocks (Verma and Verma, 2018). These considerations make it difficult to unambiguously infer the tectonic implications from evolved rocks. Nevertheless, we used 6 such parameters (two listed below; others in the Supplementary file) for statistically comparing between the SSMC and Rio Grande rift, New Mexico (RG) rocks.

The (LILE4/LREE3)_E parameter is the “four LILE to three LREE” ratio and (LILE5/HFSE5)_E parameter is the “five LILE to five HREE” ratio as follows:

$$[\text{LILE4/LREE3}]_E = \frac{[(K_{sa}/K_E) + (Rb_{sa}/Rb_E) + (Ba_{sa}/Ba_E) + (Sr_{sa}/Sr_E)]/4}{[(La_{sa}/La_E) + (Ce_{sa}/Ce_E) + (Nd_{sa}/Nd_E)]/3} \quad (1)$$

$$[\text{LILE5/LHFSE5}]_E =$$

$$= \frac{[(K_{sa}/K_E) + (Rb_{sa}/Rb_E) + (Cs_{sa}/Cs_E) + ((Ba_{sa}/Ba_E)) + (Sr_{sa}/Sr_E)]/5}{[(Ti_{sa}/Ti_E) + (P_{sa}/P_E) + (Zr_{sa}/Zr_E) + ((Hf_{sa}/Hf_E)) + (Ta_{sa}/Ta_E)]/5} \quad (2)$$

where the subscript “sa” stands for the sample and “E” for bulk silicate earth (normalization values from McDonough and Sun, 1995).

The basaltic group from the SSMC shows lower values for 2 parameters (LILE4/LREE3)_E and (LILE4/HFSE4)_E than the Central American Volcanic Arc (CAVA); the other 3 parameters show overlapping confidence interval of the mean; and the remaining parameter (LREE3/HFSE5)_E cannot be compared because of the lack of information for the CAVA (Table 5). On the other hand, the SSMC basaltic rocks show overlapping values for all 4 parameters with the RG rocks (Table 5). These results imply that the SSMC basaltic group rocks might be considered similar to the Rio Grande basic rocks but different from the CAVA.

The SSMC rhyolite-ignimbrite group samples display overlapping values for 4 out of 5 parameters with the CAVA; (LREE3/HFSE5)_E cannot be compared whereas (LILE4/LREE3)_E shows lower value for the SSMC than the CAVA. On the other hand, the SSMC evolved rocks show overlapping ranges for both parameters (LILE4/HFSE4)_E and (LREE3/HFSE5)_E that can be compared with the RG (Table 5). These results are also consistent with the similarities of the SSMC with the RG.

Thus, for both basaltic and evolved rocks, the similarity of the SSMC with the RG and at least some significant differences between the SSMC and the CAVA can be inferred. This may imply similarity of the tectonic regime of the SSMC and RG and differences of the SSMC and CAVA regions.

6.3.3. Multi-dimensional discrimination diagrams

New multi-dimensional discrimination diagrams were also used to infer the tectonic setting, because conventional bivariate and ternary diagrams for tectonic discrimination were shown to work unsatisfactorily (Verma, 2010). The failure of the older diagrams is based

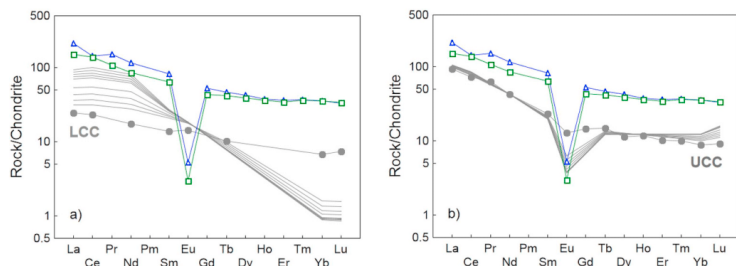


Fig. 8. (a) Chondrite-normalized rare-earth element patterns for the Lower Continental Crust (LCC) and (b) Chondrite-normalized rare-earth element patterns for the Upper Continental Crust (UCC); for both figures partial melts (solid gray lines) were generated at degree of melting $F = 0.2-0.5$, applying a batch melting equation by Zou (2007) for comparison of rhyolite-ignimbrite group rocks (average value, $n = 46$). Gray circles line correspond to LCC or UCC normalized pattern.

Table 4

Statistical information on the Nb and Ta anomaly (with respect to Ba and La) for mafic and felsic rocks from San Miguelito Complex (SSMC), San Luis Potosi Mexico (this work) and comparison with other arcs and continental rifts (taken from Verma, 2015).

Area	Magma type	$\{Nb/Nb\}_{pm}$		$\{Ta/Ta\}_{pm}$	
		Mean \pm standard deviation (number of samples) $\bar{x} \pm s$ (n)	99% Confidence limits (CL) of the mean	Mean \pm standard deviation (number of samples) $\bar{x} \pm s$ (n)	99% Confidence limits (CL) of the mean
Sierra de San Miguelito	Basic	0.604 \pm 0.343 (8)	0.179–1.029	0.53 \pm 0.33 (4)	(0) – 1.49
Volcanic Complex	Acid	0.676 \pm 0.218 (25)	0.554–0.798	1.189 \pm 0.445 (38)	0.92–1.457
<i>Continental arcs</i>					
Central American Volcanic Arc	Basic	0.13 \pm 0.06 (59)	0.11–0.15	0.18 \pm 0.11 (34)	0.13–0.23
	Acid	0.105 \pm 0.040 (25)	0.083–0.127	0.105 \pm 0.029 (4)	0.020–0.189
Andes (Chile)	Basic	0.20 \pm 0.05 (29)	0.17–0.23	0.147 \pm 0.037 (9)	0.17–0.23
	Acid	0.177 \pm 0.031 (49)	0.165–0.189	0.194 \pm 0.034 (41)	0.180–0.208
Andes (Peru)	Acid	0.25 \pm 0.10 (37)	0.21–0.30	0.26 \pm 0.11 (33)	0.21–0.31
Andes (Ecuador)	Basic	0.079 \pm 0.009 (7)	0.065–0.090	–	–
	Acid	0.106 \pm 0.018 (210)	0.103–0.110	0.17 \pm 0.07 (16)	0.11–0.22
<i>Continental rifts or extensional areas</i>					
San Juan volcanic field, Colorado (USA)	Acid	0.12 \pm 0.05 (7)	0.06–0.19	–	–
Rio Grande rift, New Mexico (USA)	Basic	0.7 \pm 0.5 (29)	0.5–1.0	0.8 \pm 0.5	0.5–1.1
Central Mexican Volcanic Belt	Basic	0.54 \pm 0.25 (38)	0.43–0.65	0.75 \pm 0.16 (30)	0.67–0.84
	Acid	0.152 \pm 0.033 (194)	0.146–0.158	0.22 \pm 0.05 (184)	0.21–0.23
Eastern Anatolia (Turkey)	Basic	0.782 \pm 0.014 (20)	0.773–0.791	0.784 \pm 0.038 (20)	0.760–0.809
	Acid	0.50 \pm 0.13 (25)	0.43–0.58	0.680 \pm 0.017 (24)	0.580–0.779

Table 5

Statistical information of subduction signal for the Sierra de San Miguelito complex (SSMC), Rio Grande Rift (RG; data compiled from Duncker et al., 1991; Gibson et al., 1992; McMillan et al., 2000) and Central American Volcanic Arc (CAVA; data compilation by Verma, 2015).

Parameter	SSMC			RG			CAVA			
	n	\bar{x}	u_{99}	n	\bar{x}	u_{99}	99% confidence limit of the mean	n	\bar{x}	u_{99}
Basic										
(LILE4/LREE3) _E	8	1.67	0.82	21	1.44	0.272	2.7–3.2			
(LILE5/HFSE5) _E	4	3.30	7.2	1	3.75	–	4.1–5.7			
(LILE4/HFSE4) _E	8	1.47	0.58	45	1.884	0.064	4.2–5.7			
(LREE3/HFSE5) _E	4	1.25	1.32	22	2.13	0.48	–			
(LILE4/HREE3)	4	10.29	0.366	–	–	–	7.8–9.1			
(LILE4/LREE4)	4	2.07	2.09	–	–	–	2.8–3.03			
Acid										
(LILE4/LREE3) _E	29	2.80	0.396	–	–	–	3.4–4.5			
(LILE5/HFSE5) _E	29	13.48	2.0	–	–	–	3.7–16.0			
(LILE4/HFSE4) _E	37	15.71	0.61	4	10.87	4.14	4.3–16.0			
(LREE3/HFSE5) _E	30	2.5	0.68	1	3.03	–	–			
(LILE4/HREE3)	37	13.93	0.58	–	–	–	6.0–28.0			
(LILE4/LREE4)	29	2.772	0.119	–	–	–	2.1–7.0			

on the following criteria: (a) use of limited databases to construct them; (b) the problem of closed or constant sum compositional variables not attended in older diagrams; and (c) eye-fitted tectonic field boundaries

in the older diagrams. All these problems have been solved in the multidimensional solutions (Armstrong-Altrin, 2015; Verma, 2015, 2019; Verma and Armstrong-Altrin, 2016; Verma et al., 2012, 2013, 2015). Therefore, new multi-dimensional discrimination diagrams for basic and acid rocks, which are based on log-ratio transformation, linear discrimination and canonical analysis, were used for tectonic interpretation.

The new multi-dimensional discrimination diagrams (Verma and Agrawal, 2011; Verma et al., 2013) indicated a within-plate setting for the SSMC volcanic rocks (Fig. 9). The findings of new multi-dimensional discrimination diagrams are coherent with the syn-transform extension regime and intraplate volcanism that was produced during the Oligocene to Miocene within the southern Basin and Range province (western USA, northwestern Mexico; Dickinson, 2002; Cosca et al., 2014).

More constraints, especially the study of basic volcanic rocks and underlying crust from this and surrounding areas and appropriate well-constrained thermochemical models, are required before a comprehensive petrogenetic and tectonic model could be put forth. This should constitute future studies.

7. Conclusions

The following conclusions, based on the results and observations, can be summarized:

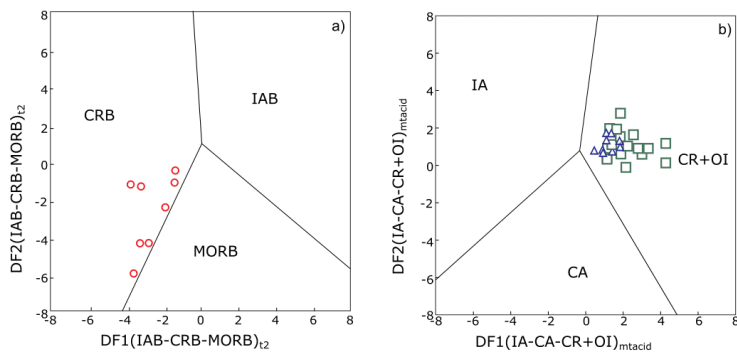


Fig. 9. (a) Discriminant – function multidimensional diagrams based on log – transformed ratios of major and trace elements for tectonic discrimination of the basic units from the SSMC (Verma and Agrawal, 2011). (b) Discriminant – function multidimensional diagrams based on log – transformed ratios of major and trace elements for tectonic discrimination of the acid units from the SSMC (Verma et al., 2013).

1. The San Miguelito volcanic complex is constituted by the basaltic components of trachybasalt, basalt, and foidite compositions and the rhyolite-ignimbrite component of rhyolitic and trachyte compositions.
2. Basaltic group rocks from the San Miguelito volcanic complex show foidite to trachybasalt composition. The REE chondrite-normalized patterns display an enrichment in LREE, a small or absent Eu-anomaly and a flat-shape for HREE. The primitive mantle normalized diagram shows slightly flat patterns in incompatible elements with small or absent anomalies of Nb, Ta, and P. In contrast, the rhyolite-ignimbrite group rocks display a trachydacite and rhyolite compositions. The REE chondrite-normalized patterns show an enrichment in LREE, a negative Eu-anomaly and a flat shape for HREE. The primitive mantle normalized diagram shows an enrichment in incompatible elements with negative anomalies in Nb, Ti, Ba and P.
3. Geochemical features indicated that the basaltic group rocks from the San Miguelito volcanic complex were derived from mantle through crustal evolution by relatively small amount of magma mixing processes with felsic volcanism. The felsic volcanic rocks were generated by partial melting of the upper continental crust.
4. New multi-dimensional tectonic discrimination diagrams indicate that these volcanic rocks are related to within-plate setting. This is consistent with Nb- and Ta anomalies as well with the syn-transformation extension regime from the Basin and Range province.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.jsames.2019.102311>.

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CAPITULO 6.-

GEOCRONOLOGÍA $^{40}\text{Ar}/^{39}\text{Ar}$ Y PETROGENÉISIS DEL COMPLEJO VOLCÁNICO DE LA SIERRA DE SAN MIGUELITO, MESA CENTRAL, MÉXICO

6.1 Presentación del trabajo de investigación publicado

El presente artículo de investigación ha sido publicao en *LITHOS* en donde el autor del presente trabajo de tesis participó activamente. Dentro de este trabajo me encuentro como primer autor y el Dr. Sanjeet Kumar Verma, la Dra. Tiffany Barry, el Dr. Surendra Pal Verma y el Dr. José Ramón Torres Hernández son los co-autores del presente artículo de investigación.

Este trabajo de investigación ha contribuido con nueva información dentro del Complejo Volcánico de la Sierra de San Miguelito (CVSSM). Dentro de este trabajo se ha presentado nueva información geocronológica para las unidades volcánicas del CVSSM por el método de $^{40}\text{Ar}/^{39}\text{Ar}$, la cual sirvió como apoyo para que, junto con la información de datación realizada por trabajos previos, se elaborara una limitación entre los distintos episodios magmáticos dentro del CVSSM. De igual forma, en el presente trabajo se reporta nuevos datos isotópicos (Sr-Nd-Pb), que junto a información isotópica de la provincia de Sierras y Cuencas (Basin and Range), sirvieron para la generación de modelos que muestran el comportamiento de la fuente de las rocas volcánicas del CVSSM.

A su vez, en el presente trabajo se muestra que el CVSSM no exhibe un bimodalismo en la composición de las rocas, como se pensaba en trabajos previos, sino, el CVSSM exhibe un rango composicional amplio (rocas básicas, intermedias y ácidas). De igual forma, en el trabajo se reporta nuevos modelos cuantitativos de procesos de fusión parcial del manto para las rocas básicas, modelos de asimilación y cristalización fraccionada para las rocas intermedias y modelos de fusión parcial cortical para las rocas ácidas, los cuales brindaron una mejor comprensión en la evolución de las rocas volcánicas del CVSSM.

Por último, una contribución importante dentro de este trabajo es la discusión entre las limitaciones petrogenéticas de la información de edades, datos geoquímicos e isotópicos a través del área de estudio dentro del contexto tectónico del desarrollo de la sección sur de la provincia de Sierras y Cuencas (Basin and Range). Es de igual importancia destacar que para el momento en que el presente trabajo ha sido publicado se ha generado un nuevo esquema propuesto por Verma (2020), el cual llega a invalidar los procedimientos existentes de discriminación tectónica. Este nuevo esquema será aplicado para futuros trabajos dentro del CVSSM.

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Research Article

$^{40}\text{Ar}/^{39}\text{Ar}$ geochronology and petrogenesis of the Sierra de San Miguelito Volcanic Complex, Mesa Central, Mexico

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ABSTRACT

The southern part of the Mesa Central (MC) province, Mexico, is formed of several Cenozoic volcanic complexes. The Sierra de San Miguelito Complex (SSMC) is in the south-eastern part of the MC. The SSMC consists of: (1) mafic volcanic rocks of porphyritic texture and trachybasalt/basalt compositions; (2) intermediate volcanic rocks of porphyritic texture and basaltic-trachyandesite, basaltic andesite and andesite compositions; and (3) silicic volcanic rocks of porphyritic texture and rhyolite composition. New $^{40}\text{Ar}/^{39}\text{Ar}$ dating results, in combination with major- and trace-element data, and Sr-Nd-Pb isotope data, are used to investigate the petrogenesis and geodynamic evolution of SSMC. The $^{40}\text{Ar}/^{39}\text{Ar}$ radiometric age data constrains the magmatic events in the SSMC to between 34 and 21 Ma. Chondrite-normalized rare-earth element patterns are distinct for each volcanic succession; mafic and intermediate lavas have relatively flat light rare earth element (LREE) and large ion lithophile element (LILE) patterns, whereas the silicic volcanic rocks show enrichment in LREE and high field strength elements (HFSE). Within each volcanic phase, the total rare-earth element concentrations increase from mafic to silicic, and the size of the negative Eu anomalies progressively increase (Eu/Eu* from 0.02 to 1.04). The initial $^{87}\text{Sr}/^{86}\text{Sr}$ ratios are widely distributed (from 0.70344 to 0.71973) whereas the initial $^{143}\text{Nd}/^{144}\text{Nd}$ ratios are somewhat low and show a narrower range (0.51245 to 0.51287), indicating the mafic magmas derived from a slightly heterogeneous source. Geochemical modelling of the mafic volcanic rocks reveals two sources of magma: (1) a parental magmas generated from melting underlying lithospheric mantle; and (2) a second lithospheric melt contaminated by lower crust. Intermediate magmas evolved from assimilation and fractional crystallization (AFC) processes of both lithospheric melts, at shallower levels. The silicic volcanic rocks in the area, however, were probably derived from partial melting of sedimentary rocks within the upper-middle continental crust. New multidimensional tectonic discrimination diagrams, combined with the magmatic model, indicates that volcanic activities in the region were generated in an extensional environment.
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1. Introduction

The Basin and Range (BR) province is one of the most prominent tectonic features across the western part of the North American plate (Fig. 1a). The BR extends from Canada through the western USA to north-western Mexico (Fig. 1a) and is a composite Cenozoic taphrogen with a present configuration and internal geometry derived from

multiple phases of extension (Dickinson, 2002). Late Paleocene, Mid-Miocene and Pliocene syn-transform extension within the southern BR (western USA and north-western Mexico) have been related to rapid migration of the Rivera triple junction past Baja California to its current location between the tip of the Baja Peninsula and mainland Mexico (Fig. 1a; Dickinson, 2004; Cosca et al., 2014). Features of Middle to Late Cenozoic extension and magmatism generated through BR have been extensively studied (e.g. Berglund et al., 2012; Cosca et al., 2014; Dickinson, 2002, 2004; Jones et al., 1992). Nevertheless, its origin, dynamics and evolution are still a much-debated topic of discussion within the earth sciences (e.g., Atwater, 1970; Coney and Harms, 1984; Cosca et al., 2014; Glazner and Bartley, 1984; Henry and

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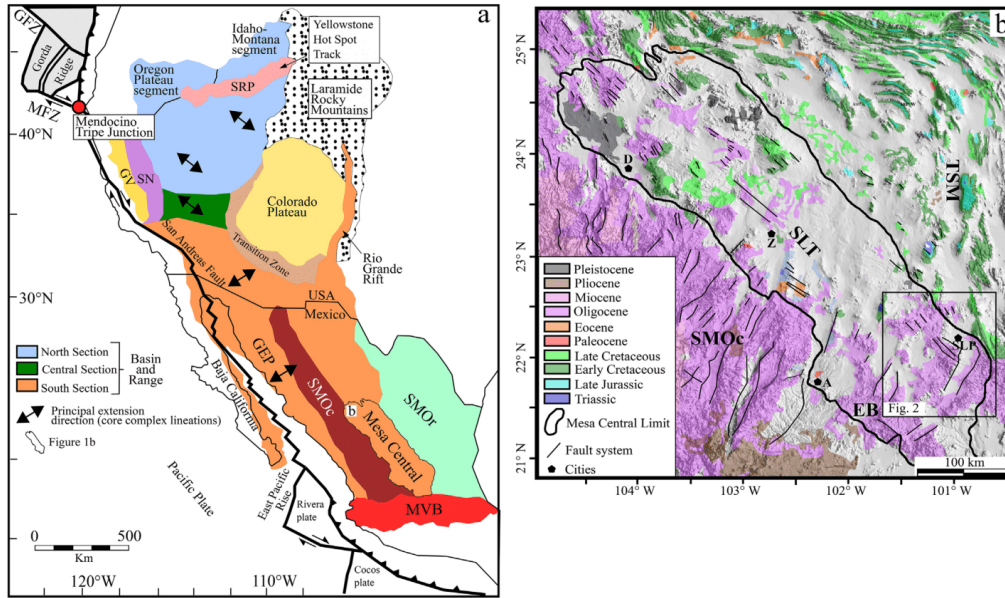


Fig. 1. (a) Present-day regional tectonic map of western North America and Basin and Range province with subdivisions of the northern, central and southern Basin and Range from Jones et al. (1992) and Sonder and Jones (1999) (modified from Cosca et al., 2014); (b) Simplified geological map of the Mesa Central province (modified from Nieto-Samaniego et al., 2007) showing Triassic metamorphic and sedimentary basement; Late Jurassic sediments and sandstones; Early Cretaceous limestone; Late Cretaceous sediments and sandstones; Paleocene plutonic rocks; Eocene silicic volcanism; Oligocene silicic and pyroclastic volcanism; Miocene mafic volcanism; Pliocene mafic volcanism; and Pleistocene mafic volcanism. Abbreviations are as follows: GEP-Gulf Extensional Province; MFZ-Mendocino Fracture Zone; MVB-Mexican Volcanic Belt; RFZ-Rivera Fracture Zone; GFZ-Gorda Fracture Zone; SRP-Snake River Plain; SN-Sierra Nevada; GV-Great Valley; STFS-San Luis Tepehuanes Fault Systems; TSM-Taxco-San Miguel de Allende Fault System; EB-El Bajío Fault System; SMOc-Sierra Madre Occidental province; SMOr-Sierra Madre Oriental; USA-United States of America; D-Durango city; Z-Zacatecas city; A-Aguascalientes city; SLP-San Luis Potosí city.

Aranda-Gómez, 2000; Parsons, 1995; Severinghaus and Atwater, 1990; Wernicke et al., 1988).

The Mesa Central (MC) province, Mexico, sits within the southern limits of the BR. The province is within an elevated plateau that extends across central-northern Mexico and is bound to the north and east by the Sierra Madre Oriental (SMOr) and to the south and west by the Sierra Madre Occidental (SMOc; Fig. 1b; Nieto-Samaniego et al., 2007). Within MC there are several fault systems (Fig. 1b): (i) the NW-SE trending El Bajío (EB) in the south; (ii) the Taxco-San Miguel de Allende (TSM) in the west; and (iii) the San Luis-Tepehuanes (SLT) extending within the Mesa Central. The MC is divided into two main regions by ~1600 km long lineament towards the NW direction: (a) the northern region, which is characterized by advanced stages of erosion, alluvial-lacustrine basin development, and only a low volume of magmatic activity during the Oligocene and Quaternary; whereas (b) the southern region, is mainly covered by Paleogene-Neogene volcanic rocks and cross-cut by several normal faults (Fig. 2a; Nieto-Samaniego et al., 1996; Nieto-Samaniego et al., 1999; Nieto-Samaniego et al., 2007). Several geological, geophysical and geochemical studies have been carried out within the southern region of the MC (i.e., Aguirre-Díaz and Labarthe-Hernández, 2003; Labarthe-Hernández et al., 1982; López-Loera and Tristán-González, 2013; Nieto-Samaniego et al., 2007; Orozco-Esquivel et al., 2002; Sieck et al., 2019), but the origin, evolution, timing, and cause of the volcanism is still poorly constrained.

One of the mostly poorly understood parts of the MC is the bimodal Sierra de San Miguelito Complex (SSMC; Fig. 2b) in the south (i.e., Labarthe-Hernández et al., 1982; Rodríguez-Ríos et al., 2007).

Limited geological work has been done on the area of SSMC but the essential information, such as robust age constraints and detailed geochemical and radiogenic isotope data, is still missing. So far, the volcanic rocks have predominantly been dated by the K–Ar method that has several substantial drawbacks (Clay et al., 2015), including much larger uncertainties than Ar–Ar ages (Lee, 2015). Thus, the timing and origin of the highly evolved explosive rhyolitic volcanism in the SSMC, relative to the spatially associated basaltic volcanism remains enigmatic.

In this work, we present new $^{40}\text{Ar}/^{39}\text{Ar}$ ages along with whole-rock chemistry and Sr–Nd–Pb isotope data of the magmatic units across the area. We present the age, geochemical and isotopic constraints on the mafic, intermediate and silicic rocks in the area and discuss their petrogenesis in the context of the development of the southern parts of the Basin and Range.

2. Geological framework

The southern region of the MC is mainly characterized by widespread Cenozoic faults that commonly bound basins filled with fluvial and lacustrine deposits. These faults reflect a complex tectonic stress regime that was generated during the Mid- to Late Cenozoic and marks a continuous extension to the BR (Fig. 1b). The region is underpinned by a Mesozoic basement comprising marine calcareous rocks of the SMOr, and volcanic-flysch sequences of the Sierra de Guanajuato Complex (Centeno-García, 2017; Orozco-Esquivel et al., 2002).

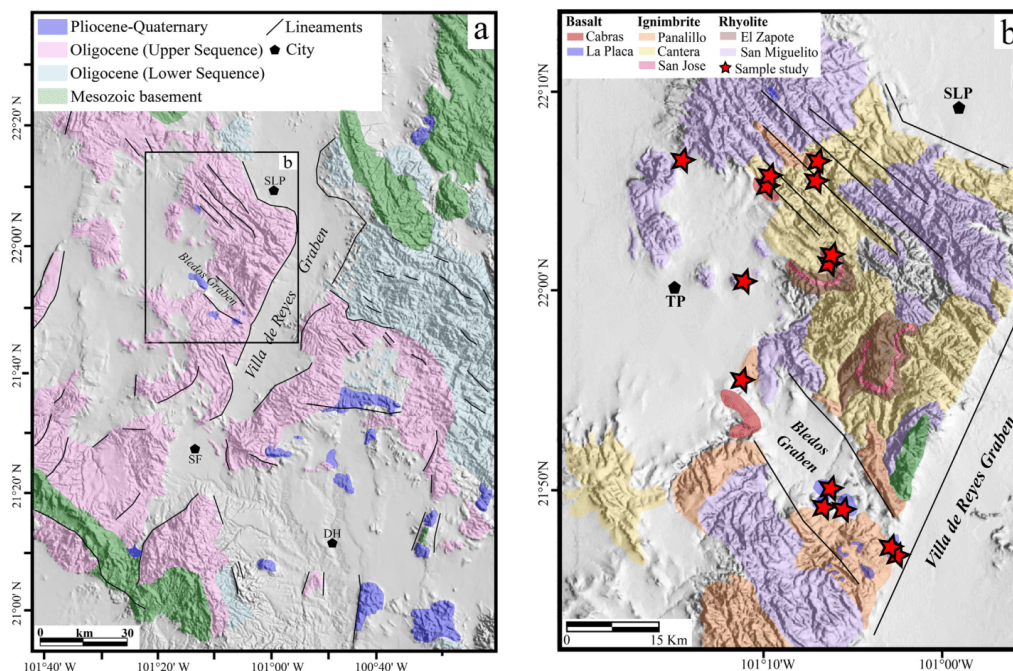


Fig. 2. (a) Simplified geological map showing the division of the stratigraphic units and the location of the main stratigraphy of the southern Mesa Central province (Modified after Orozco-Esquivel, et al., 2002); (b) Geological map of the Sierra de San Miguelito complex (SSMC). Basalt from late Oligocene to Miocene; Ignimbrite and Rhyolite from early Oligocene to late Oligocene. Abbreviations are as follows: SLP-San Luis Potosí city, SF-San Felipe city, DH-Dolores Hidalgo city, TP-Tepehate city.

Prior to the Oligocene, there is little evidence of Cenozoic cover sequence within the MC. During the Paleocene, a few post-Laramide granite intrusions occurred within the southern part of the MC (Angeles-Moreno et al., 2017). By the Eocene, continental sandstones and conglomerates were deposited along with intermittent pyroclastic deposits and mafic-intermediate lavas (Aranda-Gómez and McDowell, 1998; Labarthe-Hernández et al., 1982). Regional extension occurred across the southern part of the MC, beginning in the Eocene, but continuing through the Miocene, forming NW-SE and NE-SW oriented grabens that suggest episodes of extension during this period (Nieto-Samaniego et al., 1996). These basins were later filled with alluvial and lacustrine deposits (Nieto-Samaniego et al., 1996).

During the Oligocene, a voluminous magmatic event took place within the southern region of the MC, forming a thick volcanic succession of mafic, intermediate and silicic volcanic rocks. This magmatic event has been divided into a lower and upper sequence (Fig. 2a; Orozco-Esquivel et al., 2002). The lower sequence comprises mainly of altered intermediate volcanic rocks and semi-altered pyroclastic deposits. The ages of the lower sequence have been reported as 32.8 ± 0.9 to 29.5 ± 1.5 Ma by K–Ar technique (Cerca-Martínez et al., 2000; Labarthe-Hernández et al., 1982). The upper sequence consists predominantly of silicic domes, the majority associated with lava and pyroclastic deposits. The domes are aligned along subsidiary faults or fractures parallel to the direction of the major fault systems through the area (Orozco-Esquivel et al., 2002). The ages of this sequence have previously been reported as 30.1 ± 0.8 to 27.0 ± 0.7 Ma by K–Ar technique (Nieto-Samaniego et al., 1996), despite it being situated on top the lower sequence. No clearly defined caldera structures have been

identified for the upper sequence, but rare pyroclastic dikes, underlying the domes, have been observed with a similar orientation to the regional faults (Torres-Hernández et al., 2006). The uppermost part of the upper sequence is represented by a small number of isolated outcrops of mafic and intermediate volcanic rocks that erupted during the late Oligocene-Miocene (Orozco-Esquivel et al., 2002).

The SSMC is situated within the upper sequence and is mainly bounded by the Villa de Reyes Graben (Fig. 2a; Nieto-Samaniego et al., 1999, 2007). The volcanic activity of the SSMC was emplaced through fault systems that exhibit a “domino style geometry” with strike direction of 300° – 340° dip to the SW and NW systematically with dip angles between 12° and 31° (Xu et al., 2004). The stratigraphy of the SSMC covers the Epoch of Oligocene to early Miocene (Fig. 3). Lithologically, the units can be categorized into three types: (a) mafic volcanic rocks (Cabras unit); (b) intermediate volcanic rocks (La Placa unit); (c) silicic volcanic rocks (San Miguelito, San José, El Zapote, Cantera and Panalillo units).

2.1. Mafic volcanic rocks

The mafic volcanic rocks consist of the Cabras unit (Fig. 3). This unit is found as isolated outcrops of lava, displaying a thickness of <25 m, and is mainly characterized by an aphanitic-porphyrific texture, scarce olivine phenocrysts, and rare vesicles in a vitreous matrix (Torres-Sánchez et al., 2019; Tristán-González et al., 2009). Tristán-González et al. (2009) and Aguillón-Robles et al. (2014) presented whole-rock K–Ar ages of 22 to 21 Ma for this unit.

		This work			
		Literature Age	Plateau age	Isochron age	
Miocene	23.03 Ma	Cibros unit	21.5 ± 0.5 to 22.1 ± 0.5 Ma ⁵⁻⁶	22.21 ± 0.29 Ma	21.93 ± 0.59 Ma
		Panalillo unit	25.4 ± 0.6 to 31.9 ± 0.7 Ma ^{1,5}	31.05 ± 0.37 Ma	29.50 ± 2.30 Ma
Late Oligocene	27.82 Ma	Placa unit	26.9 ± 0.4 to 29.0 ± 0.7 Ma ⁵⁻⁶	30.13 ± 0.42 Ma	30.47 ± 0.99 Ma
		Cantera unit	29.0 ± 1.5 Ma ⁵	32.10 ± 0.38 Ma	32.50 ± 0.39 Ma
Early Oligocene		Zapote unit	27.4 ± 0.4 to 31.2 ± 0.7 Ma ⁴⁻⁵	33.48 ± 0.43 Ma	33.40 ± 0.68 Ma
		San José unit	29.0 ± 1.5 Ma ⁵	33.95 ± 0.44 Ma	34.09 ± 0.48 Ma
		San Miguelito unit	21.1 ± 0.3 to 32.7 ± 1.0 Ma ¹⁻⁶	33.17 ± 0.67 Ma	34.80 ± 5.3 Ma
	33.9 Ma	Rupelita unit			

Fig. 3. Schematic stratigraphic column of the volcanic rocks from the SSMC. Ages from this study were taken from Table 4. Literature age are taken from the following: ¹Labarthe-Labarthe-Hernández et al. (1982); ²Tuta et al. (1988); ³Aguillón-Robles et al. (1994); ⁴Aguillón-Robles et al. (2009); ⁵Tristán-González et al. (2009); ⁶Aguillón-Robles et al. (2014).

2.2. Intermediate volcanic rocks

The intermediate volcanic rocks comprise of the La Placa unit (Fig. 3); previous studies (Aguillón-Robles et al., 2014; Tristán-González et al., 2009) have reported whole-rock K—Ar ages of 29 to 26 Ma. This unit consists of massive lavas flows that are only exposed in isolated outcrops, displaying a thickness range from 10 to 15 m, and characterized by scarce olivine phenocrysts in a vitreous matrix.

2.3. Silicic volcanic rocks

The most voluminous rock type in the SSMC is the silicic volcanic rocks represented by the San Miguelito, San José, El Zapote, Cantera and Panalillo units (Fig. 3). The San Miguelito and El Zapote units consist of exogenous lava domes, which during the final stages of development erupted thick breccias on the top. The overall thickness of the San Miguelito and El Zapote units is variable (130–800 m) and is higher at the center of the domes. Previous studies (Aguillón-Robles et al., 1994, 2014; Tristán-González et al., 2009) reported whole-rock K—Ar dates of 33 to 21 Ma for the San Miguelito and El Zapote units. In some places, between the San Miguelito and El Zapote units, lies the San José unit, though elsewhere El Zapote sits directly on top of the San Miguelito unit. The San José unit is 3–15 m thick and consists of ash and a pyroclastic deposit which is mildly welded (Tristán-González et al., 2009).

Above the El Zapote unit, is the Cantera unit - a highly welded pyroclastic deposit characterized by grayish white to pink pumice and ash, abundant fiamme and lithic fragments of rhyolite and sandstone compositions, in a devitrified glassy matrix. It's thickness varies between 2 and 30 m thick. The unit covers much of the SSMC and overlays the San Miguelito unit. Previously, the Cantera unit has been dated at about 29 Ma by K—Ar method (Caballero-Miranda et al., 2009; Tristán-González et al., 2009).

Finally, the youngest silicic rocks in the area are the Panalillo unit, previously dated at 29 to 26 Ma (Torres-Hernández et al., 2006;

Tristán-González et al., 2009). It is mainly composed of a pumice-ash pyroclastic deposit which is lithic-rich and is slightly less welded than the Cantera below. It has an average thickness of 20–30 m (Torres-Sánchez et al., 2019; Tristán-González et al., 2009).

3. Materials and methods

3.1. Sampling and petrography

For this study, representative samples of the main lithological units from the SSMC were selected. The locations of weathering, deformation and veins were avoided during sampling and only fresh samples were collected for ⁴⁰Ar/³⁹Ar dating and geochemical analysis. Stratigraphic and lithological boundaries for volcanic rocks were drawn, largely on geomorphologic characteristics tested in the field and later refined by petrological, geochemical and geochronological analyses. Modal compositions were determined at the Institute of Scientific and Technological Research of San Luis Potosí (IPICYT, San Luis Potosí, Mexico) by point counting on thin sections using a Leica petrographic microscope and a PELCON Automatic point counter (Table S1).

3.2. ⁴⁰Ar/³⁹Ar geochronology

Radiometric ⁴⁰Ar/³⁹Ar dating was undertaken using feldspar crystals (silicic samples) and whole rock pieces (mafic samples). Samples selected for radiometric dating were crushed, sieved, and washed repeatedly with de-ionized water, and the 250–350 μm fragments were selected. These fragments were again cleaned ultrasonically in acetone and de-ionized water, dried using the hot plate, and packaged in aluminum foil packets of 10 × 10 mm size prior to irradiation.

Irradiation was carried out at the McMaster Nuclear Reactor (McMaster University, Canada) for 101 h. Cadmium shielding was used, and the samples were held in position 8D. Neutron flux was monitored using biotite mineral standard GA1550 (99.738 ± 0.104 Ma;

Renne et al., 2011). Standards were packed for irradiation either side of the unknown samples and analyzed using the single grain fusion method, in a 1059 nm CSI fiber laser and MAP215–50 mass spectrometer. J values were calculated by linear extrapolation between two measured J values, and a 0.5% error on J is used. The values for J for each sample are reported in Table S3 (see the supplementary material file).

Irradiated samples were loaded into an ultra-high vacuum system and a 1059 nm CSI fiber laser was focused into the sample chamber and used to step-heat the sample. After passing through a liquid-nitrogen trap, extracted gases were cleaned for 5 min using three SAES AP-10 getters, two running at 450 °C and one at room temperature. Following this, the gases were let into a MAP 215–50 mass spectrometer for measurement, with the mass discrimination value for $^{40}\text{Ar}/^{36}\text{Ar}$ measured at 283. System blanks were measured before and after every two sample analyses. Gas clean-up and inlet was fully automated, with measurement of ^{40}Ar , ^{39}Ar , ^{38}Ar , ^{37}Ar , and ^{36}Ar , each for ten scans, and the final measurements are extrapolations back to the inlet time.

3.3. Whole rock geochemistry

Fifteen samples were jaw crushed and then powdered in an agate bowl obtaining a 400 μm mesh fine powder. Major elements were determined on fusion beads prepared from pre-ignited powders, which were fused with lithium metaborate flux (80% lithium metaborate and 20% lithium tetraborate) in a ratio of 1:5. Trace elements were determined on pressed powder pellets made with a mixture of 7 g of sample and 7% PVA (Polyvinyl Alcohol) solution. Analyses were carried out at the University of Leicester, United Kingdom, in a PANalytical Axios Advanced x-ray fluorescence (XRF) spectrometer. Internal standards BH-1, WS-1, BCS375, BCS 376, MRG-1 and NIM-D were analyzed to monitor precision and accuracy of the results. The analytical precision (2σ) and accuracy were considered between 2 and 5% for major and < 2% for trace elements. Rare earth element concentrations were determined at the University of Leicester by a Thermo Scientific ICAP-Qc quadrupole ICP mass spectrometer. Samples were prepared using a standard HF-HNO₃ digestion. Seven geochemical reference materials (BHVO-1, NIM-G, MRG-1, JSD-1, JSD-2, JSD-3, and JR-1) were used to calibrate and evaluate the analytical data quality.

3.4. Isotopic analyses

Twelve samples from the SSMC were analyzed for Sr, Nd, and Pb isotope compositions. Samples were prepared and analyzed following the procedures of Kempton et al. (1995) and Royse et al. (1998). The Sr, Nd, and Pb isotope compositions were analyzed as metal species on single Ta, double Re—Ta and single Re filaments, respectively, using a Finnigan MAT 262 multi-collector mass spectrometer at the NERC Isotope Geosciences Laboratory (NIGL), Keyworth, UK. Samples were leached for ~1 h in 6 M HCl at 50 °C prior to digestion by a standard HF-HNO₃ procedure. Blanks for Sr, Nd, and Pb were less than 125 pg, 275 pg, and 325 pg, respectively. $^{87}\text{Sr}/^{86}\text{Sr}$ was normalized during run time to $^{87}\text{Sr}/^{86}\text{Sr} = 0.1194$ and $^{143}\text{Nd}/^{144}\text{Nd}$ was normalized to a value of $^{143}\text{Nd}/^{144}\text{Nd} = 0.7219$. Sample data are reported relative to accepted values of 0.710177 for $^{87}\text{Sr}/^{86}\text{Sr}$ in the NBS987 standard and 0.512128 for $^{143}\text{Nd}/^{144}\text{Nd}$ in the La Jolla standard. Based on repeated runs of NBS981, the reproducibility of the Pb isotope ratios was ~0.1%. Pb isotopes were corrected relative to the average standard Pb isotopic compositions of Todt et al. (1996). Measured values were age corrected based on the new $^{40}\text{Ar}/^{39}\text{Ar}$.

4. Results

4.1. Petrography

According to field and microscopic observations, the mafic and intermediate volcanic rocks from the SSMC (Fig. 4a-b; Table S1) are characterized by porphyritic textures. The main mineral assemblages represent euhedral-subhedral phenocrysts of plagioclase (10–64%), subhedral clinopyroxene (4–20%), subhedral-anhedral orthopyroxene (1–6%) and a lesser amount of subhedral-anhedral olivine (3–8%). The phenocrysts range in size from 0.1 to 0.5 mm, in a vitreous matrix (Fig. 4a-b; Table S1). Additionally, the rocks display the following characteristics: (a) diverse disequilibrium textures that include normal and sieved plagioclases crystals in the same rock samples; (b) rounded and embayed crystals of quartz with reaction rims of microcrysts of pyroxene; and (c) plagioclase with complex mineral zoning, such as reverse and oscillatory zoning, or normally zoned crystals, in the same rock sample (Fig. 4a-b).

In contrast, the silicic volcanic rocks display porphyritic textures with mineral assemblages that consist of phenocrysts of anhedral quartz (15–46%), euhedral to subhedral sanidine (10–22%), euhedral to subhedral plagioclase (2–43%), subhedral orthopyroxene (1–18%) and subhedral clinopyroxene (3–35%) embedded in a vitreous matrix showing a phenocryst size ranging from 0.2–0.7 mm of diameter (Fig. 4c-g).

4.2. $^{40}\text{Ar}/^{39}\text{Ar}$ geochronology

A total of 12 samples were analyzed; four dates were obtained from whole rock fragments (mafic-intermediate volcanic rocks), and eight from sanidine crystals (silicic volcanic rocks). Table 4 presents the summary of the $^{40}\text{Ar}/^{39}\text{Ar}$ plateau and correlation dating results from the SSMC. Spectra and correlation diagrams are shown in Fig. 8a–n, whereas step-heating results are reported in Table S3. Errors on the age spectrum and correlation diagrams represent the analytical precision at $\pm 2\sigma$ level.

The sanidine phenocrysts of rhyolite sample SLP17–11 (San Miguelito volcanic unit) show a spectrum with argon-loss defined by steps 1–3. A slightly flat age spectrum is defined by steps 4–6 with a plateau age of 33.17 ± 0.67 Ma (Fig. 5a). Moreover, this sample yielded an isochron age of 34.80 ± 5.4 Ma (Fig. 5b) with MSWD = 13, due to the spread of the error of the isochron data of this sample; therefore the isochron age should not be considered reliable. We use the more robust plateau age instead.

Sanidine phenocrysts of the San José unit (sample SLP17–29) yielded a slightly disturbed spectrum with a plateau age of 33.95 ± 0.44 Ma (Fig. 5c) over fourteen steps with 98.1% of released ^{39}Ar . Similarly, this sample yielded an isochron age of 34.09 ± 0.48 Ma (Fig. 5d) with a MSWD = 1.7. The isochron indicates that some presence of excess argon, which is also supported by the trend of the age spectrum.

^{40}Ar – ^{39}Ar analyses of sanidine phenocrysts from the rhyolitic sample SLP17–28 (El Zapote volcanic unit) produced a nearly ideal flat age spectrum, with some low ages at 11–14 steps. Steps 8–14 define a plateau age of 33.48 ± 0.43 Ma (Fig. 5e) with 90.2% of released ^{39}Ar . This sample yielded an isochron age of 33.40 ± 0.68 Ma (Fig. 5f) with MSWD = 2.2. The isochron age is nearly correlated to the plateau age, lending confidence to the result.

Analyses of sanidine crystals from the Cantera unit (sample SLP17–24) shows a nearly flat spectrum with exception of some slightly higher ages on the first two and the last three steps, and a lower age in step 8. Step 3–7 yielded a flat spectrum age of 32.10 ± 0.38 Ma (Fig. 5g) with 86% of released ^{39}Ar . The isochron for this samples yielded an age of 32.50 ± 0.39 Ma (Fig. 5h) with MSWD = 1.3. The isochron age and plateau age are nearly correlated, hence, both ages can be considered reliable.

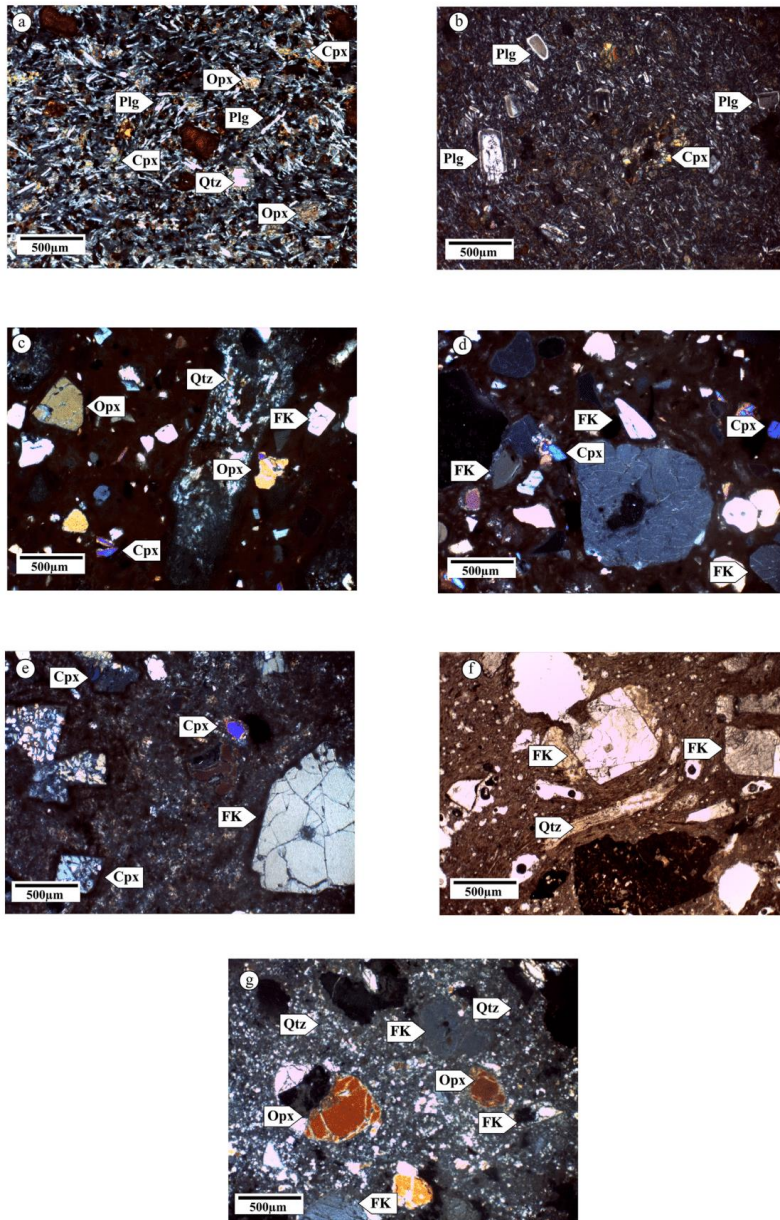


Fig. 4. Microphotographs of the SSMC volcanic rocks. (a) Phenocrysts of plagioclase, orthopyroxene, clinopyroxene and quartz with reaction rims from the Cabras unit; (b) Phenocrysts of plagioclase with sieved textures, microcrystals of clinopyroxene from the Placa unit; (c) Phenocrysts of potassium feldspar, orthopyroxene, clinopyroxene and microcrystals of quartz from the Panalillo unit; (d) Phenocrysts of potassium feldspar, orthopyroxene and clinopyroxene from the Cantera unit; (e) Phenocrysts of potassium feldspar and clinopyroxene from El Zapote unit; (f) Phenocrysts of potassium feldspar and bands filled with quartz from the San José unit; (g) Phenocrysts of potassium feldspar, orthopyroxene with microcrystals of quartz from the San Miguelito unit. Abbreviations of mineral names are as follows: FK-potassium feldspar; Plg-plagioclases; Opx-Orthopyroxene; Cpx-clinopyroxene.

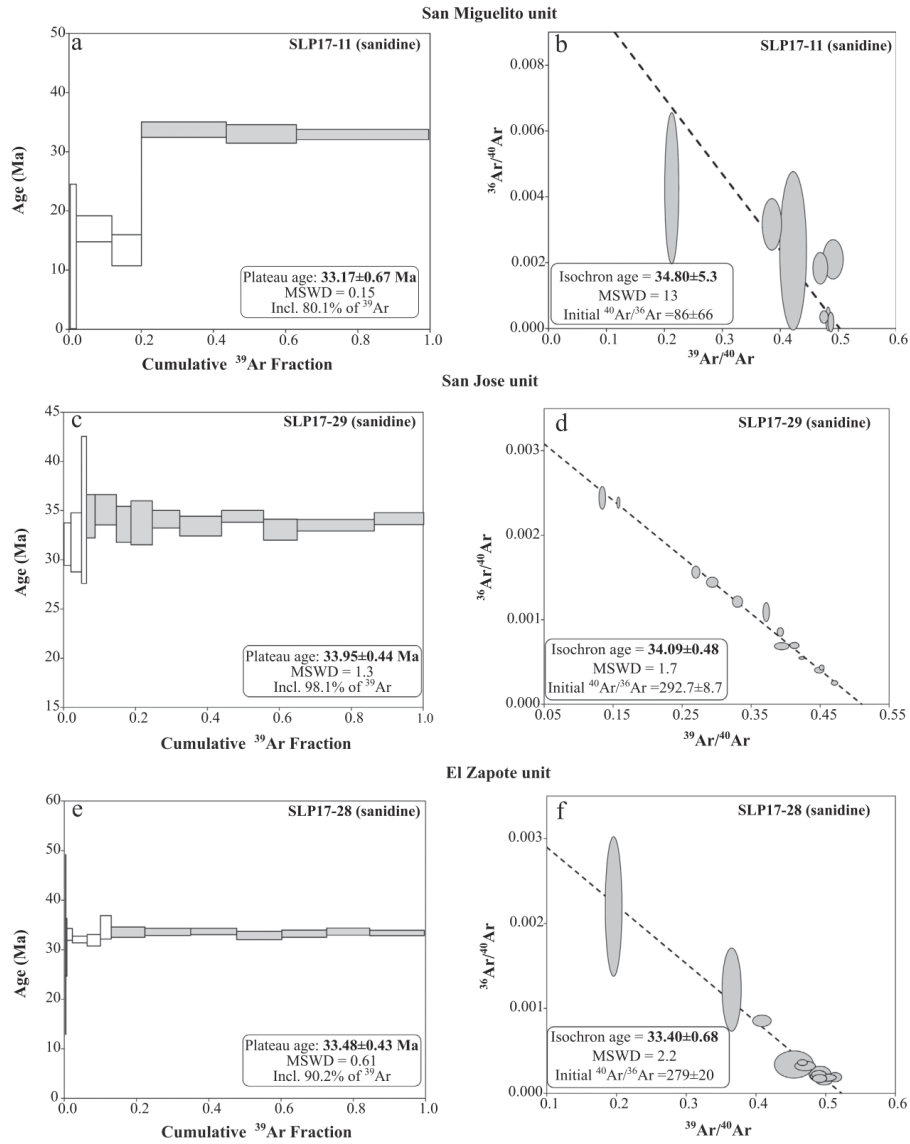


Fig. 5. $^{40}\text{Ar}/^{39}\text{Ar}$ age spectrum and respective isotope isochron diagram for the volcanic rocks. See text for discussion.

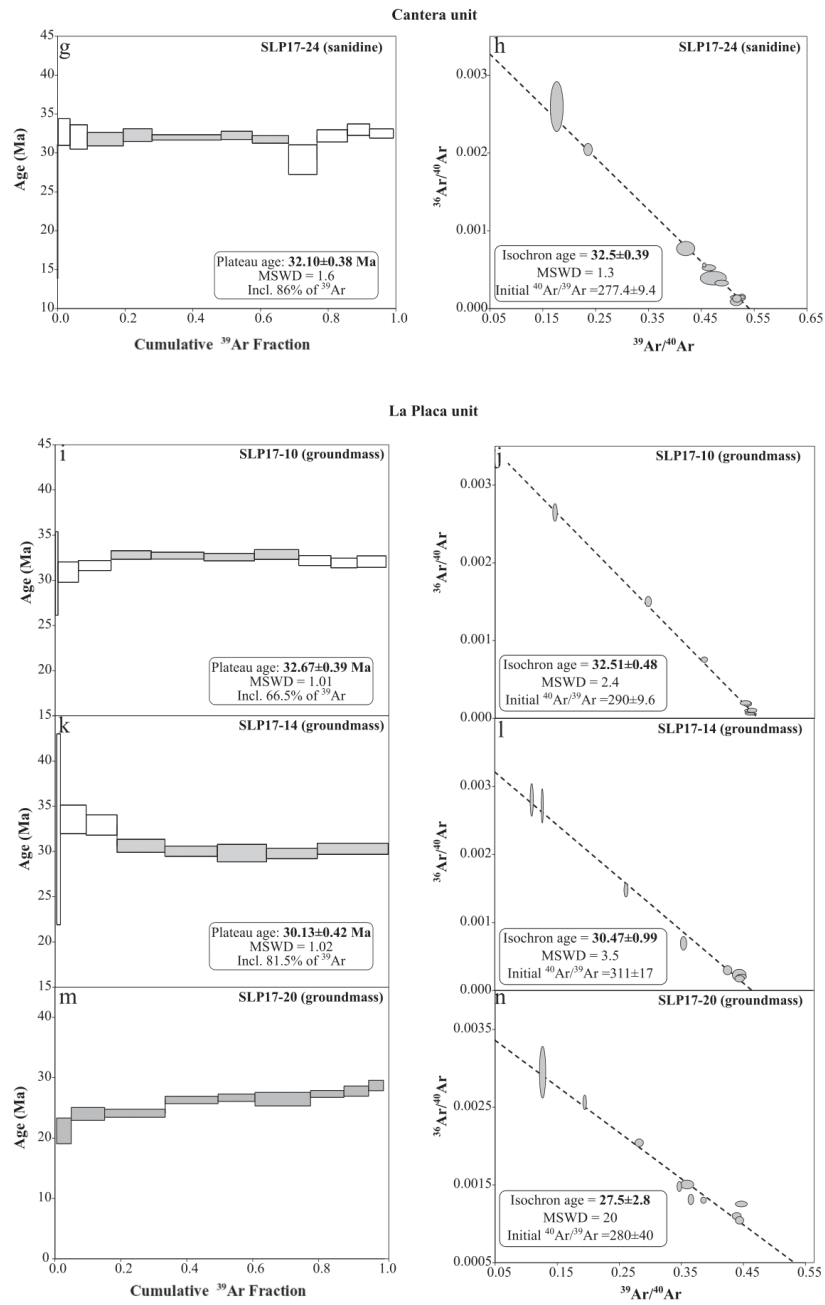


Fig. 5 (continued).

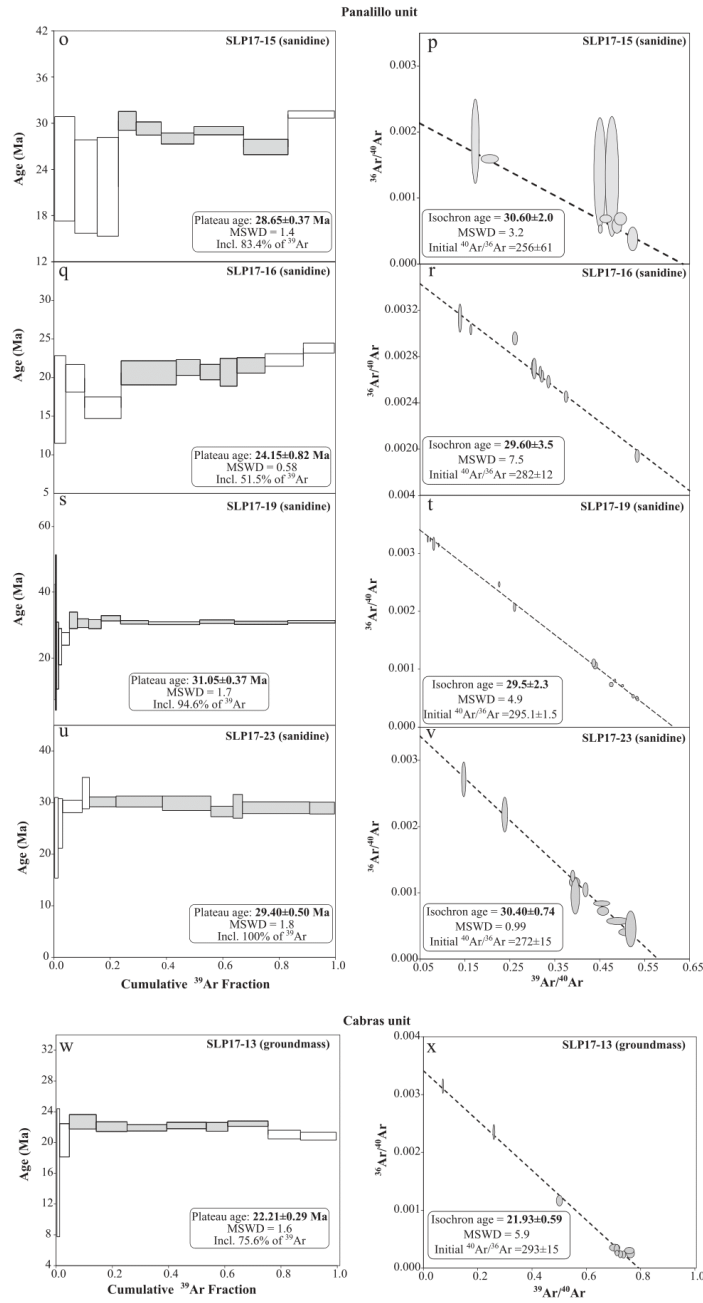


Fig. 5 (continued).

Groundmass from andesite and basalt samples of La Placa unit (SLP17–10; SLP17–14; SLP17–20) yielded nearly flat spectrum with some lower ages in the initial steps. For this unit, the spectrum ages yielded a range between 32.67 ± 0.39 and 30.13 ± 0.42 Ma (Fig. 5i, k, m), with ~66–81% of released ^{39}Ar . The isochrons for this unit show a range ages between 27.00 and 32.67 ± 0.42 Ma (Fig. 5j, l, n) with ~3.5–20 MSWD. The isochrons indicate some excess argon, which may be related to the age spectrum form. However, the concordance between the plateau and the isochron ages gives a confidence to the age of this unit.

^{40}Ar – ^{39}Ar analyses from sanidine of the samples SLP17–15 and SLP17–16 (Panalillo unit) shows a disturbed spectrum with a trend of argon loss profile with a plateau age of 28.65 ± 0.37 Ma and 24.15 ± 0.82 Ma (Fig. 5o, q, s, u) respectively and a ^{39}Ar release of 83.4 and 51.5%, with an isochron age of 30.60 ± 2.0 Ma and 29.60 ± 3.5 Ma (Fig. 5p, r, t, v). However, for the same unit, the samples SLP17–19 and SLP17–23 yielded a plateau age of 31.05 ± 0.37 Ma and 29.40 ± 0.50 Ma, with 94.6 and 100% of ^{39}Ar released respectively; these samples show an ideal flat age spectrum. The isochron of the samples SLP17–19 and SLP17–23 presents an age of 29.50 ± 2.30 Ma with MSWD = 4.9 and 30.40 ± 0.74 Ma with MSWD = 0.99, respectively.

The groundmass of SLP17–13 sample (Cabras unit; Fig. 5w) yielded a plateau age of 22.21 ± 0.29 Ma with 75.6% of ^{39}Ar released. The spectrum for this sample is slightly saddle-shaped. The isochron shows an isochron age of 21.93 ± 0.59 Ma (Fig. 5x) with MSWD = 5.9.

4.3. Whole-rock geochemical compositions

New major and trace element geochemical data for 15 samples from the SSMC are reported in Table 1. Besides 15 samples, additional literature geochemical data for 68 samples (Table S2 in the Supplementary Excel file) from the SSMC were compiled from the following references: Orozco-Esquivel et al. (2002, n = 4 samples); Leroy et al. (2002, n = 4 samples); Rodríguez-Ríos and Torres Aguilera (2009, n = 15 samples); Torres-Hernández et al. (2014, n = 8 samples); Aguillón-Robles et al. (2014, n = 14 samples); and Torres-Sánchez et al. (2019, n = 23 samples).

All data were processed in the IgRoCS software (Verma and Rivera-Gómez, 2013) to automatically determine the magma and rock types under the Middlemost (1989) option for Fe-oxidation adjustment, which allowed us to strictly follow the IUGS recommendations for rock classification and nomenclature (Le Bas et al., 1986). Thus, all major element data were treated in exactly the same manner. The use of 100% adjusted data on an anhydrous basis and after Fe-oxidation adjustment helps minimize the effect of analytical errors and element mobility and makes the use of the TAS diagram more consistent with the IUGS scheme.

The new data from the SSMC combined with the literature data (Orozco-Esquivel et al., 2002; Leroy et al., 2002; Rodríguez-Ríos and Torres Aguilera, 2009; Torres-Hernández et al., 2014; Aguillón-Robles et al., 2014; and Torres-Sánchez et al., 2019), are presented in the conventional TAS diagram (Fig. 6a; Le Bas et al., 1986). The samples plot in a wide compositional range including basalt, trachybasalt, basaltic andesite, andesite and rhyolite (Fig. 6a). These samples show a range of calc-alkaline to shoshonitic affinity (Fig. 6b), although caution is required in the use of a diagram without having Ca in it (Sheth et al., 2002).

From the new data (Table 1), we summarize the following characteristics. The mafic rock samples record values of $(\text{SiO}_2)_{\text{adj}} = 48.33$ – 51.48% m/m, $(\text{MgO})_{\text{adj}} = 5.71$ – 6.26% m/m, $(\text{TiO}_2)_{\text{adj}} = 1.97$ – 2.86% m/m, and Mg-number (Mg#) = 48.99–55.24. The Cr and Ni contents are in the ranges of 92.02–234.35 $\mu\text{g/g}$ and 3.50–83.25 $\mu\text{g/g}$, respectively. The intermediate rocks display more variable contents of $(\text{SiO}_2)_{\text{adj}} = 57.46$ – 62.24% m/m, $(\text{MgO})_{\text{adj}} = 2.81$ – 5.71% m/m, $(\text{TiO}_2)_{\text{adj}} = 0.88$ – 1.50% m/m. Their Mg# ranges from 53.04 to 64.13, Cr from 20.72 to 239.0 $\mu\text{g/g}$, Ni from 3.5 to 42.7 $\mu\text{g/g}$. The silicic rocks

have high $(\text{SiO}_2)_{\text{adj}} = 76.56$ – 82.24% m/m, $(\text{MgO})_{\text{adj}} = 0.03$ – 0.18% m/m, $(\text{TiO}_2)_{\text{adj}} = 0.59$ – 0.24% m/m, and Mg# = 8.42–27.61, with Cr and Ni contents are in the ranges of 11.36–34.82 and 0.9–3.6 $\mu\text{g/g}$, respectively.

The chondrite-normalized rare earth element (REE; McDonough and Sun, 1995) plots of the new data (Table 1) are shown in Fig. 7a–c. The mafic volcanic rocks display a relatively flat pattern with a slight enrichment in light REE (LREE; $(\text{La}/\text{Sm})_{\text{CN}} = 1.91$ – 2.05 ; $(\text{La}/\text{Yb})_{\text{CN}} = 3.76$ – 6.44 ; Fig. 7a), with a small negative or positive Eu anomaly ($\text{Eu}/\text{Eu}^* = 1.03$ – 1.09). Total REE concentrations of the mafic volcanic rocks vary from about 101.66 to 152.84 $\mu\text{g/g}$. The intermediate volcanic rocks show total REE content ranging from about 125.5 to 181.3 $\mu\text{g/g}$ (one sample having very low REEs; Fig. 7b). The intermediate rocks also display enrichment in LREE (Fig. 6b; $(\text{La}/\text{Sm})_{\text{CN}} = 2.02$ – 2.43 ; $(\text{La}/\text{Yb})_{\text{CN}} = 5.13$ – 6.41), but also show a relatively small negative Eu anomaly ($\text{Eu}/\text{Eu}^* = 0.12$ – 0.82). The silicic volcanic rocks display total REE content of 37.45–439.75 $\mu\text{g/g}$, enriched LREE patterns (Fig. 7c; $(\text{La}/\text{Sm})_{\text{CN}} = 1.21$ – 5.82 ; $(\text{La}/\text{Yb})_{\text{CN}} = 1.16$ – 13.10) but also show large negative Eu anomalies ($\text{Eu}/\text{Eu}^* = 0.03$ – 0.36).

Primitive mantle-normalized multi-element (McDonough and Sun, 1995) diagrams of mafic volcanic rocks from SSMC display almost flat patterns (Fig. 7d). In general, the mafic volcanic rocks are enriched in highly incompatible LILE (i.e., Rb, U, LREE) relative to the moderately incompatible HFSE (e.g., Nb, Ta, Th and HREE). Normalized trace element patterns are enriched in incompatible elements with low Nb anomalies ($\text{Nb}/\text{Nb}^* = 0.64$). The intermediate rocks display enrichment in LILE, relative to the incompatible HFS (Fig. 7e) and enrichment in incompatible elements with low Nb ($\text{Nb}/\text{Nb}^* = 0.22$ – 0.33), Ti ($\text{Ti}/\text{Ti}^* = 0.39$ – 0.51), Ba and P anomalies. The silicic volcanic rocks from the SSMC are enriched in incompatible LILE compared to incompatible HFS elements (Fig. 7f). Normalized patterns of silicic rocks display an enrichment in incompatible elements with low Nb anomalies ($\text{Nb}/\text{Nb}^* = 0.21$ – 35.38) and strong Ti ($\text{Ti}/\text{Ti}^* = 0.01$ – 0.13), Ba, Sr, and P anomalies.

The Nb and Ti anomalies are important to understanding the crustal contamination because mafic rocks from extensional or rift tectonic settings have lesser anomalies than intermediate and silicic rocks (Verma, 2020a), which shows that mafic magmas are likely to have lesser crustal assimilation effects as compared to intermediate and silicic rocks. Intermediate and silicic rocks from the SSMC are likely to have more residence time in the crust as compared to mafic rocks. The lesser crustal residence time is especially true for relatively primary mafic magmas having high MgO, Mg#, Ni, and Cr.

4.4. Whole-rock Sr-Nd-Pb isotope data

Whole-rock Sr, Nd, Pb isotopic data for the SSMC volcanic rocks are listed in Table 2. Due to the scarcity of isotope information in the study area, literature data from the southern region of the BR were compiled and plotted in all isotope diagrams (Fig. 8a–d). Isotopic data from the BR were taken from: Duncker et al. (1991, n = 14); Johnson and Thompson (1991, n = 17); Kempton et al. (1991, n = 35); Gibson et al. (1992, n = 20); Bradshaw et al. (1993, n = 14); Davis and Hawkesworth (1995, n = 19); Rogers et al. (1995, n = 26); McMillan et al. (2000, n = 26) and Christiansen et al. (2007, n = 8).

Strontium-Nd and initial $\epsilon_{\text{Nd}}(t)$ values for SSMC volcanic rocks are plotted in Fig. 8a, together with isotope data from the BR volcanic rocks. The mafic volcanic rocks of the SSMC display initial $^{87}\text{Sr}/^{86}\text{Sr}$ isotopic ratios (I_{Sr}) of 0.70344 and 0.70496 and positive $\epsilon_{\text{Nd}}(t)$ values of +5.2 and +0.9 (Table 2). The intermediate volcanic rocks from the SSMC show I_{Sr} that range from 0.70540 to 0.70581 and negative $\epsilon_{\text{Nd}}(t)$ values of –1.8 to –2.5 (Table 2). For silicic volcanic rocks the I_{Sr} ratios range from 0.70719 to 0.71973 and negative $\epsilon_{\text{Nd}}(t)$ values ranging from –0.2 to –2.7 (Table 2). The mafic volcanic rocks plot within the mantle array and overlap the mafic BR volcanic rocks. Intermediate-silicic volcanic rocks plot below the bulk earth line, which suggests

Table 1

Representative whole rock compositions of volcanic rocks from the Sierra de San Miguelito Volcanic Complex, San Luis Potosí, Mexico (major elements in weight percent (also known as % m/m) and trace and rare earth elements in µg/g).

Sample Rock unit	SLP17-11 San Miguelito	SLP17-12	SLP17-29 San José	SLP17-28 El Zapote	SLP17-24 Cantera	SLP17-10 La Placa	SLP17-14	SLP17-20	SMB17-01	SMB17-02
Rock (TAS)	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Andesite	Basalt	Andesite	Andesite	Andesite
Long. (°W)	101°14'06"	101°10'51"	101°05'44"	101°05'45"	101°06'34"	101°09'23"	101°02'11"	101°05'43"	101°09'24"	101°09'2"
Lat. (°N)	22°06'23"	22°00'03"	22°01'23"	22°05'14"	22°05'26"	22°05'09"	21°46'46"	21°49'48"	22°05'10"	22°05'07"
SiO ₂	77.63	76.53	79.15	77.07	78.01	62.88	51.23	56.89	61.86	61.27
TiO ₂	0.13	0.06	0.11	0.11	0.10	0.89	1.96	1.49	0.91	0.92
Al ₂ O ₃	12.57	13.75	11.54	12.69	12.20	16.73	16.34	15.79	15.57	16.12
Fe ₂ O ₃	1.39	1.34	1.25	1.24	1.28	6.55	11.80	8.23	6.78	7.02
MnO	0.02	0.02	0.01	0.01	0.01	0.11	0.17	0.14	0.11	0.12
MgO	0.18	0.07	0.04	0.07	0.17	2.84	6.23	5.65	3.25	3.07
CaO	0.45	0.29	0.37	0.37	0.38	5.46	8.84	6.94	5.56	6.18
Na ₂ O	2.26	3.41	2.08	2.44	1.59	2.70	2.73	2.41	2.66	2.54
K ₂ O	5.43	4.57	5.12	5.44	5.38	3.15	0.94	1.71	3.11	2.89
P ₂ O ₅	0.02	0.01	0.02	0.01	0.01	0.22	0.28	0.39	0.23	0.23
LOI	0.63	0.76	0.96	0.89	1.23	0.53	0.31	1.19	0.60	0.76
Sum	100.70	100.81	100.65	100.35	100.37	102.07	100.84	100.83	100.65	101.12
Q	42.06	38.01	46.42	41.02	47.14	17.35	1.50	12.30	16.80	16.60
Or	32.09	27.02	30.38	32.35	32.10	18.43	5.58	10.21	18.47	17.11
Ab	19.13	28.87	17.67	20.78	13.58	22.61	23.21	20.60	22.62	21.53
An	2.10	1.37	1.71	1.78	1.84	23.98	29.70	27.49	21.46	24.09
Ne	2.20	2.69	1.96	2.15	3.12					
C	1.50	1.30	1.04	1.11	1.42	12.04	22.71	18.95	12.14	11.74
Di-Mg	0.63	0.60	0.56	0.56	0.58	2.25	2.62	2.89	2.37	2.44
Di-Fe	0.25	0.11	0.21	0.21	0.19	1.67	3.74	2.86	1.74	1.75
Hy-Mg	0.05	0.02	0.05	0.02	0.02	0.50	0.65	0.91	0.54	0.53
Hy-Fe	27.11	13.05	8.42	13.95	27.61	53.04	55.24	64.14	55.53	53.25
Mt	6.95	17.22	28.12	15.94	6.78	2.08	1.70	1.31	1.88	2.06
Il	42.06	38.01	46.42	41.02	47.14	17.35	1.50	12.30	16.80	16.60
Ap	32.09	27.02	30.38	32.35	32.10	18.43	5.58	10.21	18.47	17.11
Mg#	19.13	28.87	17.67	20.78	13.58	22.61	23.21	20.60	22.62	21.53
FeO/MgO	2.10	1.37	1.71	1.78	1.84	23.98	29.70	27.49	21.46	24.09
Ba	197.987	3.993	84.426	56.488	93.939	553.860	243.011	815.147	541.437	561.683
Co	1.174	0.681	0.745	0.879	0.684	16.790	33.135	27.095	2.581	17.910
Cr	11.381	11.365	34.817	12.732	14.384	103.450	92.022	238.981	20.717	108.275
Cs	9.426	3.656	4.574	11.994	7.751	3.245	0.950	0.583	0.583	2.853
Cu	2.102	4.915	2.438	3.041	2.502	8.883	13.950	10.996	7.900	8.300
Ga	22.550	48.024	21.436	23.497	22.506	19.931	25.484	19.267	19.600	20.900
Hf	6.364	11.285	6.022	6.772	6.043	6.308	4.290	6.412	5.817	5.578
Nb	22.111	124.387	21.488	23.509	23.281	13.252	13.638	12.944	12.872	12.091
Ni	1.355	1.576	2.337	1.025	2.036	6.921	3.483	42.683	6.800	3.500
Pb	26.661	51.671	28.743	29.351	27.019	10.886	4.164	8.495	10.723	9.958
Rb	213.350	290.797	144.259	250.091	181.507	66.264	15.721	60.949	50.935	42.022
Sc	15.384	8.264	1.316	7.121	2.242	24.478	34.975	24.756	21.682	23.251
Sr	27.560	2.458	14.007	16.071	23.189	276.837	311.042	390.274	252.612	294.668
Th	28.959	44.084	27.782	30.295	25.802	9.562	2.751	7.037	9.073	8.010
U	4.414	6.664	5.752	4.297	5.005	2.766	0.599	1.242	2.721	2.653
V	7.029	2.210	16.231	1.605	2.670	91.165	140.290	116.680	105.382	99.617
Y	45.551	22.808	43.756	13.331	33.373	25.781	28.204	37.810	22.901	27.477
Zn	52.222	163.443	52.300	59.009	63.020	82.635	82.039	91.125	79.200	81.000
Zr	170.859	166.115	158.115	165.630	152.773	237.990	182.985	266.598	235.385	217.393
La	40.62	6.26	50.45	19.69	22.90	24.98	16.50	33.71	21.63	22.46
Ce	96.46	11.68	80.28	58.29	58.65	56.28	36.23	68.91	46.83	46.21
Pr	11.61	1.34	15.58	3.43	7.02	7.42	4.85	9.06	6.69	6.66
Nd	48.15	5.11	67.79	11.13	30.04	32.71	23.30	41.11	5.04	30.33
Sm	10.73	1.76	14.24	2.13	6.80	7.21	5.48	8.64	1.05	6.43
Eu	0.51	0.09	0.29	0.27	1.62	1.99	1.99	2.33	0.26	1.58
Gd	10.35	2.38	12.77	2.58	7.41	7.06	5.87	8.72	6.41	6.69
Tb	1.68	0.58	2.07	0.46	1.45	1.04	0.97	1.29	0.99	1.01
Dy	9.91	4.24	12.32	3.23	9.39	6.07	5.72	7.39	5.51	6.01
Ho	1.91	0.93	2.38	0.68	1.91	1.16	1.15	1.46	1.08	1.19
Er	5.59	3.01	6.81	2.25	5.52	3.30	3.30	4.25	3.14	3.50
Tm	0.69	0.44	0.75	0.29	0.62	0.34	0.39	0.51	0.28	0.33
Yb	5.01	3.68	5.89	2.75	5.14	3.00	2.99	3.57	2.86	3.07
Lu	0.75	0.55	0.88	0.41	0.78	0.45	0.46	0.58	0.45	0.47
TREE	229.90	37.45	254.17	103.42	146.27	144.05	101.66	181.26	125.46	127.95
(La/Yb) _{CN}	5.51	1.16	5.82	4.87	3.03	5.66	3.75	6.41	5.13	4.97
(La/Sm) _{CN}	2.47	2.18	2.26	5.82	2.04	2.21	1.91	2.43	2.02	2.14
(Gd/Yb) _{CN}	1.67	0.52	1.75	0.76	1.17	1.90	1.59	1.97	1.81	1.76
(La/Yb) _{PM}	5.52	1.16	5.83	4.88	3.03	5.67	3.76	6.42	5.14	4.98
(La/Sm) _{PM}	2.48	2.19	2.27	5.84	2.04	2.22	1.92	2.43	2.02	2.15
(Gd/Yb) _{PM}	1.68	0.53	1.76	0.76	1.17	1.91	1.59	1.98	1.82	1.76
(Eu/Eu*)	0.15	0.13	0.07	0.36	0.11	0.70	1.09	0.82	0.12	0.73
(Nb/Nb*)	0.70	35.38	0.69	1.76	1.37	0.32	0.64	0.22	0.33	0.29
(Ti/Ti*)	0.04	0.06	0.03	0.13	0.04	0.39	1.02	0.51	0.43	0.42

Sample Rock unit	SLP17–15 Panalillo	SLP17–16	SLP17–19	SLP17–23	SLP17–13 Cabras
Rock (TAS)	Rhyolite	Rhyolite	Rhyolite	Rhyolite	Basalt
Long. (°W)	101°02'13"	101°05'11"	101°06'08"	101°06'35"	101°10'50"
Lat. (°N)	21°46'47"	21°48'46"	21°49'02"	22°06'06"	21°55'17"
SiO ₂	75.93	82.28	76.62	78.25	47.76
TiO ₂	0.20	0.06	0.24	0.15	2.83
Al ₂ O ₃	12.24	9.74	11.99	11.23	15.59
Fe ₂ O ₃	2.43	0.92	2.34	1.66	13.73
MnO	0.03	0.02	0.02	0.01	0.20
MgO	0.12	0.08	0.03	0.02	5.64
CaO	1.13	0.51	0.44	0.30	8.87
Na ₂ O	2.53	2.55	2.51	2.61	3.39
K ₂ O	5.02	3.94	5.12	5.49	1.27
P ₂ O ₅	0.05	0.01	0.04	0.02	0.71
LOI	0.64	0.47	0.92	0.70	0.39
Sum	100.32	100.57	100.27	100.43	100.38
Q	38.88	50.82	41.11	41.03	0.00
Or	29.81	23.27	30.50	32.57	7.59
Ab	21.51	21.57	21.41	22.17	29.03
An	5.31	2.46	1.94	1.36	23.85
Ne	0.71	0.38	1.63	0.50	-
C	2.18	0.95	1.80	1.29	2.58
Di-Mg	1.10	0.41	1.06	0.75	3.07
Di-Fe	0.38	0.11	0.46	0.29	5.44
Hy-Mg	0.12	0.02	0.09	0.05	1.66
Hy-Fe	12.42	19.98	3.55	3.34	48.99
Mt	18.22	10.35	70.19	74.68	2.19
Il	38.88	50.82	41.11	41.03	0.00
Ap	29.81	23.27	30.50	32.57	7.59
Mg#	21.51	21.57	21.41	22.17	29.03
FeO ⁺ /MgO	5.31	2.46	1.94	1.36	23.85
Ba	518.985	23.390	499.326	37.639	304.315
Co	2.026	0.853	1.115	0.593	43.183
Cr	18.347	14.968	11.565	12.659	234.352
Cs	8.323	9.347	1.596	1.843	0.333
Cu	3.919	1.670	4.026	2.862	36.438
Ga	18.087	19.209	24.249	23.983	26.990
Hf	6.578	4.276	11.171	9.380	6.068
Nb	15.766	19.431	34.377	40.481	25.028
Ni	1.861	3.599	0.904	1.674	83.257
Pb	22.503	16.621	21.886	26.949	3.066
Rb	157.642	237.321	85.363	121.351	10.359
Sc	13.620	4.740	0.896	1.349	26.952
Sr	64.817	12.268	29.233	5.279	425.171
Th	23.395	36.907	17.217	25.063	2.130
U	5.197	8.373	4.350	5.222	0.740
V	8.874	32.621	9.936	37.868	183.768
Y	49.555	75.276	35.823	39.647	32.536
Zn	53.115	27.636	93.980	85.772	98.530
Zr	239.542	85.699	428.562	274.831	266.834
La	89.25	20.22	45.42	46.26	26.45
Ce	195.11	52.90	104.44	98.96	55.06
Pr	22.48	7.46	14.60	14.99	7.73
Nd	91.68	34.91	64.06	64.10	37.28
Sm	16.08	10.37	12.40	12.52	8.18
Eu	1.23	0.11	0.94	0.16	2.75
Gd	15.27	11.60	11.87	11.87	8.42
Tb	2.00	2.30	1.86	1.91	1.24
Dy	11.05	14.47	10.65	11.44	7.00
Ho	1.99	2.98	2.06	2.26	1.30
Er	5.57	8.94	5.95	6.63	3.47
Tm	0.60	1.08	0.63	0.69	0.38
Yb	4.63	8.56	5.35	6.17	2.79
Lu	0.70	1.24	0.82	0.92	0.45
TREE	439.75	159.82	265.58	262.77	152.84
(La/Yb) _{CN}	3.59	1.21	2.24	2.29	2.05
(La/Sm) _{CN}	2.67	1.10	1.79	1.56	2.44
(Gd/Yb) _{CN}	13.13	1.61	5.78	5.11	6.44
(La/Yb) _{PM}	3.60	1.21	2.25	2.30	2.06
(La/Sm) _{PM}	2.68	1.10	1.80	1.56	2.44
(Gd/Yb) _{PM}	0.25	0.03	0.24	0.04	1.03
(Eu/Eu ⁺)	0.21	1.63	0.69	1.53	0.84
(Nb/Nb ⁺)	0.04	0.01	0.06	0.04	1.07
(Ti/Ti ⁺)	13.10	1.60	5.77	5.10	6.43

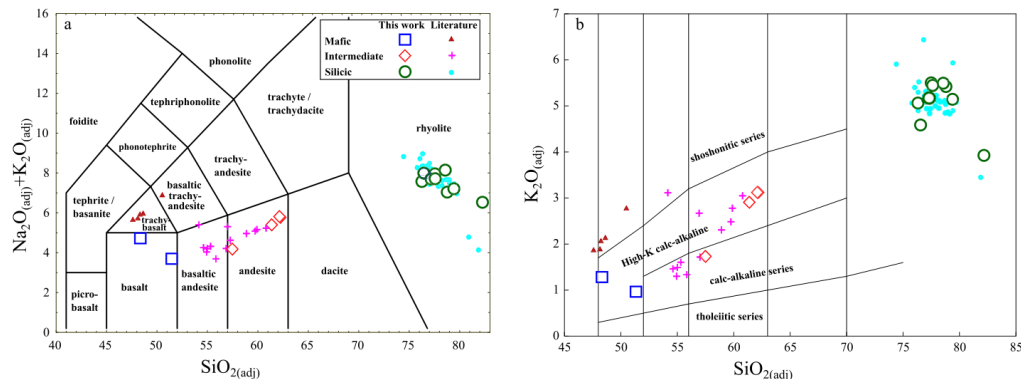


Fig. 6. (a) Total Alkali–Silica classification diagram for the volcanic rocks from the SSMC (TAS; Le Bas et al., 1986); (b) $K_2O_{(adj)}$ – $SiO_{2(adj)}$ diagram of Peccerillo and Taylor (1976). Literature data were taken from the following sources: Orozco-Esquivel et al. (2002); Leroy et al. (2002); Rodríguez-Ríos and Torres-Aguilera (2009); Torres-Hernández et al. (2014); Aguillón-Robles et al. (2014); and Torres-Sánchez et al. (2019).

these rocks may have originated from partial melting of the continental crust.

Lead isotopic compositions of the SSMC volcanic rocks are summarized in Table 3. The rocks are plotted in Fig. 8b, together with the BR isotopic data. The graph also depicts the Northern Hemisphere reference line (NHRL; Zindler and Hart, 1986). The variations in Pb isotopic ratios for the mafic volcanic rocks are 18.925 and 18.962 for $(^{206}Pb/^{204}Pb)_i$, 15.303 and 15.329 for $(^{207}Pb/^{204}Pb)_i$, and 38.571 and 38.909 for $(^{208}Pb/^{204}Pb)_i$ (Table 3). For intermediate rocks, the initial values for $(^{206}Pb/^{204}Pb)_i$, $(^{207}Pb/^{204}Pb)_i$, $(^{208}Pb/^{204}Pb)_i$ ratios show a range of 18.891–18.934, 15.149–15.248, 38.864–38.902 (Table 3), respectively. The silicic volcanic rocks from the SSMC display $(^{206}Pb/^{204}Pb)_i$, $(^{207}Pb/^{204}Pb)_i$, $(^{208}Pb/^{204}Pb)_i$ ratios between 18.889 and 19.005, 14.521–15.217, and 38.831–39.016 (Table 3), respectively. In the Pb isotope diagrams (Fig. 8b), the SSMC mafic volcanic rocks plot above the NHRL with a trend towards depleted mantle (DM) just as the mafic rocks from the BR. The intermediate and silicic volcanic rocks from the SSMC partially overlap and plot above the NHRL with a trend towards enriched mantle type II (EMII) like the silicic volcanic rocks from the BR.

5. Discussion

5.1. Implications of $^{40}Ar/^{39}Ar$ data

For several years, it has been shown that the K–Ar method presents a major disadvantage over the Ar–Ar method. Compared with K/Ar dating, the $^{40}Ar/^{39}Ar$ technique has the advantage of requiring a much lower quantity of sample and uses only a single sample aliquot for analysis (Jourdan, 2012). Consequently, K–Ar ages tend to have much larger uncertainties than Ar–Ar ages (Lee, 2015). Therefore, when comparing $^{40}Ar/^{39}Ar$ ages with ages obtained by another method, it should be kept in mind that only the total uncertainty should be considered (Couil e et al., 2003).

A database of K–Ar ages have been compiled from the SSMC (Table S3). Data were compiled from: a) Labarthe-Hern andez et al. (1982; $n = 6$); b) Tuta et al. (1988; $n = 1$); c) Aguill on-Robles et al. (1994; $n = 1$); d) Aguill on-Robles et al. (2009; $n = 1$); e) Trist an-Gonz alez et al. (2009; $n = 23$); f) Aguill on-Robles et al. (2014; $n = 4$). New $^{40}Ar/^{39}Ar$ radiometric data demonstrate that volcanism in the SSMC starts from late Eocene, about 34 Ma, with the emplacement of the San Miguelito volcanic unit. The youngest age is obtained from the

Cabras volcanic unit (about 21.9 Ma), indicating that the volcanism of the SSMC was active until the late Miocene.

The new $^{40}Ar/^{39}Ar$ data display three possible phases of magmatism in the SSMC (Fig. 9). The first phase is characterized by the generation of rhyolitic domes (San Miguelito and El Zapote unit); these domes were accompanied with effusive events of rhyolitic ignimbrites (San Jos e and Cantera unit). The onset of the magmatic activity is constrained by Ar–Ar age of about 34 Ma (Fig. 9). The second phase, defined as re-activation of magmatic sources, is occupied by the eruption of intermediate volcanic rocks (La Placa unit), followed by, effusive magmatism of rhyolitic ignimbrites, between ~32.5 and 29.5 Ma (Panalillo unit). The third and final phase is characterized by emplacement of mafic volcanic rocks (Cabras unit), which represent a last period of activity, at about 22 Ma; this age is well constrained by $^{40}Ar/^{39}Ar$ ages, however, K–Ar shows high levels of uncertainty (Fig. 9).

The evidence of these three main phases observed in the SSMC leads to important observations of the evolution of the southern region of the MC: (1) high volumes of rhyolite ignimbrite volcanism between the late Eocene and the middle Oligocene; (2) the southern region of MC is comprised of three different compositional volcanic activities since middle Oligocene to early Miocene, with SiO_2 varying widely from about 37.4% m/m to 79.9% m/m (Table S2). The bimodal volcanism, as proposed previously (Rodr guez-R os and Torres-Aguilera, 2009; Aguill on-Robles et al., 2014), does not seem to be valid.

5.2. Petrogenetic implications

Since the negative Nb anomaly in the volcanic rocks likely reflects a source, (Nb/Yb) vs. (Th/Yb) (Fig. 10a) can be used to investigate source characteristics (Pearce, 2008). In the diagram, it can be inferred that the mafic sample (SLP17–13) from the SSMC is mantle derived, while SLP17–14 (mafic rock) displays a lower continental crust signature; in contrast, the intermediate and silicic volcanic rocks of the SSMC display an interaction with middle and upper continental crust.

In order to explore the evolution of the magmatic rocks of the SSMC, petrogenetic mechanisms have been tentatively quantified using different methods. These different models aim at testing the relationship between the more evolved magma and the less differentiated magmas of the SSMC.

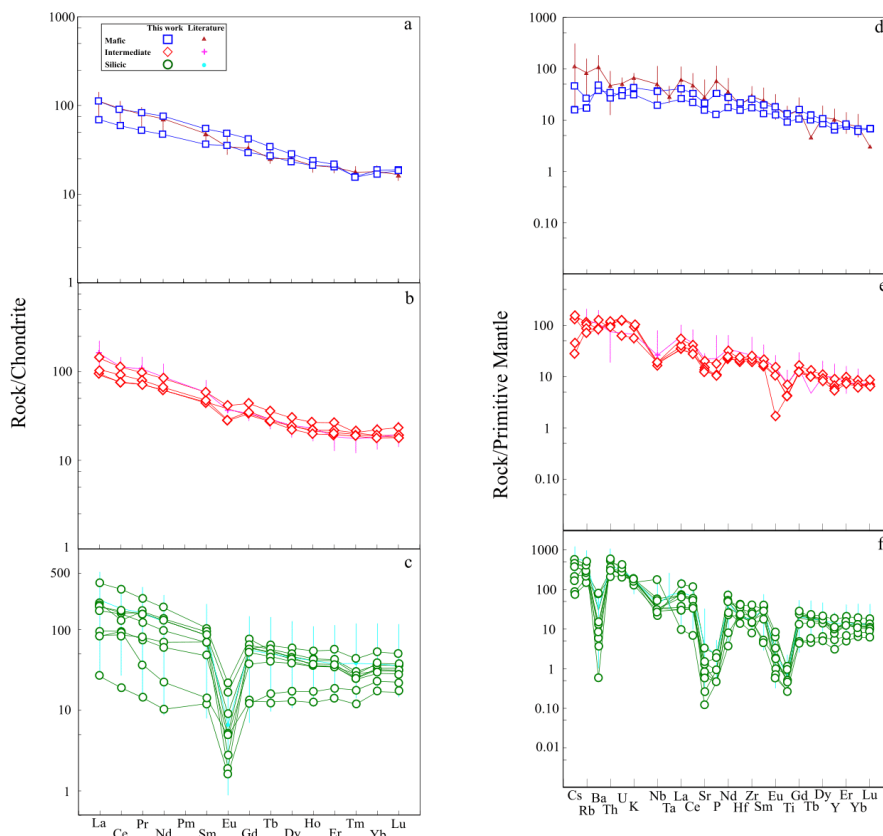


Fig. 7. (a–c) Chondrite-normalized rare-earth element (REE) diagrams for the SSMC rocks (chondrite values for normalization taken from McDonough and Sun, 1995); (d–f) Primitive-mantle normalized multi-element diagrams for the SSMC rocks (primitive mantle values for normalization taken from McDonough and Sun, 1995). Symbols are same as in Fig. 5. Literature data were taken from the following sources: Orozco-Esquivel et al. (2002); Leroy et al. (2002); Rodríguez-Ríos and Torres-Aguilera (2009); Torres-Hernández et al. (2014); Aguilón-Robles et al. (2014) and Torres-Sánchez et al. (2019).

5.2.1. Origin of the mafic volcanic rocks

Thus far, the origin of the mafic volcanic rocks from the SSMC has been poorly discussed. Some authors described that these types of rocks in the area have presented a process of magma mixing during their rise to the surface (e.g., Rodríguez-Ríos and Torres-Aguilera, 2007; Torres-Sánchez et al., 2019). Therefore, it is worthwhile to explore the petrogenetic possibilities for these volcanic rocks.

The mafic samples (SLP17–13 and SLP17–14) from the SSMC plot nearby in the $\text{SiO}_2(\text{adj})$ vs $^{87}\text{Sr}/^{86}\text{Sr}_{(i)}$ diagram (Fig. 10b), although the sample SLP17–14 is shifted towards the mantle-crust mixing lines. A small progressive signature of $^{87}\text{Sr}/^{86}\text{Sr}_{(i)}$ with $\text{SiO}_2(\text{adj})$ from the sample SLP17–13 towards SLP17–14 may suggest an AFC process. Moreover, sample SLP17–14 could indicate a possible consequence of mixing between mantle-derived magma and crustal melt. This trend converges to a mantle-derived magma composition with an appropriate $^{87}\text{Sr}/^{86}\text{Sr}_{(i)}$ isotope ratio (0.70344–0.70496; Table 2). On the other hand, this may indicate derivation of the mafic samples (SLP17–13 and SLP17–14) from a heterogeneous mantle source.

In this context, an additional model has been tested. The initial Sr-Nd-Pb isotope ratios are useful to distinguish both mantle source and crustal involvement during magmatism. However, the $^{206}\text{Pb}/^{204}\text{Pb}$ ratio effectively distinguishes between the various mantle reservoirs, whereas the other isotope values can be used to infer the crustal component (Harangi et al., 2007). Therefore, following the model proposed by Harangi et al. (2007) (Fig. 10c), it can be interpreted that mafic samples from SSMC show two different sources.

The sample SLP17–13 shows a trend to an enriched-mid ocean ridge basalt (*E*-MORB) mantle without any style of mixing process (Fig. 10c). On the other hand, SLP17–14 yields a trend towards a mantle-lower crust mixing line (Fig. 10c), which may suggest a small crustal component in the genesis of this sample.

To support the idea of two different sources in the mafic rocks from the SSMC, a partial melting model of the mantle has been developed following the batch model equations reported by Zou (2007). Data of lherzolite xenolith (sample SD7; Dávalos-Elizondo et al., 2016) were used to represent lithospheric mantle composition. A mineralogical

Table 2

Whole-rock Sr and Nd isotopic data for volcanic rocks from the Sierra de San Miguelito Complex. (age data for in situ-growth corrections were taken from Table 4). For ϵ_{Nd} , CHUR values for $^{143}Nd/^{144}Nd = 0.512630$ and $^{147}Sm/^{144}Nd = 0.1960$ were taken from Faure (1977). Rb, Sr, Sm and Nd concentrations were taken from Table 1).

Sample	Rock unit	Magma type	Age	Rb (ppm)	Sr (ppm)	$^{87}Rb/^{86}Sr$	$^{87}Sr/^{86}Sr$	$\pm 2\sigma$	$^{87}Sr/^{86}Sr$ (i)	Sm (ppm)	Nd (ppm)	$^{147}Sm/^{144}Nd$	$^{143}Nd/^{144}Nd$	$\pm 2\sigma$	$^{143}Nd/^{144}Nd$ (i)	ϵ_{Nd}
SLP17-13	Cabras	Basic	22.21	10.36	425.17	0.07049	0.70346	6	0.70344	8.18	37.28	0.13271	0.51289	10	0.51287	+5.2
SLP17-14	La Placa	Basic	30.13	15.72	311.04	0.14622	0.70502	6	0.70496	5.48	23.30	0.14210	0.51266	18	0.51264	+0.9
SLP17-10	La Placa	Inter.	32.67	66.26	276.84	0.69248	–	–	–	7.21	32.71	0.13330	0.51249	9	0.51246	-2.5
SLP17-20	La Placa	Inter.	27.50	60.95	390.27	0.45181	0.70557	6	0.70540	8.64	41.11	0.12707	0.51252	7	0.51249	-2.0
SMB17-01	La Placa	Inter.	30.00	50.94	252.61	0.58333	0.70606	7	0.70581	1.05	5.04	0.12628	0.51249	13	0.51247	-2.4
SMB17-02	La Placa	Inter.	30.00	42.02	294.67	0.41257	0.70592	20	0.70575	6.43	30.33	0.12822	0.51252	7	0.51249	-1.8
SLP17-11	San Miguelito	Acid	33.17	213.35	27.56	22.39592	0.71995	7	0.70940	10.73	48.15	0.13474	0.51249	6	0.51246	-2.5
SLP17-12	San Miguelito	Acid	33.00	290.80	2.46	342.26587	–	–	–	1.76	5.11	0.20808	0.51262	8	0.51258	-0.2
SLP17-15	Panalillo	Acid	28.65	157.64	64.82	7.03621	0.71005	6	0.70719	16.08	91.68	0.10608	0.51248	8	0.51245	-2.7
SLP17-16	Panalillo	Acid	28.65	237.32	12.27	55.96519	0.73354	7	0.71078	10.37	34.91	–	–	–	–	–
SLP17-19	Panalillo	Acid	31.05	85.36	29.23	8.44796	0.71258	8	0.70886	12.40	64.06	0.11702	0.51254	8	0.51252	-1.4
SLP17-23	Panalillo	Acid	29.4	121.35	5.28	66.50392	0.74749	11	0.71973	12.52	64.10	0.11812	0.51254	8	0.51251	-1.4
SLP17-24	Cantera	Acid	32.1	181.51	23.19	22.64472	0.72170	7	0.71138	6.80	30.04	0.13698	0.51249	8	0.51247	-2.3
SLP17-28	El Zapote	Acid	33.48	250.09	16.07	45.02054	0.73799	6	0.71659	2.13	11.13	0.11587	0.51249	8	0.51247	-2.3
SLP17-29	San José	Acid	33.95	144.26	14.01	29.79568	–	–	–	14.24	67.79	0.12706	0.51248	9	0.51245	-2.6

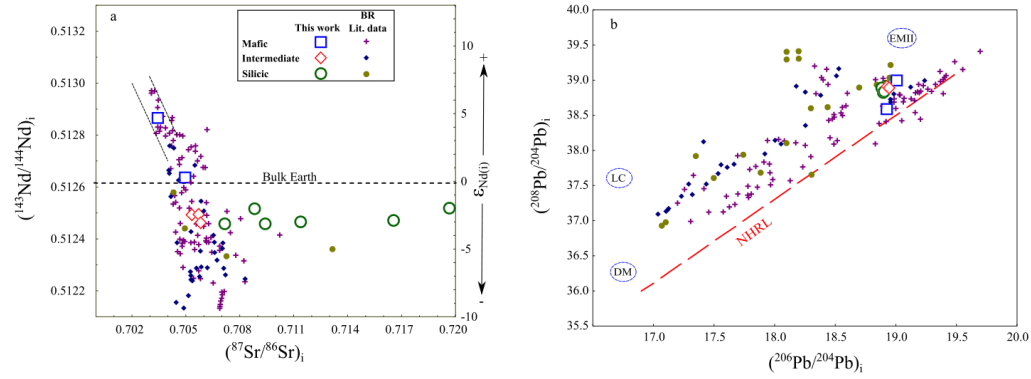


Fig. 8. Sr, Nd (modified after Verma, 1984) and Pb isotope plots of SSMC volcanic rocks compared to southern Basin and Range igneous rocks. (a) $^{143}Nd/^{144}Nd$ versus $^{87}Sr/^{86}Sr$ and ϵ_{Nd} plot of SSMC volcanic rocks; (b) $^{206}Pb/^{204}Pb$ versus $^{208}Pb/^{204}Pb$ plot SSMC volcanic rocks. Abbreviations: BR-Basin and Range; LC-Lower Crust; DM-Depleted Mantle; EMII-Enriched Mantle type-II; NHRL-Northern Hemisphere Reference Line. Compositions from DM, EMII are taken from Faure (1986) and NHRL values are taken from Zindler and Hart (1986). Literature data were taken from the following sources: Duncker et al. (1991); Johnson and Thompson (1991); Kempton and Fitton (1991); Gibson et al. (1992); Bradshaw et al. (1993); Davis and Hawkesworth (1995); Rogers et al. (1995); McMillan et al. (2000); and Christiansen et al. (2007).

Table 3

Whole-rock Pb isotopic data for volcanic rocks from the Sierra de San Miguelito Complex (age data for correction were taken from Table 4; element concentrations of Th, U and Pb were taken from Table 1).

Sample	Rock unit	Magma type	Age	Th (ppm)	U (ppm)	Pb (ppm)	$^{206}Pb/^{204}Pb$	$^{207}Pb/^{204}Pb$	$^{208}Pb/^{204}Pb$	$^{206}Pb/^{204}Pb_{(i)}$	$^{207}Pb/^{204}Pb_{(i)}$	$^{208}Pb/^{204}Pb_{(i)}$
SLP17-13	Cabras	Basic	22.21	2.13	0.74	3.066	18.928	15.614	38.571	18.925	15.329	38.571
SLP17-14	La Placa	Basic	30.13	2.751	0.599	4.164	18.965	15.665	38.909	18.962	15.303	38.909
SLP17-10	La Placa	Inter.	32.67	9.562	2.766	10.886	18.939	15.667	38.902	18.934	15.149	38.902
SLP17-20	La Placa	Inter.	27.5	7.037	1.242	8.495	18.894	15.664	38.864	18.891	15.248	38.864
SMB17-01	La Placa	Inter.	30	9.073	2.721	10.723	18.936	15.666	38.894	18.932	15.205	38.893
SMB17-02	La Placa	Inter.	30	8.01	2.653	9.958	18.938	15.665	38.889	18.933	15.227	38.889
SLP17-11	San Miguelito	Acid	33.17	28.959	4.414	26.661	18.900	15.666	38.885	18.897	15.016	38.885
SLP17-12	San Miguelito	Acid	33	44.084	6.664	51.671	18.905	15.661	38.831	18.903	15.154	38.831
SLP17-15	Panalillo	Acid	28.65	23.395	5.197	22.503	18.909	15.665	38.881	18.906	15.122	38.881
SLP17-16	Panalillo	Acid	28.65	36.907	8.373	16.621	19.013	15.680	39.016	19.005	14.521	39.016
SLP17-19	Panalillo	Acid	31.05	17.217	4.35	21.886	18.896	15.660	38.858	18.892	15.217	38.858
SLP17-23	Panalillo	Acid	29.4	25.063	5.222	26.949	18.897	15.659	38.861	18.894	15.162	38.861
SLP17-24	Cantera	Acid	32.1	25.802	5.005	27.019	18.907	15.666	38.889	18.903	15.112	38.889
SLP17-28	El Zapote	Acid	33.48	30.295	4.297	29.351	18.892	15.668	38.876	18.889	15.045	38.876
SLP17-29	San José	Acid	33.95	27.782	5.752	28.743	18.922	15.669	38.913	18.918	15.078	38.913

Table 4
Summary of $^{40}\text{Ar}/^{39}\text{Ar}$ ages of volcanic rocks from the Sierra de San Miguelito volcanic complex.

Sample	Plateau age (Ma) \pm 2 σ	^{39}Ar released (%)	Isochron age (Ma) \pm 2 σ	MSWD
<i>a) San Miguelito Unit</i>				
SLP17-11	33.17 \pm 0.67	80.1	34.80 \pm 5.30	13
<i>b) San José Unit</i>				
SLP17-29	33.95 \pm 0.44	98.1	34.09 \pm 0.48	1.7
<i>c) El Zapote Unit</i>				
SLP17-28	33.48 \pm 0.43	90.2	33.40 \pm 0.68	2.2
<i>d) Cantera Unit</i>				
SLP17-24	32.10 \pm 0.38	86	32.50 \pm 0.39	1.3
<i>e) La Placa Unit</i>				
SLP17-10	32.67 \pm 0.39	66.5	32.51 \pm 0.48	2.4
SLP17-14	30.13 \pm 0.42	81.5	30.47 \pm 0.99	3.5
SLP17-20	–	–	27.5 \pm 2.8	20
<i>f) Panalillo Unit</i>				
SLP17-19	31.05 \pm 0.37	94.6	29.50 \pm 2.30	4.9
SLP17-23	29.40 \pm 0.50	100	30.40 \pm 0.74	0.99
SLP17-15	28.65 \pm 0.37	83.4	30.60 \pm 2.00	3.2
SLP17-16	24.15 \pm 0.82	51.5	29.60 \pm 3.5	7.5
<i>g) Cabras Unit</i>				
SLP17-13	22.21 \pm 0.29	75.6	21.93 \pm 0.59	5.9

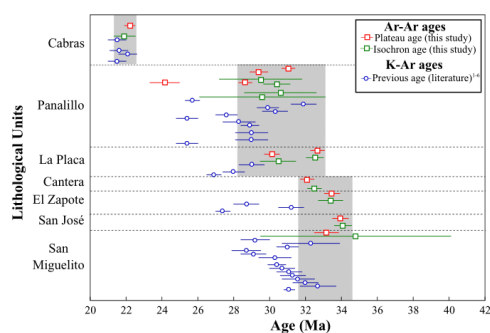


Fig. 9. Distribution of age from each SSMC unit. Gray shadow represents three main eruptive events of the SSMC. Previous age was taken from: ¹ Labarthe-Hernández et al. (1982); ² Tuta et al. (1988); ³ Aguillón-Robles et al. (1994); ⁴ Aguillón-Robles et al. (2009); ⁵ Tristán-González et al. (2009); ⁶ Aguillón-Robles et al. (2014).

arrangement of 0.45 Ol + 0.35 Opx + 0.2 Cpx was considered for the partial melting model. Melting degrees for the partial melting model that were used as $F = 0.01$ to 0.2 (Fig. 10d). The mafic sample (SLP17-13) could be produced by approximately 3–7% of partial melting of the mantle of a lherzolite composition, whereas sample SLP17-14 indicates a trend towards the partial melting line for granulite (Fig. 10d), suggesting lower crustal contamination of a mantle-derived melt. Such a melt regime could exist during an extensional phase within the area. In this context, it is noteworthy that the two mafic samples (SLP17-13 and SLP17-14) displayed different ages i.e., 30 Ma and 22.2 Ma, respectively. Thus, the differences in their initial isotopic composition from the older mafic sample (SLP17-14; 30 Ma) to the youngest mafic sample (SLP17-13; 22.2 Ma), may, on the other hand, represent a temporal switch over from melting the lithospheric- to asthenosphere-derived mafic magmas. However, more detailed sampling and analysis of mafic magmas of different ages is required to resolve these different petrogenetic possibilities. We do not consider

one sample each for a statistically representative evidence to pursue further these arguments.

5.2.2. Origin of the silicic volcanism

The origin of the silicic volcanic rocks from the SSMC have been discussed by several authors (i.e., Orozco-Esquivel et al., 2002; Rodríguez-Ríos and Torres-Aguilera, 2009; Aguillón-Robles et al., 1994, 2014), that proposed that the silicic volcanic rocks were derived from the partial melting of granulites from the lower continental crust. Recently, Torres-Sánchez et al. (2019) have proposed a first quantitative model for these volcanic rocks that indicated the silicic volcanic rocks were derived from melts of the upper continental crust. According to Xu et al. (2008), if the magmas are derived from partial melt of mafic granulites from the lower crust, the magma produced would have a metaluminous and calc-alkaline signature and are thus inadequate to account for the silicic volcanic rocks from the SSMC. Therefore, in this work we propose a new quantitative model for the partial melting processes that derived the generation of the silicic volcanism in the SSMC.

A crustal partial melting model was developed applying the batch melting equation reported by Zou (2007). Compositions of intermediate granulite were taken from Schaaf et al. (1994) (sample LP89). The mineralogical arrangements of continental crust reservoirs considered for the model are: 0.4Plg + 0.3Cpx + 0.2Gt + 0.05Opx + 0.05Amph. Melting degrees for the partial melting model were $F = 0.1$ to 0.9 (Fig. 10d). The bivariate model (Fig. 10d) displays that high grades of melting of intermediate granulites can explain the different silicic volcanic rocks of the SSMC (Fig. 10d).

5.2.3. Assimilation-fractional crystallization processes (AFC)

Fractional crystallization processes are generally combined with the assimilation of the wall rocks surrounding the magma chamber (DePaolo, 1981). The amount of assimilated material correlates to the quantity of magma that has crystallized through cooling (DePaolo, 1981).

For a better understanding of the processes that involved the evolution of the intermediate rocks from the SSMC an initial quantitative treatment was carried out to represent the assimilation-fractional crystallization (AFC) processes applying the equations proposed by DePaolo (1981). The isotopic composition of a basaltic magma (sample SLP1713, $\text{SiO}_2 = 47.76\%$ m/m) was taken to represent an initial magma (C_0), as well as the composition of metasediments (sample LP51; Schaaf et al., 1994), as a representative of the assimilated wall-rock (C_A). A mineralogical arrangement of 0.45Plg + 0.1Amph + 0.2Bt + 0.1Ap + 0.05Ol + 0.05Ksp + 0.05Zr was considered for the fractional crystallization process. Partition coefficients were taken from Villemant et al. (1981) and McKenzie and O'Nions (1991). Isotopic AFC models (Fig. 11) were calculated considering the ratios for $r = 0.1$, $r = 0.3$, $r = 0.5$, $r = 0.7$ and $r = 0.9$ of the assimilation to the fractional crystallization, as well as a fraction of magma remaining $F = 0.9$ to 0.1.

The AFC modelling yields a better fit to isotopic data, indicating that most of the intermediate rocks of the SSMC experience variable degrees of assimilation ($r > 0.3$) of metasediments combined with fractional crystallization. The model reveals ~35 to 40% of assimilation of metasediments from the continental crust (Fig. 11).

5.3. Tectonic settings

In this work, we use new multi-dimensional discrimination diagrams to infer the plate tectonic setting, due to conventional bivariate and ternary diagrams revealing several problems related to statistical treatment of compositional data. The most important reason for this failure is related to the use of limited databases, problems of closed or constant sum compositional variables, and eye-fitted tectonic field boundaries and distribution of compositional data (Verma et al., 2012;

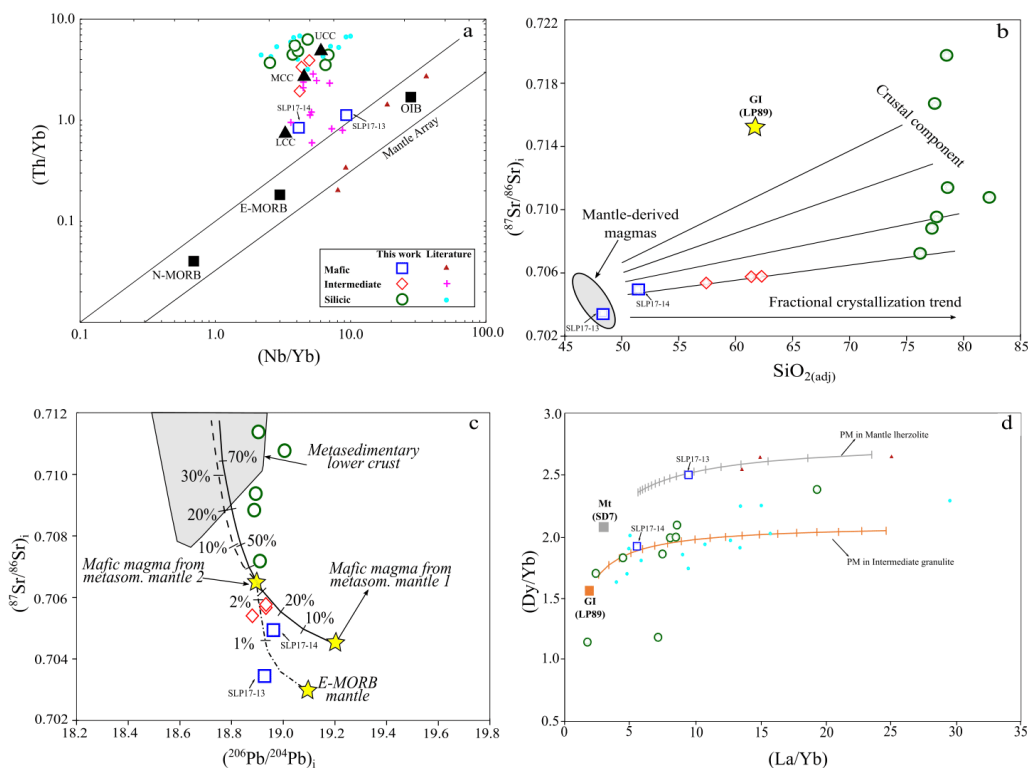


Fig. 10. a) (Th/Yb)₁–(Nb/Yb)₁ diagram for volcanic rocks from the SSMC; (b) SiO_{2(adj)} vs ⁸⁷Sr/⁸⁶Sr₍₁₎ for the SSMC volcanic rocks. The positive correlation trends can be explained either by AFC processes or mixing between mantle-derived mafic magmas and crustal components, whereas the near-horizontal trend indicates closed-system fractional crystallization.; (c) Petrogenetic modelling for the genesis of the SSMC mafic volcanic rocks based on the variation of ⁸⁷Sr/⁸⁶Sr₍₁₎ vs ²⁰⁶Pb/²⁰⁴Pb₍₁₎. Model is taken and modify after Harangi et al. (2007); (d) (La/Yb)₁–(Dy/Yb)₁ partial model, values of granulite and mantle taken from Schaaf et al. (1994). Each small bar represents 5% of melting for the orange line (granulite melting) and 1% of melting for the gray line (mantle melting). Literature data were taken from the following sources: Orozco-Esquivel et al. (2002); Leroy et al. (2002); Rodríguez-Ríos and Torres-Aguilera (2009); Torres-Hernández et al. (2014); Aguilón-Aguillón-Robles et al. (2014) and Torres-Sánchez et al. (2019). Abbreviations: E-MORB–Enriched Mid-Ocean Ridge; GI–Intermediate Granulite; LCC–Lower Continental Crust; MCC–Middle Continental Crust; Mt–Mantle composition; N-MORB–Normal Mid-Ocean Ridge; OIB–Ocean Island Basalt; UCC–Upper Continental Crust.

Verma et al., 2013). Therefore, new multidimensional discrimination diagrams proposed by Agrawal et al. (2008), Verma and Verma (2013) and Verma et al. (2013) for mafic, intermediate and silicic rocks were used for tectonic interpretation.

In the multidimensional discrimination diagrams by Agrawal et al. (2008) the mafic volcanic rocks from the SSMC were discriminated as within-plate (CR + OI; Fig. 12a) setting. This was also the case for intermediate and silicic rocks (Fig. 12b–c; Verma and Verma, 2013 and Verma et al., 2013). Conventional Sr–Nd isotope diagram (Fig. 12d) was used for comparison of the SSMC volcanic rocks with compiled literature data from continental rifts (Verma and Verma, 2018). The mafic volcanic rocks from the SSMC display a trend towards the mantle array, as well as the evolved rocks (intermediate and silicic volcanic rocks) continental rift rocks. The Nb and Ta anomalies in the primitive mantle-normalized diagrams for the SSMC rocks also support these findings. More detailed sampling and additional data in future could be evaluated from the most recent multidimensional tectonomagmatic models (Verma, 2020b) for better understanding the evolution of the tectonic regime through time from the Oligocene to the Quaternary.

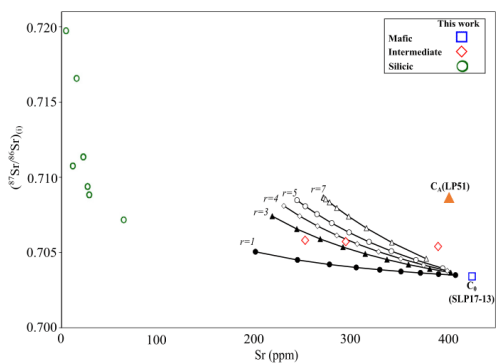


Fig. 11. Sr vs ⁸⁷Sr/⁸⁶Sr₍₁₎ assimilation and fractional crystallization model for the intermediate volcanic rocks from the SSMC, values of metasediments were taken from Schaaf et al. (1994). Abbreviations: Co-Initial magma; CA–assimilated wall-rock.

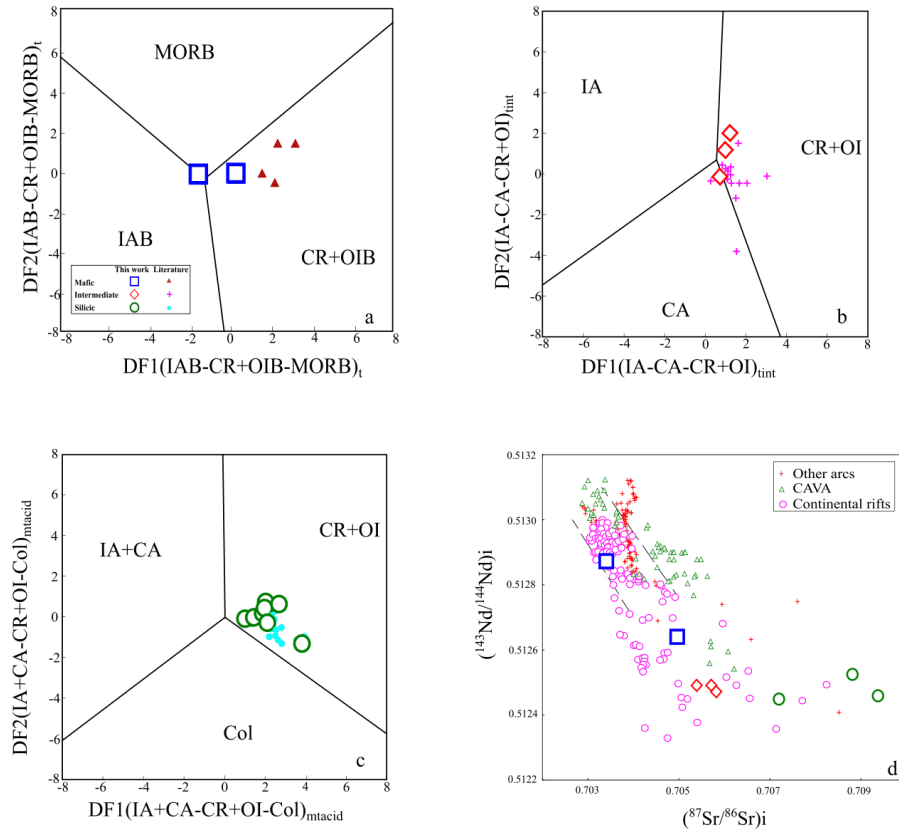


Fig. 12. a) Major-trace element based multidimensional discrimination diagram for mafic rocks (Verma and Agrawal, 2011); b) Major-trace element based multidimensional discrimination diagram for intermediate rocks (Verma and Verma, 2013); c) Immobile trace element based multidimensional discrimination diagram for silicic rocks (Verma et al., 2013) from the SSMC. d) Conventional Sr–Nd isotope diagram for the SSMC. Abbreviations: CAVA-Central American Volcanic Arc; Col-Collision; CR + OI-combined continental rift and ocean island; CR + OIB-combined continental rift and ocean island basalt; IA-Island Arc; IA + CA-combined island arc and continental arc; IAB-Island Arc Basalt; MORB-Mid-Ocean Ridge.

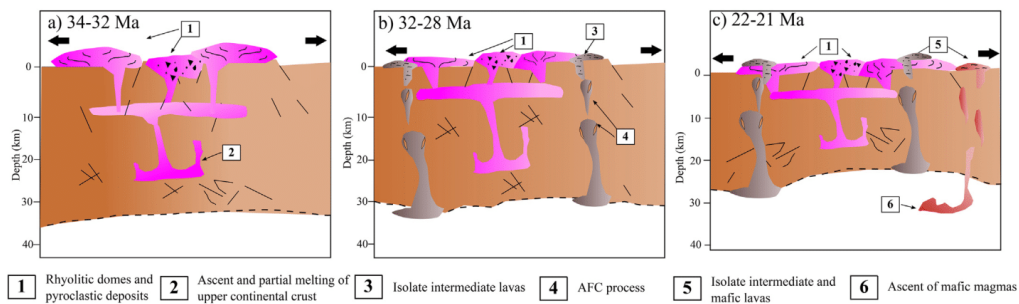


Fig. 13. Schematic diagram showing the evolution of the SSMC volcanic rocks between 34 Ma and 21 Ma. a) Generation and ascent of rhyolites and ignimbrites volcanic rocks during the late Eocene to early Oligocene; b) Ascent and assimilation and fractional crystallization of intermediate volcanic rocks during the late Oligocene; c) Emplacement of mafic volcanic rocks at the early Miocene. Black arrows represent continental crust extensional regime; thickness values of the continental crust were taken from Nieto-Samaniego et al. (2007); the figure was modified after Aguilón-Robles et al. (2014) and Sieck et al. (2019).

5.4. Geodynamic implications

The petrological observations indicate that the Cenozoic units from the SSMC were derived from a lithospheric mantle source that was heterogeneously enriched by continental contributions. Some authors suggest that the volcanic rocks from the SSMC were generated by a transitional subduction-intraplate setting from 42 to 31 Ma (Aguillón-Robles et al., 2014). Similarly, Tristán-González et al. (2009) proposed that Eocene-Oligocene (42–32 Ma) magmatism was associated with a typical convergence regime of continental volcanic arc and the Miocene (22–20 Ma) volcanic rocks were generated in an intra-plate regime. Rodríguez-Ríos et al. (2007) suggested that the silicic volcanic rocks evolved through a partial melting of a Precambrian lower continental crust (granulite facies).

However, the present study reveals that silicic volcanic rocks of the SSMC were generated through melting of intermediate granulites at 34–32 Ma. Meanwhile, intermediate volcanic rocks were erupted during the early Oligocene (~32–29 Ma) derived from lithospheric mantle melts with contaminated by metasediments. Mafic volcanism was produced at 21 Ma through low degree partial melting of the lithospheric mantle. The observations and data from the present study may be used to hypothesize the magmatic evolution of the SSMC volcanic rocks during late Eocene to early Miocene by extensional regimes (Fig. 13).

If the geological observations are considered, it is apparent that the Eocene to Miocene volcanism in the study area was developed synchronously with the tectonic exhumation of the volcanic rocks from the BR and southern region of the MC. These observations may be used to discuss the nature of the Cenozoic extension systems that produced the volcanic rocks across the southern section of the BR.

6. Conclusions

From the integrated petrography, geochemistry, geochronological and isotopic data obtained from the SSMC, the following conclusions can be drawn:

1. The magmatic rocks of the SSMC have a wide composition range of trachybasalt, basalt, basaltic trachyandesite, basaltic andesite, andesite and rhyolite. They display porphyritic textures with disequilibrium features in the mafic volcanic rocks. Although there is a wide range of compositions documented in this study, there is no evidence for synchronous bimodality therefore does not show true bimodal volcanism.
2. Geochemical features indicate that the mafic volcanic rocks from the SSMC were derived from two sources: one derived from the lithospheric mantle alone, and a second source involving a lower crustal component. On the other hand, intermediate volcanic rocks derived from assimilation and fractional crystallization processes of the mafic melt. It is likely that the silicic volcanic rocks were derived from partial melting of mid continental crust granulites.
3. Volcanism was active in the SSMC between 34 and 21 Ma. Magmas were erupted through three episodes: an initial phase characterized by domes and rhyolite ignimbrites, followed by a second phase of volcanism represented by intermediate rocks intercalated with pyroclastic rocks of rhyolite composition; a third phase is characterized by isolated mafic lavas.
4. Magmatic rocks of the SSMC were probably formed in a within-plate (CR + OI) setting, during BR related extension, in the late Eocene to Miocene time, although more detailed sampling and analyses are required to better document the time-related tectonic evolution.

Declaration of Competing Interest

This paper has not been published elsewhere, nor is it under consideration of any other journal. It is an original contribution, authors have contributed to its development, and there is no conflict of interest.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.lithos.2020.105613>.

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CAPITULO 7.-

ANÁLISIS QUÍMICO DE MINERALES

En este estudio se aplicó un método de combinación de técnicas tanto de Microsonda Electrónica (Electron Probe Microanalysis, EMPA), para llevar el análisis cuantitativo de los minerales y de Microscopía de Barrido Electrónico (Scanning Electron Microscope, SEM) para la toma de imágenes de retrodispersión de electrones de alta resolución para los minerales analizados.

El análisis químico aplicando el método EMPA se lleva a cabo bombardeando la muestra con un haz de electrones acelerados a 10-20 kV. El haz electrónico de 10 μm de diámetro se enfoca sobre una superficie pulida de la muestra. Esta emite rayos-X característicos de cada elemento los cuales son analizados en un detector de dispersión de energía o en uno de dispersión de longitud de onda. El análisis cuantitativo con EMPA se encuentra limitado generalmente a elementos de $Z > 11$. Los elementos con concentraciones $> 0.05\%$ son detectados sin dificultad, aunque no es posible distinguir iones con diferentes estados de oxidación ([Johnson y Maxwell, 1981](#)).

Los resultados obtenidos para el presente estudio se obtuvieron en el Departamento de Geología de la Open University de Milton Keynes, Inglaterra. El equipo utilizado fue una microsonda electrónica Cameca SX100. Se seleccionaron 5 muestras de las unidades (2 muestras de composición félsicas, 2 muestras de composición intermedia y 1 muestra de composición básica) del Complejo Volcánico de la Sierra de San Miguelito.

La medición se llevó a cabo acelerando una corriente de 10 nA con un voltaje de 15 kV. Los tiempos de conteo de los picos fueron de 10 a 40s y para la corrección de matriz se utilizó una rutina de PhiRhoZ. Los siguientes elementos fueron medidos utilizando una espectroscopia de longitud de onda dispersiva: K, Al, Si, Fe, Mn, Ca, P, Ti, Na and Mg.

La microsonda fue calibrada usando los estándares minerales naturales de: Feldespato (K,Al,Si), hematita (Fe), bustamita (Mn, Ca), apatito (P), rutilo (Ti), jadeíta (Na) y fosterita (Mg). Detalles adicionales del análisis de microsonda son reportados en el apartado de Anexos A7.

En los siguientes apartados se presentará los resultados obtenidos por este método, como también, el procesamiento de los mismos. A su vez, los datos obtenidos se reportan en la Tabla A8 en la sección de anexos.

7.1 Composiciones minerales

A partir de la abundancia mineral conocida de las rocas volcánicas del Campo Volcánico de la Sierra de San Miguelito, las composiciones minerales fueron obtenidas de fenocristales de plagioclasas, clinopiroxenos y olivinos para cada uno de los tipos de rocas (máficas, intermedias y ácidas; Tabla A8).

Las composiciones de plagioclasas en las rocas máficas e intermedias indican que estas muestran un rango composicional de An₃₂₋₆₉ a Ab₂₉₋₄₂ (Figura 7.1a) Por otro lado las rocas ácidas muestran un rango composicional de plagioclasas de An₂₉₋₂₈ a Ab₆₅₋₆₆ (Figura 7.1a), de igual forma, en las rocas ácidas se puede observar la presencia de feldespatos alcalinos, los cuales muestran concentraciones de albita (Ab) y ortoclasa (Or) con abundancias de 65-66 y 5.16 a 5.40 respectivamente.

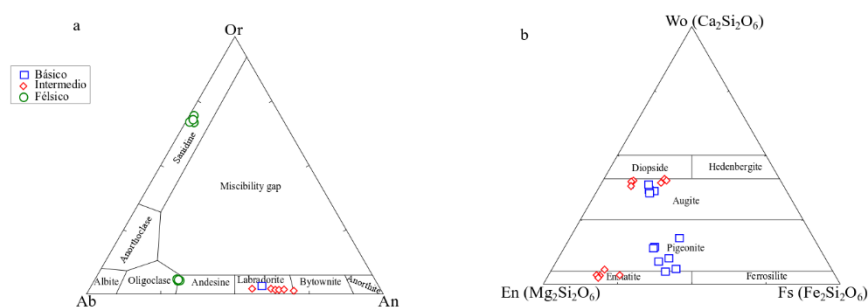


Figura 7.1 a) Diagrama de clasificación de feldespatos para las rocas del Complejo Volcánico de la Sierra de San Miguelito; b) Diagrama de clasificación de piroxenos para las rocas del Complejo Volcánico de la Sierra de San Miguelito.

Los feldespatos alcalinos muestran un contenido el cual difiere a la de los fenocristales de plagioclasa (Ab_{29-42} y Or_{2-25}), que sugiere que la cristalización de los dos estilos de feldespatos no se encuentra en equilibrio (Figura 7.1^a).

Por otro lado, las rocas máficas e intermedias muestran dos estilos de piroxenos (Figura 7.1b). Los clinopiroxenos tanto de las rocas máficas muestran una composición de augita (Wo_{36-39} y En_{44-45} ; Figura 7.1b), mientras que los ortopiroxenos correspondientes muestran composiciones de enstatita (En_{76-79} y Wo_{3-5} ; Figura 7.1b). El rango composicional de las rocas intermedias (En_{44-51} y Wo_{35-39}) marca una similitud a las rocas máficas, lo cual indican que cristalizaron en equilibrio.

7.2 Geotermometría

El cálculo de temperaturas en equilibrio tanto de las rocas ácidas y básicas del Campo Volcánico de la Sierra de San Miguelito fue desarrollado siguiendo las ecuaciones de termómetros propuestas por Putirka (2008) tanto para plagioclasa-feldespatos alcalinos-líquido, termómetros de ortopiroxeno y clinopiroxeno. La ecuación correspondiente de cada uno de estos se reportan en la sección de Anexos en la Tabla A9.

Sin embargo, se debe de prestar atención en el uso de estas ecuaciones empíricas que no llegan a presentar incertidumbres en los coeficientes de los términos, la evaluación de estas se dispone a tratar en futuras investigaciones.

Las temperaturas de equilibrio en ortopiroxenos en las unidad intermedia de La Placa presenta valores promedio de 967° a 905°C (total de análisis puntuales de 10), en contrario, las rocas básicas (unidad Cabras) las cuales muestran un valor promedio de 1241°C (total de análisis puntuales de 7). De igual forma, estas temperaturas fueron calculadas siguiendo un valor de presión experimental de 1GPa.

Dentro del diagrama de Rhodes (Figura 7.2a), se observa que los valores de ortopiroxenos de las rocas intermedias muestran una tendencia hacia una remoción de ortopiroxeno lo cual puede marcar una fraccionación dentro de la evolución del magma lo cual es coherente con los modelos de asimilación y cristalización fracciona plantados previamente. Por el contrario, la unidad básica del CVSSM (Figura 7.2a) muestra una tendencia hacia una diferenciación, lo cual remarca que el proceso principal de generación de las rocas básicas dentro del complejo es un proceso de fusión parcial.

Las rocas intermedias muestran temperaturas de clinopiroxeno de 973-1014°C, a partir de presiones experimentales de 1GPa (total de análisis puntuales de 9), por otro lado, las rocas básicas muestran un valor de 1392°C (total de análisis puntuales de 6). Este último valor de temperatura fue calculado siguiendo un valor de presión experimental de 1GPa.

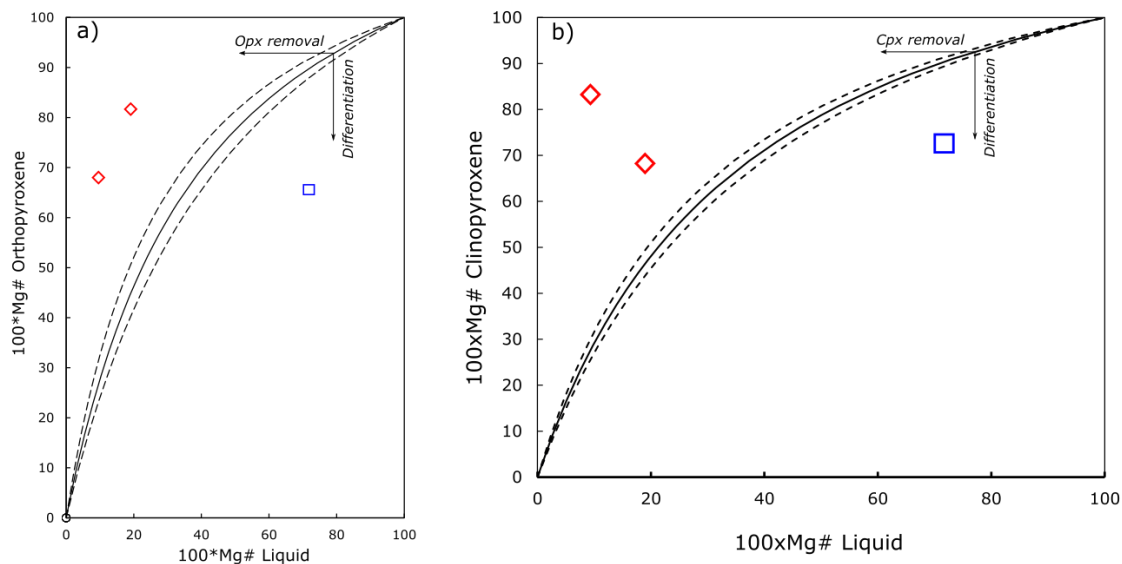


Figura 7.2 a) Diagrama de Rhodes para ortopiroxenos de las rocas intermedias y básicas del Complejo de la Sierra de San Miguelito; b) Diagrama de Rhodes para clinopiroxenos de las rocas intermedias y básicas del Complejo de la Sierra de San Miguelito. Valores promedio de las rocas intermedias (rombo color rojo) y básicas (cuadrado en color azul).

El diagrama de Rhodes (Figura 7.2b) para los clinopiroxenos muestra que las rocas intermedias tienden en una dirección de remoción de clinopiroxeno lo cual, igual que con los ortopiroxenos, demuestran una fraccionación dentro de la evolución del magma, como lo indica el modelo de asimilación y cristalización fraccionada. En contraste a esto, la unidad de composición básica (Figura 7.2b) muestra una tendencia hacia una diferenciación, lo cual confirma que el proceso principal de generación en este estilo de rocas fue un proceso de fusión parcial.

Por otro lado, los valores de temperatura de equilibrio en feldespatos alcalinos de las unidades félsicas muestran un rango promedio de 854° a 880°C, estas temperaturas fueron calculadas siguiendo un valor de presión experimental de 1 GPa, a partir de un total de 12 análisis puntuales en el centro y borde del cristal.

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CAPITULO 8.-

CONCLUSIONES

La elaboración del presente trabajo aporta nueva información petrogenética para la comprensión de la evolución de las rocas magmáticas, como también limita los distintos episodios magmáticos dentro de la región. Las principales limitaciones en este trabajo se presentan en la corta extensión de los afloramientos de las rocas de composiciones básicas e intermedias, como también la accesibilidad de ciertas regiones del área de estudio.

A pesar de las limitaciones el presente trabajo demuestra que el CVSSM no presenta un comportamiento bimodal, si no que exhibe un rango composicional amplio que incluye rocas de composición básicas, intermedias y ácidas; como también, un contraste en modelos petrogenéticos propuestos por trabajos previas. En estos trabajos previos se proponía que las rocas félsicas fueron desarrolladas por procesos de fusión parcial de la corteza continental inferior, sin embargo, en este trabajo se reporta que las rocas félsicas se originaron a partir de la fusión parcial en niveles medio-superior de la corteza continental; como también, no se había propuesto modelos cuantitativos que ayuden a comprender los procesos magmáticos que generaron el magmatismo básico e intermedio. De igual forma la información generada en el presente trabajo apoyan las cuatro hipótesis enlistadas en la sección 1.2.

Los resultados se simplifican en:

1. El CVSSM se constituye por rocas de composición amplia. Entre las rocas básicas se encuentran rocas de composición traquibasáltica, basáltica y foidítica, a su vez, las rocas intermedias presentan composiciones traquiandesítica basáltica, andesita basáltica y andesítica. Por último las rocas ácidas del CVSSM muestran composiciones riolíticas y traquíticas.

2. Los tres tipos de rocas presentes en el CVSSM muestran texturas porfíricas, como a su vez, se muestran texturas de desequilibrio en las rocas básicas e intermedias. El ensamble mineralógico principal de las rocas básicas del CVSSM se caracteriza por fenocristales de plagioclasa, olivino, clinopiroxeno y ortopiroxeno, por otra parte, las rocas intermedias presentan fenocristales similares con la excepción de la abundancia de ellos. De igual forma, los fenocristales principales de las rocas ácidas del CVSSM son feldespatos alcalinos (sanidino), plagioclasas, piroxenos (ortopiroxenos y clinopiroxenos).
3. La información geocronológica muestra que el vulcanismo del CVSSM inició hace 34 Ma hasta los 21 Ma. Los magmas del CVSSM se formaron a través de tres episodios, un primer episodio caracterizado por estructuras de domos y de piroclásticos de composiciones riolíticas; la segunda fase de vulcanismo es representada por rocas intermedias intercaladas con rocas piroclásticas de composición riolítica; y por último la tercera fase consiste en lavas de composición máfica aisladas.
4. Las características geoquímicas de las rocas máficas del CVSSM muestran un enriquecimiento en tierras raras ligeras, como también, un enriquecimiento en elementos traza menos incompatibles, junto con valores positivos de $\epsilon_{Nd(t)}$ (+5.2 - +0.9), lo cual marca una fuente del manto. A su vez, las rocas intermedias del CVSSM muestran patrones de tierras raras y traza similares a las rocas básicas, con la diferencia de pequeñas anomalías negativas de Eu, como también, estas rocas muestran valores negativos de $\epsilon_{Nd(t)}$ (-2.5 a -1.8), lo cual indica una contribución de la corteza en la evolución de este estilo de rocas. Por último, las rocas félsicas muestran patrones de enriquecimiento en tierras raras ligeras con pronunciadas anomalías negativas de Eu, lo que marca un fraccionamiento de feldespatos, de igual forma, las composiciones isotópicas de las rocas félsicas muestran valores negativos de $\epsilon_{Nd(t)}$ (-2.6 a -0.2), lo que nos indica una contribución de la corteza en la generación evolutiva de este estilo de rocas.

5. La modelación de procesos magmáticos de las rocas del CVSSM revelan que las rocas máficas evolucionaron por la fusión parcial de una fuente de manto litosférico, principalmente de fuentes lherzolíticas a niveles de fusión (~15-20%) aproximadamente a los 22–21 Ma, mientras que las rocas intermedias se generaron por fraccionamiento de una líquido parental máfico el cual presentó distintos niveles de asimilación ($r > 0.3$) con una corteza compuesta por metasedimentos en un periodo de 32–28Ma. Por último, los modelos revelan que las rocas félsicas se generaron a partir de un proceso de fusión parcial de metasedimentos a niveles de corteza media y superior en un periodo de tiempo que abarca de los 34 a 32Ma.
6. Las rocas magmáticas del CVSSM se indican que fueron generadas a partir de un ambiente tectónico intra-placa durante el Eoceno hasta el Mioceno, lo cual puede ser correlacionado con las anomalías de Nb y Ta. Esto puede ser relacionado al régimen extensional de la provincia del Basin and Range.

8.1 Recomendaciones y futuros trabajos

El presente trabajo marca una iniciativa para futuros trabajos dentro del Complejo Volcánico de la Sierra de San Miguelito. A partir de los resultados obtenidos en el presente trabajo, se recomienda que para próximas investigaciones se marquen un enfoque en el estudio a profundidad de las rocas máficas de la región, ya que la comprensión de estas rocas junto con información de la corteza subyacente en el área, brindará modelos petrogenéticos y tectónicos integrales.

De igual forma, la aplicación de modelos cuantitativos de inversión en rocas máficas del complejo, permitirán evaluar los procesos de fusión parcial que dieron origen a estos estilos de magmas. Es importante señalar que modelos más confiables pueden ser preparados si se cuenta con una base más amplia de coeficientes de partición y un mayor número de datos analíticos confiables de elementos traza. Estos modelos podrían ser de alta confiabilidad, y representativos, si se cuantifica los coeficientes de partición en las rocas magmáticas del complejo por medio de mediciones directas, por ejemplo, por el método de *Laser Ablation ICP-MS*.

A.1 Análisis Petrográfico

Se elaboraron 20 láminas delgadas para efectuar el análisis petrográfico. La preparación de las mismas se llevó a cabo en el Laboratorio de Preparación de la Facultad de Ciencias de la Tierra, UANL.

El procedimiento que se llevó a cabo para la elaboración de las secciones delgadas es el siguiente:

- a) **Corte:** el fragmento de la roca fue cortada con una sierra de disco diamantado (Fig. A1a) para obtener cubos de una medida de 4x2x3 cm.
- b) **Pulido:** se seleccionó una de las caras del cubo para ser pulido utilizando un disco metálico (Fig. A1b), al cual se le agrega agua y polvo abrasivo de SiC (Carburo de Silicio) de una granulometría de 200 μm , hasta obtener una superficie plana. La superficie obtenida fue pulida en una placa de vidrio, utilizando agua y polvos abrasivos de distinta granulometría (400, 600, 800 μm).
- c) **Pegado:** la superficie pulida se pegó sobre un portaobjetos de vidrio con un agente cementante incoloro e isótropo (mezcla de resina epóxica KÖRAPOX 439; mezcla de resina cristalizados en relación 2:1).
- d) **Corte final:** una vez pegado el cubo de roca en el portaobjetos se cortó con un disco de diamante más fino hasta obtener un espesor de ~ 1 mm.
- e) **Pulido final:** la sección obtenida se sometió a un pulido (Fig. Ac, d) como el anteriormente mencionado hasta obtener un espesor aproximado de 25-30 μm . El proceso fue controlado principalmente siguiendo los colores de interferencia de la tabla de Michel Levy para los minerales base presentes (p ej. plagioclasas, cuarzo).

La mineralogía se determinó en base a las características reportadas en literatura especializada (Heinrich, 1970; Philpotts, 1989; McKenzie et al., 1996; Vernon, 2004; Higgins, 2006). De igual forma, las composiciones modales fueron determinadas por conteo en láminas delgadas usando un microscopio petrográfico Leica y un contador semi-Automático PELCON, esto fue determinado en el Instituto Potosino de Investigación Científica y Tecnológica, San Luis Potosí (IPICYT).



Figura A.1 a) Cortadora con disco de diamante; b) Discos metálicos giratorios para pulido de muestras; c) Vidrios para pulido ; d) Láminas delgadas terminadas.

A.2 Separación de minerales

La separación de minerales se llevó a cabo a partir de la trituración de 12 rocas a partir de un triturador de quijada ASC (Fig. A2a), obteniendo fragmentos de aproximadamente de 3-4 mm, los cuales pasaron por tres distintas cribas de 500, 1000, 2000 μm respectivamente (Fig. A2b). Una vez separada la granulometría deseada se pasa a secar la muestra húmeda con una lámpara de luz roja (Fig. A2c, d) por aproximadamente 30 minutos.

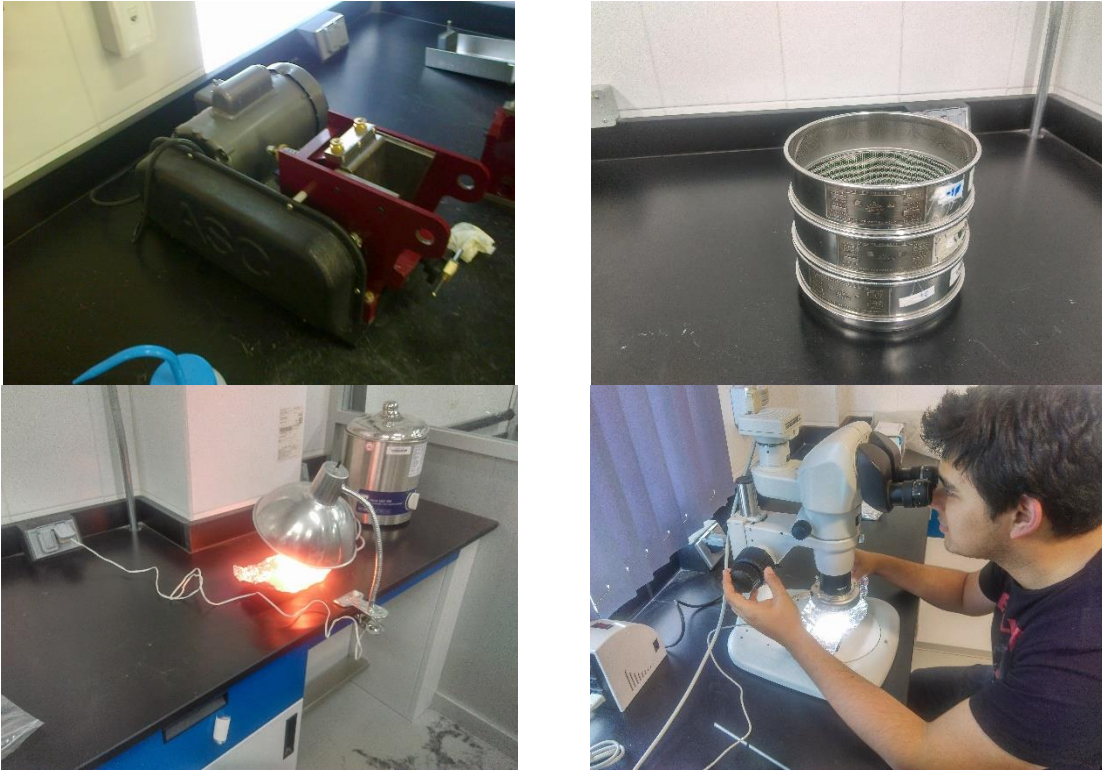


Figura A2.- a) Trituradora de quijada ASC. ; **b)** Cribas de 500, 1000, 2000 μm de espesor; **c)** Muestra cribada bajo lámpara con luz roja para un secado; **d)** Triturado y cribado terminado para separado manual.

Al finalizar el secado, se separaron cristales de sanidino de aproximadamente de 1-3 mm de longitud para las rocas félsicas y fragmentos de roca de ~ 1-4 mm de longitud para las rocas máficas con ayuda de un microscopio estereoscópico Nikon. Los fragmentos y minerales separados fueron seleccionados considerando la menor cantidad de imperfecciones en los cristales y fragmentos, como a su vez, se recolectaron la mayor cantidad de minerales sin estar rotos.

A.3 Trituración y molienda

Las muestras recolectadas pasaron por un procesamiento de molienda, el cual tiene el fin de realizar el análisis geoquímico de elementos mayores, tierras raras, traza e isótopos

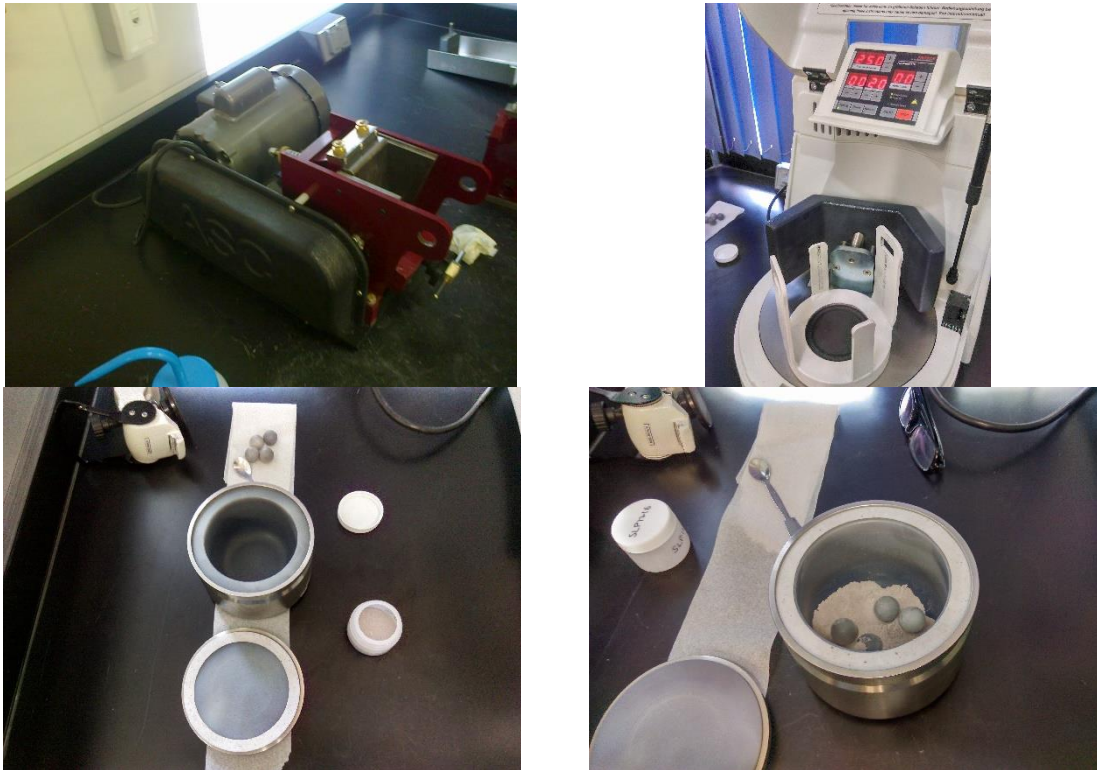


Figura A3.- a) Trituradora de quijada ASC; **b) y c)** Molino de ágata PLANETARY MONO MILL PULVERISETTE, FRITSCH; **d)** Polvo de roca terminado.

En este proceso se retiró la capa superficial a cada roca, a fin de contar con una muestra con la menor cantidad de alteración posible, considerando una cantidad aproximada de 500 g.

La roca una vez libre de alteración, se procesó en una triturada de quijada ASC (Fig. A3a), para así reducir los fragmentos a un tamaño aproximado de 0.3-0.5 cm, los cuales se pasaron a pulverizar utilizando un molino de ágata PLANETARY MONO MILL PULVERISETTE, FRITSCH (Fig. A3b, c), generando polvo de roca con un tamaño de partícula aproximado de 75-38 μm .

A.4 Análisis químico de elementos mayores, tierras raras y traza

Los análisis de elementos mayores, tierras raras y traza se dividieron en dos partes. La primera parte consistió en el análisis de 23 muestras, las cuales se prepararon siguiendo la metodología previamente mencionada. Los elementos mayores fueron determinados por fluorescencia de rayos-X de longitud de onda dispersiva (WD-XRF) Rigaku ZSX Primus II en el Instituto Potosino de Investigación Científica y

Tecnológica (IPICYT), San Luis Potosí. Los procedimientos analíticos, exactitud y precisión son dados por Verma et al. (2018). Las concentraciones de elementos traza y tierras raras fueron determinados en los laboratorios ALS Chemex, Canadá usando el método *ME-MS81*, para más información de este método se puede consultar en www.alsglobal.com/geochemistry. Las concentraciones de elementos traza y tierras raras se obtuvieron por Espectrometría de Masas con Plasma Acoplado Inductivamente, con una precisión analítica basa en materiales de referencia geoquímica SY-2 Y OREAS146.

En la segunda parte de análisis, se seleccionaron 15 muestras, las cuales fueron trituradas siguiendo la metodología previamente descrita. Los análisis de esta segunda parte fueron analizados en la Universidad de Leicester, Inglaterra. Los elementos mayores fueron determinados en perlas de fusión preparadas a partir de polvos pre-calentadas las cuales fueron fundidas con metaborato de litio (80% de metaborato de Li y 20% de tetraborato de Li) en una relación de 1:5. Los elementos traza fueron determinados a partir de pellets de polvo prensados elaboradas con una mezcla de 7g de muestra y una solución de 7% PVA (Alcohol Polivinilo). Los análisis se llevaron a cabo en la Universidad de Leicester, Inglaterra usando una fluorescencia de rayos-X PANalytical Arios. Los estándares internos utilizados fueron BH-1, WS-1, BCS375, BCS 376, MRG-1 y NIM-D los cuales a su vez fueron analizados para el monitoreo de la precisión y exactitud de los resultados. La precisión analítica y exactitud fue considerada entre 2-5% para los elementos mayores y <2% para elementos traza. Las concentraciones de tierras raras fueron determinadas de igual forma en la Universidad de Leicester, Inglaterra por un espectrómetro de masas ICP quadruple Thermo Scientific ICAP-Qc. Las muestras fueron preparadas usando una digestión estándar de HF-HNO₃. Siete materiales de referencia (BHVO-1, NIM-G, MRG-1, JSd-1, JSd-2, JSD-3, y JR-1) fueron usados para la calibración y control de calidad del análisis

A.5 Análisis de datación ³⁹Ar/⁴⁰Ar

La irradiación de 12 muestras fue llevada a cabo en el reactor Nuclear McMaster (Universidad de Canadá) por cerca de 101 horas. Se utilizó blindaje de

cadmio y las muestras se mantuvieron en la posición 8D. El flujo de neutrones fue monitoreado usando el estándar de mineral biotita GA1550 (99.738 ± 0.104 Ma; [Renne et al., 2011](#)).

Los análisis se llevaron a cabo en el laboratorio de Ar/Ar de la Open University, Milton Keynes, Inglaterra. Los estándares fueron empaquetados para irradiación a cada lado de las muestras y analizados utilizando el método de fusión de grano único, con una fibra de laser CSI de 1059 nm y un espectrómetro de masas MAP215-50. Los valores J fueron calculados por extrapolación lineal entre dos valores J medidos y un error de 0.5%.

Las muestras irradiadas fueron cargadas en un sistema de vacío y una fibra de laser CSI de 1059 nm fue enfocado dentro de la cámara de muestras y fue usado para el calentamiento escalonado por muestra. Los sistemas de blancos fueron medidos antes y después de cada dos análisis de muestra. La limpieza y entrada de gas es totalmente automatizada con la medición de ^{40}Ar , ^{39}Ar , ^{38}Ar , ^{37}Ar , y ^{36}Ar por cada 10 escaneos.

A.6 Análisis isotópicos (Sr-Nd-Pb)

Se seleccionaron 12 muestras representativas de las unidades litológicas del Complejo Volcánico de la Sierra de San Miguelito, en las cuales se analizaron las composiciones isotópicas de Sr, Nd y Pb.

Las muestras fueron preparadas y analizadas siguiendo los procedimientos de Kempton (1995) y Royse et al., (1998). Las composiciones isotópicas de Sr, Nd, y Pb fueron analizados como especímenes metálicos en filamentos individual de Ta, filamento doble Re-Ta y filamento individual de Re, respectivamente, usando un espectrómetro de masas multi-colector Finnigan MAT 262 (Fig. A4) en el NERC Laboratorio Isotópico de Geociencias (NIGL), Keyworth, Inglaterra.



Figura A4.- Espectrómetro de masas multi-colector Finnigan MAT 262, NERC Laboratorio Isotópico de Geociencias (NIGL), Keyworth, Inglaterra

Las muestras fueron lixiviadas aproximadamente 1 hora en 6M HCl a 50°C antes de la digestión por un procedimiento estándar de HF- HNO₃. Blancos para Sr, Nd y Pb fueron menos que 125 pg, 275pg y 325 pg, respectivamente.

Las relaciones de ⁸⁷Sr/⁸⁶Sr fueron normalizadas durante el tiempo de corrida a ⁸⁷Sr/⁸⁶Sr = 0.1194 y la relación ¹⁴³Nd/¹⁴⁴Nd al valor de ¹⁴³Nd/¹⁴⁴Nd = 0.7219. Los datos de muestra se informan en relación con los valores aceptados de los estándares NBS987 de ⁸⁷Sr/⁸⁶Sr = 0.710177 y 0.512128 de ¹⁴³Nd/¹⁴⁴Nd para el estándar La Jolla. Basado en corridas repetidas de NBS981, la reproducibilidad de relaciones de isótopos de Pb es ~ 0.1%. Las relaciones de Pb fueron corregidas relativamente al estándar promedio de composiciones isotópicas de Pb de Todt et al., (1996). Los valores medidos fuer corregidos en edad a partir de los datos obtenidos por datación de Ar.

A.7 Análisis químico de minerales

En este estudio se aplicó un método de Microsonda Electrónica (Electron Probe Microanalysis, EMPA), para llevar el análisis cuantitativo de los minerales para la toma de imágenes de retrodispersión de electrones de alta resolución para los minerales analizados.

Los análisis se llevaron a cabo en el Departamento de Geología de la Open University, Milton Keynes, Inglaterra. El equipo utilizado fue una microsonda electrónica Cameca SX100 (Fig. A5) Se seleccionaron 15 muestras del CVSSM, las cuales abarcaron la litología presente en el área, como también el ensamblaje mineralógico que se determinó en el análisis petrográfico.

La medición se llevó a cabo acelerando una corriente de 10 nA con un voltaje de 15 kV. Los tiempos de conteo de los picos fueron de 10 a 40s y para la corrección de matriz se utilizó una rutina de PhiRhoZ. Los siguientes elementos fueron medidos utilizando una espectroscopia de longitud de onda dispersiva: K, Al, Si, Fe, Mn, Ca, P, Ti, Na and Mg. La microsonda fue calibrada usando los estándares minerales naturales de: Feldespato (K,Al,Si), hematita (Fe), bustamita (Mn, Ca), apatito (P), rutilo (Ti), jadeíta (Na) y fosterita (Mg). Los límites de detección de la microsonda electrónica Cameca SX100 dependen del elemento y de la matriz de la muestra analizada, en donde puede variar desde algunas decenas hasta algunas centenas de ppm (típicamente en orden de 0.02%).



Figura A5.- Microsonda electrónica Cameca SX100, Open University, Milton Keynes, Inglaterra

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TABLAS

Tabla A.1.- Información petrográfica de las rocas volcánicas del Complejo Volcánico de la Sierra de San Miguelito.

Muestra	Unidad	Lat (N)°	Long(W)°	Textura	Fenocristales							Matriz	Tipo de desequilibrio		
					OI	Plg	Opx	Cpx	Qz	FldK	Op		Qz+R	PI-N-S	
SLP-01	Cabras	21°55'17"	101°11'50"	P	4	10	4	6				26	50	*	*
SLP1713	Cabras	21°55'17"	101°10'50"	P	5	10	3	8				24	50	*	*
SLP1724	Cantera	22°05'26"	101°06'34"	P			3	6	24	20	6		41		
SLP1724	Cantera	22°05'26"	101°06'34"	P			3	6	24	20	6		41		
SLP-07	El Zapote	22°04'12"	101°04'45"	P		44	4	4	26	15			7		
SLP1728	El Zapote	22°05'14"	101°05'45"	P		42	6	3	26	15			8		
SLP-02	La Placa	21°46'45"	101°03'11"	P	6	68		20			5		1		
SLP-03	La Placa	21°49'47"	101°04'42"	P	3	19	7	3			1		67	*	*
SLP1714	La Placa	21°46'46"	101°02'11"	P	8	64	1	20			6		1		
SLP1720	La Placa	21°49'48"	101°05'43"	P	3	18	6	4			2		67	*	*
SLP1715	Panalillo	21°46'47"	101°02'13"	P	2	10	18	35	15	15	1		4		
SLP-04	Panalillo	21°48'45"	101°04'11"	P	2	2	9	11	32	14	4		26		
SLP1719	Panalillo	21°49'02"	101°06'08"	P			4		16	22	9		49		
SLP1723	Panalillo	22°06'06"	101°06'35"	P		43	1		38	15			3		
SLP1715	Panalillo	21°46'47"	101°02'13"	P	2	10	18	35	15	15	1		4		
SLP1716	Panalillo	21°48'46"	101°05'11"	P		2	10	12	32	14	4		26		
SLP1719	Panalillo	21°49'02"	101°06'08"	P			4		16	22	9		49		
SLP1723	Panalillo	22°06'06"	101°06'35"	P		43	1		38	15			3		
SLP-05	San José	22°01'20"	101°04'44"	P			4	2	16	21	5		52		
SLP1729	San José	22°01'23"	101°05'44"	P			6	3	16	20	5		50		
SLP-06	San Miguelito	22°05'23"	101°13'06"	P			3	1	41	15	30		10		
SLP1712	San Miguelito	22°00'03"	101°10'51"	P		3	1		46	10	36		4		
SLP1711	San Miguelito	22°06'23"	101°14'06"	P			5		40	15	30		10		

Datos modales están representados en porcentaje. Textura: P= porfírica. OI: olivino; Plg: plagioclasas; Opx: ortopiroxeno; Cpx: clinopiroxeno; Qz: cuarzo; FldK: feldespato potásico; Op: minerales opacos. Tipo de desequilibrio: Qz+R = cuarzo con aros de reacción, PI-N-S = plagioclasas con textura normal y de criba.

Tabla A.2.- Composición de elementos mayores de las rocas volcánicas del Complejo Volcánico Sierra de San Miguelito. Mediciones elaboradas en el Instituto Potosino de Investigación Científica y Tecnológica, San Luis Potosí.

Muestra	SLP-01	SLP-02	SLP1718	SLP-03	SLP17-01	SLP17-02	SLP17-06	SLP17-07
Unidad	Cabras	La Placa	La Placa	La Placa	Panalillo	Panalillo	Cantera	Panalillo
Roca(TAS)	Foidita, melaefelinita	Foidita, melaefelinita	Traquibasalto Potásico	Basalto alcalino	Riolita	Riolita	Riolita	Riolita
Long. (°W)	101°11'50"	101°03'11"	101°05'12'	101°04'42"	101°06'34"	101°09'23"	101°02'11"	101°05'43"
Lat. (°N)	21°55'17"	21°46'45"	21°49'47"	21°49'47"	22°05'26"	22°05'09"	21°46'46"	21°49'48"
n	12	12	12	12	12	12	12	12
SiO ₂	39.11 ± 1.50	37.38 ± 1.44	44.76 ± 1.5	46.34 ± 1.54	71.01 ± 1.94	72.73 ± 1.91	66.91 ± 168	71.6 ± 1.94
TiO ₂	3.9991 ± 0.039	1.7453±0.0225	1.7345 ± 0.021	2.1155 ± 0.0256	0.1137 ± 0.011	0.196 ± 0.011	0.187 ± 0.0113	0.1045 ± 0.0112
Al ₂ O ₃	15.76 ± 0.314	18.556 ± 0.305	17.139 ± 0.297	15.616 ± 0.302	11.245 ± 0.304	11.388 ± 0.301	18.445 ± 0.303	12.726 ± 0.319
Fe ₂ O ₃ ¹	14.898 ± 0.131	14.657 ± 0.12	10.977 ± 0.105	11.721 ± 0.111	1.608 ± 0.078	1.613 ± 0.078	2.385 ± 0.079	1.611 ± 0.079
MnO	0.38551 ± 0.003	0.34943 ± 0.003	0.34366 ± 0.003	0.34362 ± 0.003	0.01435 ± 0.002	0.01786 ± 0.002	0.04815 ± 0.00234	0.02757 ± 0.0023
MgO	4.513 ± 0.255	6.025 ± 0.251	5.177 ± 0.249	4.488 ± 0.249	0.841 ± 0.263	0.845 ± 0.236	0.339 ± 0.236	0.722 ± 0.236
CaO	14.355 ± 0.175	15.963 ± 0.159	12.486 ± 0.143	11.74 ± 0.148	2.134 ± 0.119	2.217 ± 0.119	1.035 ± 0.118	2.052 ± 0.119
Na ₂ O	2.082 ± 0.077	2.333 ± 0.073	2.642 ± 0.075	2.142 ± 0.074	0.482 ± 0.069	0.613 ± 0.069	2.95 ± 0.074	0.45 ± 0.069
K ₂ O	1.984 ± 0.048	1.403 ± 0.045	2.406 ± 0.048	2.255 ± 0.048	5.366 ± 0.065	4.012 ± 0.056	6.774 ± 0.065	4.842 ± 0.061
P ₂ O ₅	2.4266 ± 0.043	0.8962 ± 0.013	0.6307 ± 0.010	0.7638 ± 0.013	0.0442 ± 0.007	0.0592 ± 0.007	0.1049 ± 0.0068	0.0501 ± 0.0067
LOI	0.496	0.692	1.703	2.481	7.146	6.31	0.824	5.818
Sum	100.005	99.999	99.998	100.005	100.004	100.001	100.002	100.003
Q					44.63	50.71	20.74	47.33
Or	11.93		14.59	13.81	34.19	25.34	40.43	30.42
Ab	0.44		5.67	16.65	4.40	5.54	25.21	4.05
An	28.28	36.73	28.53	27.28	11.10	11.34	4.49	10.47
Ne	9.48	10.91	9.36	1.15				
C					0.94	2.29	4.67	3.33
Di-Mg	11.58	2.92	15.01	11.91				
Di-Fe	11.76	2.78	10.28	11.22				
Hy-Mg					2.26	2.25	0.85	1.91
Hy-Fe					1.36	1.21	1.91	1.38
Mt	3.35	3.30	3.47	2.68	0.78	0.78	1.08	0.77
Il	7.73	3.38	3.38	4.16	0.23	0.40	0.36	0.21
Ap	5.72	2.12	1.50	1.83	0.11	0.15	0.25	0.12
Mg#	41.46	49.00	54.26	47.23	60.04	60.08	28.99	56.28
FeO/MgO	2.97	2.19	1.91	2.35	1.72	1.72	6.33	2.01

Tabla A.2.- (Continuación)

Muestra	SLP17-08	SLP17-09	SLP-04	SLP17-17	SLP17-25	SLP-05	CR17-04
Unidad	Panalillo	Panalillo	Panalillo	Panalillo	Panalillo	San José	San Miguelito
Roca(TAS)	Riolita	Riolita	Riolita	Riolita	Riolita	Riolita	Riolita
Long. (°W)	101°09'24"	101°09'2"	101°04'11"	101°05'11"	101°05'45'	101°04'44"	101°12'33"
Lat. (°N)	22°05'10"	22°05'07"	21°48'45"	21°48'46"	22°05'14"	22°01'20"	22°00'48"
n	12	12	12	12	12	12	12
SiO ₂	70.98 ± 1.76	72.69 ± 1.89	78.88 ± 2.01	69.4 ± 1.81	71.98 ± 1.94	68.68 ± 1.63	74.32 ± 2.16
TiO ₂	0.2157 ± 0.0113	0.2019 ± 0.0114	0.0796 ± 0.0111	0.2344 ± 0.0115	0.1484 ± 0.0112	0.1627 ± 0.0112	0.0595 ± 0.0111
Al ₂ O ₃	14.692 ± 0.298	13.978 ± 0.309	10.895 ± 0.295	15.567 ± 0.304	10.129 ± 0.297	16.971 ± 0.291	14.004 ± 0.348
Fe ₂ O ₃ [†]	2.627 ± 0.079	2.401 ± 0.079	1.096 ± 0.078	2.484 ± 0.079	1.619 ± 0.078	2.15 ± 0.078	1.407 ± 0.079
MnO	0.02747 ± 0.00233	0.03816 ± 0.00235	0.03232 ± 0.0023	0.11047 ± 0.0023	0.0173 ± 0.0023	0.04593 ± 0.0023	0.01724 ± 0.0023
MgO	0.076 ± 0.236	0.045 ± 0.236	0.155 ± 0.236	0.737 ± 0.236	0.783 ± 0.236	0.096 ± 0.236	0.019 ± 0.236
CaO	0.606 ± 0.118	0.516 ± 0.118	0.924 ± 0.118	2.081 ± 0.119	2.397 ± 0.119	1.075 ± 0.118	0.328 ± 0.118
Na ₂ O	3.189 ± 0.077	2.946 ± 0.079	2.416 ± 0.077	3.378 ± 0.078	0.42 ± 0.069	2.822 ± 0.073	3.467 ± 0.106
K ₂ O	6.61 ± 0.064	6.458 ± 0.066	4.836 ± 0.058	4.948 ± 0.058	5.291 ± 0.063	6.647 ± 0.061	5.554 ± 0.073
P ₂ O ₅	0.0786 ± 0.0067	0.0669 ± 0.0067	0.0574 ± 0.0067	0.1068 ± 0.0068	0.0509 ± 0.0067	0.1133 ± 0.0067	0.0342 ± 0.0067
LOI	0.9	0.654	0.627	0.954	7.165	1.239	0.791
Sum	100.001	99.994	99.998	100.000	100.000	100.001	100.000
Q	25.30	29.25	44.17	24.92	46.25	24.25	32.01
Or	39.49	38.48	28.78	29.57	33.72	39.83	33.12
Ab	27.28	25.14	20.59	28.91	3.83	24.21	29.60
An	2.52	2.14	4.24	9.74	10.92	4.66	1.42
Ne							
C	1.39	1.37	0.14	1.14		3.50	1.79
Di-Mg					0.82		
Di-Fe					0.44		
Hy-Mg	0.19	0.11	0.39	1.86	1.72	0.24	0.05
Hy-Fe	2.04	1.87	0.90	2.03	1.08	1.74	1.19
Mt	1.19	1.09	0.50	1.13	0.79	0.98	0.64
Il	0.41	0.39	0.15	0.45	0.30	0.31	0.11
Ap	0.18	0.16	0.13	0.25	0.13	0.27	0.08
Mg#	7.67	5.11	28.89	46.01	58.14	11.37	3.73
FeO [†] /MgO	31.10	48.01	6.36	3.03	1.86	20.15	66.63

Tabla A.2.- (Continuación)

Muestra	SLP17-03	SLP17-04	SLP17-05	SLP-06	SLP-07	SLP17-30	SLP17-31	SM17-03
Unidad	San Miguelito	San Miguelito	San Miguelito	San Miguelito	El Zapote	Zapote	Zapote	San Miguelito
Roca(TAS)	Riolita	Riolita	Riolita	Traquidacita	Riolita	Riolita	Riolita	Riolita
Long. (°W)	101°07'00"	101°07'36"	101°07'31"	101°13'06"	101°04'45"	101°06'43"	100°58'08"	101°14'07"
Lat. (°N)	22°07'02"	22°07'04"	22°07'07"	22°05'23"	22°04'12"	22°00'40"	22°05'28"	22°06'23"
n	12	12	12	12	12	12	12	12
SiO ₂	69.43 ± 1.84	70.85 ± 1.88	68.12 ± 1.64	63.75 ± 1.67	68.57 ± 1.78	72.82 ± 1.96	72.36 ± 1.77	69.95 ± 2.02
TiO ₂	0.1305 ± 0.0112	0.1299 ± 0.0112	0.1458 ± 0.0112	0.1911 ± 0.0113	0.1436 ± 0.0112	0.1302 ± 0.0112	0.1352 ± 0.0112	0.1334 ± 0.0112
Al ₂ O ₃	17.153 ± 0.329	15.643 ± 0.317	16.264 ± 0.289	21.064 ± 0.317	18.084 ± 0.322	15.101 ± 0.326	14.863 ± 0.295	17.027 ± 0.357
Fe ₂ O ₃ ^l	1.668 ± 0.079	1.645 ± 0.078	2.052 ± 0.079	1.919 ± 0.078	1.771 ± 0.078	1.693 ± 0.079	1.627 ± 0.078	1.456 ± 0.078
MnO	0.03938 ± 0.0023	0.03955 ± 0.0024	0.0575 ± 0.0023	0.04941 ± 0.00235	0.03128 ± 0.00235	0.02178 ± 0.0023	0.03287 ± 0.0023	0.02497 ± 0.00235
MgO	0.362 ± 0.236	0.258 ± 0.236	0.194 ± 0.236	0.451 ± 0.236	0.136 ± 0.236	0.071 ± 0.236	0.046 ± 0.236	0.313 ± 0.236
CaO	0.751 ± 0.118	0.971 ± 0.118	1.828 ± 0.118	1.097 ± 0.118	0.76 ± 0.118	0.692 ± 0.118	1.018 ± 0.118	0.699 ± 0.118
Na ₂ O	2.743 ± 0.079	2.393 ± 0.078	2.764 ± 0.073	3.117 ± 0.075	3.229 ± 0.079	2.938 ± 0.082	3.346 ± 0.081	2.745 ± 0.082
K ₂ O	6.357 ± 0.069	6.519 ± 0.071	7.082 ± 0.064	7.43 ± 0.067	6.165 ± 0.064	5.812 ± 0.066	5.857 ± 0.068	6.547 ± 0.073
P ₂ O ₅	0.0531 ± 0.0067	0.0507 ± 0.0066	0.0952 ± 0.0067	0.1104 ±	0.0669 ± 0.0067	0.0608 ± 0.0067	0.0748 ± 0.0067	0.0664 ± 0.0068
LOI	1.316	1.496	1.395	0.822	1.047	0.663	0.64	1.036
Sum	100.002	99.995	99.997	100.000	100.003	100.002	99.999	99.997
Q	26.92	29.54	20.55	13.94	24.20	31.72	28.07	26.95
Or	38.11	39.16	42.51	44.33	36.86	34.62	34.88	39.14
Ab	23.55	20.58	23.75	26.63	27.64	25.06	28.53	23.49
An	3.43	4.56	8.58	4.77	3.37	3.06	4.60	3.07
Ne								
C	4.59	3.06	0.97	6.22	4.93	2.89	1.36	4.36
Di-Mg								
Di-Fe								
Hy-Mg	0.91	0.65	0.49	1.13	0.34	0.18	0.12	0.79
Hy-Fe	1.35	1.33	1.70	1.48	1.40	1.33	1.28	1.12
Mt	0.76	0.75	0.94	0.87	0.81	0.77	0.74	0.66
Il	0.25	0.25	0.28	0.37	0.28	0.25	0.26	0.26
Ap	0.12	0.12	0.22	0.26	0.16	0.14	0.17	0.16
Mg#	38.40	31.06	21.36	40.30	18.07	10.75	7.51	38.17
FeO/MgO	4.15	5.74	9.52	3.83	11.72	21.46	31.83	4.19

Tabla A.3.- Composiciones de elementos traza y tierras raras de las rocas volcánicas del Complejo Volcánico Sierra de San Miguelito. Muestras analizadas en el laboratorio ALS Chem, Canadá.

Muestra	SLP-01	SLP-02	SLP1718	SLP-03	SLP17-01	SLP17-02	SLP17-06	SLP17-07	SLP17-08	SLP17-09	SLP-04	SLP17-17
Unidad	Cabras	La Placa	La Placa	La Placa	Panalillo	Panalillo	Cantera	Panalillo	Panalillo	Panalillo	Panalillo	Panalillo
Roca(TAS)	Foidita, melaefelinita	Foidita, melaefelinita	Traquibasalto Potásico	Basalto alcalino	Riolita	Riolita	Riolita	Riolita	Riolita	Riolita	Riolita	Riolita
Long. (°W)	101°11'50"	101°03'11"	101°05'12"	101°04'42"	101°06'34"	101°09'23"	101°02'11"	101°05'43"	101°09'24"	101°09'2"	101°04'11"	101°05'11"
Lat. (°N)	21°55'17"	21°46'45"	21°49'47"	21°49'47"	22°05'26"	22°05'09"	21°46'46"	21°49'48"	22°05'10"	22°05'07"	21°48'45"	21°48'46"
Ba	376	288	691	966	392	171	219	308	70	54.7	28.5	389
Co	43	32	22	27	2					1		3
Cr	240	100	230	240		10						10
Cs	0.37	1.11	6.48	1.35	14.15	13.75	13.25	25.5	5.74	6.11	13.3	10.3
Cu	35	11	10	10	3	3	2	3	4	2	2	4
Ga	25.1	21.2	19.5	20.3	19.2	17.2	21	22.3	23.8	24.8	18.5	18.4
Hf	6	4.3	6.1	5.9	5	5.2	6	6.1	8.7	8.6	3.8	4.9
Nb	27.1	13.6	12.4	13.6	20.4	17.9	22.8	19.6	38.4	38.6	18.2	16.5
Ni	92	7	26	46	3	2	2	2				2
Pb	4	8	11	10	26	19	38	25	35	34	22	26
Rb	14.2	21.7	94	90.1	248	203	349	249	271	274	369	282
Sc	23	28	21	21	7	6	8	8	2	2	4	5
Sr	519	371	435	441	511	221	30.2	367	7.8	7.2	15.2	72.4
Ta	1.7	0.9	0.8	0.8	1.9	1.6	2	2	2.6	2.7	2.1	1.8
Th	2.1	2.56	7.14	6.24	19.8	20.9	29.9	30.5	27.8	27.4	35.2	23.8
U	0.84	0.66	1.36	1.34	4.84	4.08	7.68	6.63	4.83	5.66	9.01	6.94
V	229	161	145	138	11	8	8	5	39	33	41	13
Y	32.8	27	24.7	35.7	41.3	45.7	67	64.8	47.3	55.5	75.9	55.3
Zn	125	110	100	104	97	75	70	99	127	106	42	55
Zr	277	177	242	254	112	158	181	169	251	249	73	129
La	28.8	16.5	27	33.6	20.3	60.2	60.9	52.8	60.4	78.6	19.9	22.1
Ce	61.7	37	59	68.8	43.1	66.1	136.5	102	102.5	138	50.1	51.9
Pr	8.38	4.85	7.49	8.87	6.58	15.55	17.15	16.25	15.45	19.85	7.02	6.67
Nd	37.4	22.1	32.5	37.7	27.9	58.9	66.7	63.4	55.3	74.3	29.5	26.4
Sm	8.53	5.11	6.86	8.03	8.76	14.15	16.3	14.95	10.95	14.15	9.46	7.45
Eu	2.54	1.79	1.57	1.82	0.27	0.47	0.39	0.38	0.16	0.19	0.08	0.35
Gd	7.6	5.36	5.91	7.45	8.11	10.85	13.6	13.2	8.99	10.95	10.7	7.72
Tb	1.15	0.84	0.88	0.8	1.29	1.55	2.16	2.1	1.47	1.75	1.99	1.38
Dy	6.72	5.65	5.24	6.89	7.88	8.89	12.7	12.9	9.59	10.8	13.25	9.32
Ho	1.24	1.05	0.96	1.35	1.55	1.65	2.51	2.48	1.9	2.17	2.68	1.9
Er	3.47	2.85	2.77	3.81	4.38	4.49	6.89	6.79	5.3	5.94	7.83	5.59
Tm	0.48	0.4	0.36	0.51	0.68	0.67	1.01	1.1	0.9	0.97	1.24	0.91
Yb	2.83	2.84	2.5	3.39	4.28	4.22	6.64	6.71	5.53	6.31	7.72	5.81
Lu	0.39	0.4	0.35	0.5	0.62	0.59	0.88	0.96	0.82	0.93	1.1	0.84

Tabla A.3.- (Continuación)

Muestra	SLP17-25	SLP-05	CR17-04	SLP17-03	SLP17-04	SLP17-05	SLP-06	SLP-07	SLP17-30	SLP17-31	SM17-03
Unidad	Panalillo	San José	San Miguelito	San Miguelito	San Miguelito	San Miguelito	San Miguelito	El Zapote	Zapote	Zapote	San Miguelito
Roca(TAS)	Riolita	Riolita	Riolita	Riolita	Riolita	Riolita	Traquidacita	Riolita	Riolita	Riolita	Riolita
Long. (°W)	101°05'45"	101°04'44"	101°12'33"	101°07'00"	101°07'36"	101°07'31"	101°13'06"	101°04'45"	101°06'43"	100°58'08"	101°14'07"
Lat. (°N)	22°05'14"	22°01'20"	22°00'48"	22°07'02"	22°07'04"	22°07'07"	22°05'23"	22°04'12"	22°00'40"	22°05'28"	22°06'23"
Ba	410	119.5	6.8	21.2	34.8	167.5	255	62.1	107	47.8	254
Co	2			2		1		1	1		1
Cr		10	10								
Cs	9.05	11.75	5.18	18.1	17.35	20.4	11.9	14.8	13.25	13.5	13.05
Cu	3	2	5	2	2	2	2	3	2	2	2
Ga	15.9	23.1	42.8	24.6	23.2	23.1	22.2	23	21.6	23.1	22.7
Hf	4.6	6	8.9	4.8	4.4	4	4.8	5.8	5.6	5.4	5.6
Nb	13.3	23	100.5	26.1	24.5	25.1	20.8	23.3	21.3	25.2	21.6
Ni	1		2	1			2	1	2		2
Pb	22	35	62	37	36	33	32	40	33	38	33
Rb	183	336	397	407	390	403	319	368	330	369	315
Sc	5	7		10	10	10	7	8	8	9	8
Sr	648	19.9	5.6	16	158.5	117.5	32.8	14.5	18.1	12.9	30.2
Ta	1.2	2.1	9.7	2.5	2.4	2.5	1.9	2.2	2	2.4	2
Th	21.9	34	18.25	33.8	31.2	31	25.3	29.2	26.7	37.9	30.8
U	6.15	6.31	6.65	6.78	6.58	5.74	4.23	4.48	5.33	6.22	5.17
V	17	26					8		5		8
Y	37.6	69.7	16.9	68.7	56.3	57.8	39	14.4	26	70.1	51.3
Zn	62	64	130	71	96	75	70	76	69	65	68
Zr	132	161	135	120	108	92	121	147	144	143	152
La	55.9	76	6.6	61.7	47.2	43.5	37.9	22	44.9	73.5	54.5
Ce	88	107	16.6	132	113.5	107	86.5	63.4	90.5	127	107
Pr	14.5	20.6	1.25	17.85	13.1	12.5	10.05	3.51	13.2	20.4	16.3
Nd	53	77.4	4	67.6	49.6	46.7	37.1	10.1	49.3	76.3	62.2
Sm	10.2	15.8	1.18	15.3	11.15	10.2	8.39	1.82	10.4	16.75	13.9
Eu	0.36	0.33	0.05	0.13	0.11	0.07	0.43	0.22	0.37	0.18	0.38
Gd	7.86	13.2	1.4	12.55	9.15	8.92	7.7	1.8	7.6	13.8	11.55
Tb	1.21	2.03	0.35	1.99	1.57	1.61	1.23	0.44	1.15	2.21	1.79
Dy	7.22	12.6	2.64	13.05	10.2	10.6	7.67	2.86	6.66	13.75	10.7
Ho	1.4	2.49	0.61	2.59	2.06	2.22	1.5	0.64	1.24	2.56	2
Er	3.82	6.67	2.03	7.24	6.19	6.43	4.1	2.09	3.33	7.22	5.23
Tm	0.61	1.06	0.35	1.15	1	1.06	0.61	0.39	0.56	1.12	0.8
Yb	3.61	6.38	2.59	7.16	6.17	6.61	3.9	2.69	3.62	6.92	4.8
Lu	0.55	0.94	0.36	1.01	0.91	0.99	0.56	0.41	0.51	1.02	0.71

Tabla A.4.- Composiciones de elementos mayores, traza y tierras raras de las rocas volcánicas del Complejo Volcánico Sierra de San Miguelito. Muestras analizadas en la Universidad de Leicester, Inglaterra.

Muestra	SLP17-11	SLP17-12	SLP17-29	SLP17-28	SLP17-24	SLP17-10	SLP17-14	SLP17-20	SMB17-01	SMB17-02
Unidad	San Miguelito		San José	El Zapote	Cantera	Andesita	Basalto	La Placa	Andesita	Andesita
Roca (TAS)	Riolita	Riolita	Riolita	Riolita	Riolita	Andesita	Basalto	Andesita	Andesita	Andesita
Long. (°W)	101°14'06"	101°10'51"	101°05'44'	101°05'45"	101°06'34"	101°09'23"	101°02'11"	101°05'43"	101°09'24"	101°09'2"
Lat. (°N)	22°06'23"	22°00'03"	22°01'23"	22°05'14"	22°05'26"	22°05'09"	21°46'46"	21°49'48"	22°05'10"	22°05'07"
SiO ₂	77.63	76.53	79.15	77.07	78.01	62.88	51.23	56.89	61.86	61.27
TiO ₂	0.13	0.06	0.11	0.11	0.10	0.89	1.96	1.49	0.91	0.92
Al ₂ O ₃	12.57	13.75	11.54	12.69	12.20	16.73	16.34	15.79	15.57	16.12
Fe ₂ O ₃ ^t	1.39	1.34	1.25	1.24	1.28	6.55	11.80	8.23	6.78	7.02
MnO	0.02	0.02	0.01	0.01	0.01	0.11	0.17	0.14	0.11	0.12
MgO	0.18	0.07	0.04	0.07	0.17	2.84	6.23	5.65	3.25	3.07
CaO	0.45	0.29	0.37	0.37	0.38	5.46	8.84	6.94	5.56	6.18
Na ₂ O	2.26	3.41	2.08	2.44	1.59	2.70	2.73	2.41	2.66	2.54
K ₂ O	5.43	4.57	5.12	5.44	5.38	3.15	0.94	1.71	3.11	2.89
P ₂ O ₅	0.02	0.01	0.02	0.01	0.01	0.22	0.28	0.39	0.23	0.23
LOI	0.63	0.76	0.96	0.89	1.23	0.53	0.31	1.19	0.60	0.76
Sum	100.70	100.81	100.65	100.35	100.37	102.07	100.84	100.83	100.65	101.12
Q	42.06	38.01	46.42	41.02	47.14	17.35	1.50	12.30	16.80	16.60
Or	32.09	27.02	30.38	32.35	32.10	18.43	5.58	10.21	18.47	17.11
Ab	19.13	28.87	17.67	20.78	13.58	22.61	23.21	20.60	22.62	21.53
An	2.10	1.37	1.71	1.78	1.84	23.98	29.70	27.49	21.46	24.09
Ne	2.20	2.69	1.96	2.15	3.12					
C	1.50	1.30	1.04	1.11	1.42	12.04	22.71	18.95	12.14	11.74
Di-Mg	0.63	0.60	0.56	0.56	0.58	2.25	2.62	2.89	2.37	2.44
Di-Fe	0.25	0.11	0.21	0.21	0.19	1.67	3.74	2.86	1.74	1.75
Hy-Mg	0.05	0.02	0.05	0.02	0.02	0.50	0.65	0.91	0.54	0.53
Hy-Fe	27.11	13.05	8.42	13.95	27.61	53.04	55.24	64.14	55.53	53.25
Mt	6.95	17.22	28.12	15.94	6.78	2.08	1.70	1.31	1.88	2.06
Il	42.06	38.01	46.42	41.02	47.14	17.35	1.50	12.30	16.80	16.60
Ap	32.09	27.02	30.38	32.35	32.10	18.43	5.58	10.21	18.47	17.11
Mg#	19.13	28.87	17.67	20.78	13.58	22.61	23.21	20.60	22.62	21.53
FeO ^t /MgO	2.10	1.37	1.71	1.78	1.84	23.98	29.70	27.49	21.46	24.09
Ba	197.987	3.993	84.426	56.488	93.939	553.860	243.011	815.147	541.437	561.683
Co	1.174	0.681	0.745	0.879	0.684	16.790	33.135	27.095	2.581	17.910
Cr	11.381	11.365	34.817	12.732	14.384	103.450	92.022	238.981	20.717	108.275
Cs	9.426	3.656	4.574	11.994	7.751	3.245	0.950	0.945	0.583	2.853
Cu	2.102	4.915	2.438	3.041	2.502	8.883	13.950	10.996	7.900	8.300
Ga	22.550	48.024	21.436	23.497	22.506	19.931	25.484	19.267	19.600	20.900
Hf	6.364	11.285	6.022	6.772	6.043	6.308	4.290	6.412	5.817	5.578
Nb	22.111	124.387	21.488	23.509	23.281	13.252	13.638	12.944	12.872	12.091
Ni	1.355	1.576	2.337	1.025	2.036	6.921	3.483	42.683	6.800	3.500
Pb	26.661	51.671	28.743	29.351	27.019	10.886	4.164	8.495	10.723	9.958
Rb	213.350	290.797	144.259	250.091	181.507	66.264	15.721	60.949	50.935	42.022

Sc	15.384	8.264	1.316	7.121	2.242	24.478	34.975	24.756	21.682	23.251
Sr	27.560	2.458	14.007	16.071	23.189	276.837	311.042	390.274	252.612	294.668
Th	28.959	44.084	27.782	30.295	25.802	9.562	2.751	7.037	9.073	8.010
U	4.414	6.664	5.752	4.297	5.005	2.766	0.599	1.242	2.721	2.653
V	7.029	2.210	16.231	1.605	2.670	91.165	140.290	116.680	105.382	99.617
Y	45.551	22.808	43.756	13.331	33.373	25.781	28.204	37.810	22.901	27.477
Zn	52.222	163.443	52.300	59.009	63.020	82.635	82.039	91.125	79.200	81.000
Zr	170.859	166.115	158.115	165.630	152.773	237.990	182.985	266.598	235.385	217.393
La	40.62	6.26	50.45	19.69	22.90	24.98	16.50	33.71	21.63	22.46
Ce	96.46	11.68	80.28	58.29	58.65	56.28	36.23	68.91	46.83	46.21
Pr	11.61	1.34	15.58	3.43	7.02	7.42	4.85	9.06	6.69	6.66
Nd	48.15	5.11	67.79	11.13	30.04	32.71	23.30	41.11	5.04	30.33
Sm	10.73	1.76	14.24	2.13	6.80	7.21	5.48	8.64	1.05	6.43
Eu	0.51	0.09	0.29	0.27	0.27	1.62	1.99	2.33	0.26	1.58
Gd	10.35	2.38	12.77	2.58	7.41	7.06	5.87	8.72	6.41	6.69
Tb	1.68	0.58	2.07	0.46	1.45	1.04	0.97	1.29	0.99	1.01
Dy	9.91	4.24	12.32	3.23	9.39	6.07	5.72	7.39	5.51	6.01
Ho	1.91	0.93	2.38	0.68	1.91	1.16	1.15	1.46	1.08	1.19
Er	5.59	3.01	6.81	2.25	5.52	3.30	3.30	4.25	3.14	3.50
Tm	0.69	0.44	0.75	0.29	0.62	0.34	0.39	0.51	0.28	0.33
Yb	5.01	3.68	5.89	2.75	5.14	3.00	2.99	3.57	2.86	3.07
Lu	0.75	0.55	0.88	0.41	0.78	0.45	0.46	0.58	0.45	0.47
TREE	229.90	37.45	254.17	103.42	146.27	144.05	101.66	181.26	125.46	127.95
(La /Yb) _{CN}	5.51	1.16	5.82	4.87	3.03	5.66	3.75	6.41	5.13	4.97
(La /Sm) _{CN}	2.47	2.18	2.26	5.82	2.04	2.21	1.91	2.43	2.02	2.14
(Gd /Yb) _{CN}	1.67	0.52	1.75	0.76	1.17	1.90	1.59	1.97	1.81	1.76
(La /Yb) _{PM}	5.52	1.16	5.83	4.88	3.03	5.67	3.76	6.42	5.14	4.98
(La /Sm) _{PM}	2.48	2.19	2.27	5.84	2.04	2.22	1.92	2.43	2.02	2.15
(Gd /Yb) _{PM}	1.68	0.53	1.76	0.76	1.17	1.91	1.59	1.98	1.82	1.76
(Eu/Eu*)	0.15	0.13	0.07	0.36	0.11	0.70	1.09	0.82	0.12	0.73
(Nb/Nb*)	0.70	35.38	0.69	1.76	1.37	0.32	0.64	0.22	0.33	0.29
(Ti/Ti*)	0.04	0.06	0.03	0.13	0.04	0.39	1.02	0.51	0.43	0.42

Tabla A.4.- (Continuación)

Muestra	SLP17-15	SLP17-16	SLP17-19	SLP17-23	SLP17-13
Unidad		Panalillo			Cabras
Roca (TAS)	Riolita	Riolita	Riolita	Riolita	Basalto
Long. (°W)	101°02'13"	101°05'11"	101°06'08"	101°06'35"	101°10'50"
Lat. (°N)	21°46'47"	21°48'46"	21°49'02"	22°06'06"	21°55'17"
SiO ₂	75.93	82.28	76.62	78.25	47.76
TiO ₂	0.20	0.06	0.24	0.15	2.83
Al ₂ O ₃	12.24	9.74	11.99	11.23	15.59
Fe ₂ O ₃ ^t	2.43	0.92	2.34	1.66	13.73
MnO	0.03	0.02	0.02	0.01	0.20
MgO	0.12	0.08	0.03	0.02	5.64
CaO	1.13	0.51	0.44	0.30	8.87
Na ₂ O	2.53	2.55	2.51	2.61	3.39
K ₂ O	5.02	3.94	5.12	5.49	1.27
P ₂ O ₅	0.05	0.01	0.04	0.02	0.71
LOI	0.64	0.47	0.92	0.70	0.39
Sum	100.32	100.57	100.27	100.43	100.38
Q	38.88	50.82	41.11	41.03	0.00
Or	29.81	23.27	30.50	32.57	7.59
Ab	21.51	21.57	21.41	22.17	29.03
An	5.31	2.46	1.94	1.36	23.85
Ne	0.71	0.38	1.63	0.50	-
C	2.18	0.95	1.80	1.29	2.58
Di-Mg	1.10	0.41	1.06	0.75	3.07
Di-Fe	0.38	0.11	0.46	0.29	5.44
Hy-Mg	0.12	0.02	0.09	0.05	1.66
Hy-Fe	12.42	19.98	3.55	3.34	48.99
Mt	18.22	10.35	70.19	74.68	2.19
Il	38.88	50.82	41.11	41.03	0.00
Ap	29.81	23.27	30.50	32.57	7.59
Mg#	21.51	21.57	21.41	22.17	29.03
FeO/MgO	5.31	2.46	1.94	1.36	23.85
Ba	518.985	23.390	499.326	37.639	304.315
Co	2.026	0.853	1.115	0.593	43.183
Cr	18.347	14.968	11.565	12.659	234.352
Cs	8.323	9.347	1.596	1.843	0.333
Cu	3.919	1.670	4.026	2.862	36.438
Ga	18.087	19.209	24.249	23.983	26.990
Hf	6.578	4.276	11.171	9.380	6.068
Nb	15.766	19.431	34.377	40.481	25.028
Ni	1.861	3.599	0.904	1.674	83.257
Pb	22.503	16.621	21.886	26.949	3.066
Rb	157.642	237.321	85.363	121.351	10.359
Sc	13.620	4.740	0.896	1.349	26.952

Sr	64.817	12.268	29.233	5.279	425.171
Th	23.395	36.907	17.217	25.063	2.130
U	5.197	8.373	4.350	5.222	0.740
V	8.874	32.621	9.936	37.868	183.768
Y	49.555	75.276	35.823	39.647	32.536
Zn	53.115	27.636	93.980	85.772	98.530
Zr	239.542	85.699	428.562	274.831	266.834
La	89.25	20.22	45.42	46.26	26.45
Ce	195.11	52.90	104.44	98.96	55.06
Pr	22.48	7.46	14.60	14.99	7.73
Nd	91.68	34.91	64.06	64.10	37.28
Sm	16.08	10.37	12.40	12.52	8.18
Eu	1.23	0.11	0.94	0.16	2.75
Gd	15.27	11.60	11.87	11.87	8.42
Tb	2.00	2.30	1.86	1.91	1.24
Dy	11.05	14.47	10.65	11.44	7.00
Ho	1.99	2.98	2.06	2.26	1.30
Er	5.57	8.94	5.95	6.63	3.47
Tm	0.60	1.08	0.63	0.69	0.38
Yb	4.63	8.56	5.35	6.17	2.79
Lu	0.70	1.24	0.82	0.92	0.45
TREE	439.75	159.82	265.58	262.77	152.84
(La /Yb) _{CN}	3.59	1.21	2.24	2.29	2.05
(La /Sm) _{CN}	2.67	1.10	1.79	1.56	2.44
(Gd /Yb) _{CN}	13.13	1.61	5.78	5.11	6.44
(La /Yb) _{PM}	3.60	1.21	2.25	2.30	2.06
(La /Sm) _{PM}	2.68	1.10	1.80	1.56	2.44
(Gd /Yb) _{PM}	0.25	0.03	0.24	0.04	1.03
(Eu/Eu*)	0.21	1.63	0.69	1.53	0.84
(Nb/Nb*)	0.04	0.01	0.06	0.04	1.07
(Ti/Ti*)	13.10	1.60	5.77	5.10	6.43

Tabla A.5.- Datos isotópicos de roca total de Sr-Nd de las rocas volcánicas del Complejo Volcánico de la Sierra de San Miguelito.

Muestra	Unidad	Edad	Rb (ppm)	Sr (ppm)	$^{87}\text{Rb}/^{86}\text{Sr}$	$^{87}\text{Sr}/^{86}\text{Sr}$	$\pm 2\sigma$	$^{87}\text{Sr}/^{86}\text{Sr}_{(i)}$	Sm (ppm)	Nd (ppm)	$^{147}\text{Sm}/^{144}\text{Nd}$	$^{143}\text{Nd}/^{144}\text{Nd}$	$\pm 2\sigma$	$^{143}\text{Nd}/^{144}\text{Nd}_{(i)}$	ϵNd_t
SLP17-13	Cabras	22.21	10.36	425.17	0.07049	0.70346	6	0.70344	8.18	37.28	0.13271	0.51289	10	0.51287	+5.2
SLP17-14	La Placa	30.13	15.72	311.04	0.14622	0.70502	6	0.70496	5.48	23.30	0.14210	0.51266	18	0.51264	+0.9
SLP17-10	La Placa	32.67	66.26	276.84	0.69248		–		7.21	32.71	0.13330	0.51249	9	0.51246	-2.5
SLP17-20	La Placa	27.5	60.95	390.27	0.45181	0.70557	6	0.70540	8.64	41.11	0.12707	0.51252	7	0.51249	-2.0
SMB17-01	La Placa	30	50.94	252.61	0.58333	0.70606	7	0.70581	1.05	5.04	0.12628	0.51249	13	0.51247	-2.4
SMB17-02	La Placa	30	42.02	294.67	0.41257	0.70592	20	0.70575	6.43	30.33	0.12822	0.51252	7	0.51249	-1.8
SLP17-11	San Miguelito	33.17	213.35	27.56	22.39592	0.71995	7	0.70940	10.73	48.15	0.13474	0.51249	6	0.51246	-2.5
SLP17-12	San Miguelito	33	290.80	2.46	342.26587	–	–	–	1.76	5.11	0.20808	0.51262	8	0.51258	-0.2
SLP17-15	Panalillo	28.65	157.64	64.82	7.03621	0.71005	6	0.70719	16.08	91.68	0.10608	0.51248	8	0.51245	-2.7
SLP17-16	Panalillo	28.65	237.32	12.27	55.96519	0.73354	7	0.71078	10.37	34.91	–	–	–	–	–
SLP17-19	Panalillo	31.05	85.36	29.23	8.44796	0.71258	8	0.70886	12.40	64.06	0.11702	0.51254	8	0.51252	-1.4
SLP17-23	Panalillo	29.4	121.35	5.28	66.50392	0.74749	11	0.71973	12.52	64.10	0.11812	0.51254	8	0.51251	-1.4
SLP17-24	Cantera	32.1	181.51	23.19	22.64472	0.72170	7	0.71138	6.80	30.04	0.13698	0.51249	8	0.51247	-2.3
SLP17-28	El Zapote	33.48	250.09	16.07	45.02054	0.73799	6	0.71659	2.13	11.13	0.11587	0.51249	8	0.51247	-2.3
SLP17-29	San José	33.95	144.26	14.01	29.79568	–	–	–	14.24	67.79	0.12706	0.51248	9	0.51245	-2.6

Tabla A.6.-Datos isotópicos de roca total de Pb de las rocas volcánicas del Complejo Volcánico de la Sierra de San Miguelito.

Muestra	Unidad	Edad	Th (ppm)	U (ppm)	Pb (ppm)	$^{206}\text{Pb}/^{204}\text{Pb}$	$^{207}\text{Pb}/^{204}\text{Pb}$	$^{208}\text{Pb}/^{204}\text{Pb}$	$^{206}\text{Pb}/^{204}\text{Pb}_{(i)}$	$^{207}\text{Pb}/^{204}\text{Pb}_{(i)}$	$^{208}\text{Pb}/^{204}\text{Pb}_{(i)}$
SLP17-13	Cabras	22.21	2.13	0.74	3.066	18.928	15.614	38.571	18.925	15.329	38.571
SLP17-14	La Placa	30.13	2.751	0.599	4.164	18.965	15.665	38.909	18.962	15.303	38.909
SLP17-10	La Placa	32.67	9.562	2.766	10.886	18.939	15.667	38.902	18.934	15.149	38.902
SLP17-20	La Placa	27.5	7.037	1.242	8.495	18.894	15.664	38.864	18.891	15.248	38.864
SMB17-01	La Placa	30	9.073	2.721	10.723	18.936	15.666	38.894	18.932	15.205	38.893
SMB17-02	La Placa	30	8.01	2.653	9.958	18.938	15.665	38.889	18.933	15.227	38.889
SLP17-11	San Miguelito	33.17	28.959	4.414	26.661	18.900	15.666	38.885	18.897	15.016	38.885
SLP17-12	San Miguelito	33	44.084	6.664	51.671	18.905	15.661	38.831	18.903	15.154	38.831
SLP17-15	Panalillo	28.65	23.395	5.197	22.503	18.909	15.665	38.881	18.906	15.122	38.881
SLP17-16	Panalillo	(28.65)	36.907	8.373	16.621	19.013	15.680	39.016	19.005	14.521	39.016
SLP17-19	Panalillo	31.05	17.217	4.35	21.886	18.896	15.660	38.858	18.892	15.217	38.858
SLP17-23	Panalillo	29.4	25.063	5.222	26.949	18.897	15.659	38.861	18.894	15.162	38.861
SLP17-24	Cantera	32.1	25.802	5.005	27.019	18.907	15.666	38.889	18.903	15.112	38.889
SLP17-28	El Zapote	33.48	30.295	4.297	29.351	18.892	15.668	38.876	18.889	15.045	38.876
SLP17-29	San José	33.95	27.782	5.752	28.743	18.922	15.669	38.913	18.918	15.078	38.913

Tabla A.7.- Datos de edades $^{40}\text{Ar}/^{39}\text{Ar}$ age data para las unidades del Complejo Volcánico de la Sierra de San Miguelito

Step no	Power (%)	^{40}Ar	$\pm\sigma_{40}$	^{39}Ar	$\pm\sigma_{39}$	^{38}Ar	$\pm\sigma_{38}$	^{37}Ar	$\pm\sigma_{37}$	^{36}Ar	$\pm\sigma_{36}$	$^{40}\text{Ar}/^{39}\text{Ar}$	$\pm\sigma$	Age (Ma)	$\pm\sigma$ (Ma)
SLP17-11 (San Miguelito) sanidine, grain size 0.1-0.3 mm, 0.1062mg, $J=0.009790695 \pm 4.89535E-05$															
1	1.2	0.08104	0.00434	0.00483	0.00050	0.00001	0.00025	-0.00151	0.00403	0.00015	0.00021	7.27882	13.16002	124.00983	216.67689
2	1.5	0.22769	0.00471	0.04797	0.00056	0.00137	0.00025	0.01221	0.00404	0.00097	0.00021	-1.29137	-1.33035	-22.92000	23.76272
3	2.0	0.68898	0.00690	0.26534	0.00390	0.00743	0.00028	0.05568	0.00404	0.00218	0.00022	0.14526	0.24824	2.55999	4.37181
4	2.6	0.21664	0.00457	0.09143	0.00072	0.00150	0.00026	0.00931	0.00404	0.00051	0.00021	0.70128	0.69783	12.32573	12.22357
5	3.4	1.09766	0.00897	0.51532	0.00399	0.01106	0.00035	0.06600	0.00404	0.00201	0.00021	0.96793	0.12588	16.99040	2.20081
6	4.2	0.87296	0.01186	0.42920	0.00239	0.00755	0.00028	0.04290	0.00405	0.00183	0.00021	0.75956	0.15008	13.34636	2.62822
7	5.2	2.53212	0.00988	1.23526	0.00256	0.01399	0.00042	0.02160	0.00539	0.00048	0.00031	1.93330	0.07474	33.77807	1.30450
8	6.5	2.10912	0.00600	1.01864	0.00242	0.01205	0.00040	0.02112	0.00539	0.00061	0.00031	1.89104	0.09029	33.04634	1.57202
9	8.0	4.04445	0.00666	1.92434	0.01180	0.02295	0.00044	0.03731	0.00539	0.00140	0.00031	1.88482	0.04965	32.93863	0.87514

	MSWD	% ^{39}Ar	Initial Ratio
Plateau age	33.17	0.67	0.15
Isochron age	34.8	5.3	13
			80.1
			na
			na
			86±66

Step no	Power (%)	⁴⁰ Ar	±σ ₄₀	³⁹ Ar	±σ ₃₉	³⁸ Ar	±σ ₃₈	³⁷ Ar	±σ ₃₇	³⁶ Ar	±σ ₃₆	⁴⁰ Ar/ ³⁹ Ar	±σ	Age (Ma)	±σ (Ma)
SLP17-29 (San Jose) sanidine, grain size 0.1-0.3 mm, 0.0328mg, J=0.00973613± 4.86807E-05															
1	0.5	0.00888	0.00154	0.00080	0.00018	0.00001	0.00002	-0.00044	0.00049	0.00004	0.00002	-5.52252	-6.86829	-99.70682	127.49589
2	0.9	0.33317	0.00240	0.12370	0.00049	0.00150	0.00007	0.00254	0.00049	0.00036	0.00002	1.81540	0.06275	31.60848	1.09423
3	1.2	1.21558	0.00555	0.19103	0.00150	0.00273	0.00008	0.00476	0.00049	0.00290	0.00005	1.82595	0.08747	31.79064	1.51775
4	1.5	0.66775	0.01100	0.08937	0.00155	0.00135	0.00003	0.00295	0.00014	0.00163	0.00005	2.01781	0.21708	35.09868	3.74348
5	2.0	0.56452	0.00504	0.15207	0.00171	0.00189	0.00007	0.00376	0.00014	0.00088	0.00003	1.97997	0.06417	34.44671	1.11894
6	2.7	0.96233	0.01134	0.37882	0.00504	0.00437	0.00009	0.00638	0.00014	0.00066	0.00003	2.01889	0.04499	35.11721	0.79422
7	3.4	0.79170	0.00781	0.26101	0.00282	0.00312	0.00007	0.00359	0.00014	0.00096	0.00003	1.93237	0.05334	33.62618	0.93451
8	4.4	1.30307	0.01938	0.38188	0.00436	0.00473	0.00008	0.00594	0.00014	0.00188	0.00004	1.94107	0.06482	33.77612	1.12991
9	5.6	1.09343	0.00769	0.48988	0.00343	0.00572	0.00014	0.00612	0.00014	0.00044	0.00003	1.96282	0.02606	34.15118	0.47991
10	7.0	1.81151	0.01455	0.74657	0.00512	0.00885	0.00016	0.01047	0.00014	0.00126	0.00004	1.92234	0.02916	33.45333	0.52933
11	8.5	1.77599	0.01152	0.75081	0.00150	0.00914	0.00010	0.00966	0.00014	0.00097	0.00002	1.97942	0.01753	34.43713	0.34689
12	10.0	1.53159	0.00675	0.59982	0.00378	0.00725	0.00011	0.01120	0.00014	0.00131	0.00005	1.90119	0.03087	33.08856	0.55710
13	12.0	3.04731	0.01155	1.37643	0.00426	0.01596	0.00015	0.03779	0.00014	0.00132	0.00006	1.92688	0.01695	33.53151	0.33619
14	14.0	1.87358	0.00615	0.88109	0.00505	0.01009	0.00008	0.03943	0.00014	0.00047	0.00003	1.96619	0.01756	34.20913	0.34680

	MSWD	% ³⁹Ar	Initial Ratio
Plateau age	33.95	0.44	1.3
Isochron age	34.09	0.48	1.7
			98.1
			na
			292.7±8.7

Step no	Power (%)	⁴⁰ Ar	±σ ₄₀	³⁹ Ar	±σ ₃₉	³⁸ Ar	±σ ₃₈	³⁷ Ar	±σ ₃₇	³⁶ Ar	±σ ₃₆	⁴⁰ Ar/ ³⁹ Ar	±σ	Age (Ma)	±σ (Ma)
SLP17-28 (El Zapote) sanidine, grain size 0.1-0.3 mm, 0.0212mg, J=0.009768395 ± 4.8842E-05															
1	0.5	0.00738	0.00108	0.00096	0.00025	0.00002	0.00002	0.00135	0.00046	0.00003	0.00002	-3.10003	-7.45942	-55.47598	135.56241
2	0.9	0.07025	0.00118	0.01356	0.00026	0.00021	0.00002	0.00173	0.00046	0.00015	0.00002	1.77793	0.52476	31.06330	9.09123
3	1.2	0.11742	0.00153	0.04273	0.00037	0.00051	0.00003	0.00596	0.00047	0.00014	0.00002	1.74555	0.16836	30.50239	2.92120
4	1.5	0.41147	0.00338	0.20242	0.00207	0.00251	0.00008	0.00953	0.00047	0.00009	0.00002	1.89642	0.03463	33.11472	0.62127
5	2.0	1.06346	0.00621	0.54646	0.00264	0.00651	0.00011	0.01847	0.00047	0.00020	0.00002	1.83678	0.01929	32.08245	0.36982
6	2.7	1.20920	0.00867	0.49321	0.00560	0.00585	0.00011	0.01396	0.00047	0.00103	0.00003	1.82740	0.03353	31.92016	0.60167
7	3.4	0.87898	0.01766	0.39921	0.00621	0.00481	0.00011	0.00965	0.00023	0.00030	0.00006	1.97935	0.06893	34.54895	1.20389
8	4.4	2.65836	0.02502	1.25026	0.01111	0.01443	0.00019	0.02536	0.00023	0.00085	0.00006	1.92248	0.03015	33.56555	0.54735
9	5.6	3.41673	0.02704	1.70051	0.00807	0.01961	0.00030	0.03241	0.00023	0.00047	0.00006	1.92647	0.02093	33.63460	0.39854
10	7.0	3.50570	0.02306	1.70700	0.00668	0.01977	0.00015	0.02880	0.00023	0.00070	0.00006	1.93088	0.01886	33.71082	0.36653
11	8.5	3.33876	0.02567	1.67506	0.01357	0.01899	0.00031	0.02633	0.00023	0.00061	0.00006	1.88396	0.02393	32.89902	0.44498
12	10.0	3.38770	0.02731	1.66010	0.00553	0.01956	0.00028	0.02540	0.00023	0.00075	0.00006	1.90519	0.02083	33.26649	0.39626
13	12.0	3.25444	0.02156	1.59914	0.00881	0.01835	0.00018	0.02407	0.00023	0.00056	0.00006	1.92988	0.02067	33.69363	0.39459
14	14.0	4.33748	0.01755	2.02317	0.00948	0.02328	0.00013	0.02936	0.00023	0.00156	0.00006	1.91337	0.01544	33.40791	0.31417

	MSWD	% ³⁹Ar	Initial Ratio
Plateau age	33.48	0.43	0.61
Isochron age	33.4	0.68	2.2
			90.2
			na
			279±20

Step no	Power (%)	⁴⁰ Ar	±σ ₄₀	³⁹ Ar	±σ ₃₉	³⁸ Ar	±σ ₃₈	³⁷ Ar	±σ ₃₇	³⁶ Ar	±σ ₃₆	⁴⁰ Ar/ ³⁹ Ar	±σ	Age (Ma)	±σ (Ma)
SLP17-24 (Cantera) sanidine, grain size 0.1-0.3 mm, 0.028mg, J=0.009832925 ±4.91646E-05															
1	0.2	-0.00179	0.00225	-0.00045	0.00052	-0.00003	0.00003	-0.00042	0.00017	0.00000	0.00003	4.75258	18.38306	82.39704	311.54490
2	0.5	0.00955	0.00225	0.00007	0.00052	0.00001	0.00003	0.00026	0.00017	0.00004	0.00003	-36.79198	-283.81485	-810.14036	7888.55861
3	0.9	0.21885	0.00503	0.03799	0.00067	0.00060	0.00004	0.00775	0.00018	0.00057	0.00003	1.28869	0.24179	22.71693	4.23707
4	1.2	0.71505	0.01081	0.33857	0.00522	0.00396	0.00011	0.00518	0.00018	0.00028	0.00003	1.86532	0.04853	32.78983	0.86081
5	1.5	1.19073	0.01459	0.50088	0.00587	0.00610	0.00016	0.00798	0.00018	0.00092	0.00004	1.82948	0.04435	32.16539	0.78906
6	2.0	2.22496	0.01774	1.03458	0.00838	0.01228	0.00023	0.01616	0.00018	0.00117	0.00003	1.81387	0.02462	31.89338	0.45723
7	2.7	1.70031	0.01269	0.83209	0.00663	0.00967	0.00014	0.01247	0.00018	0.00056	0.00003	1.84329	0.02307	32.40593	0.43289
8	3.4	4.36778	0.00777	1.99192	0.00445	0.02478	0.00026	0.02515	0.00018	0.00244	0.00005	1.82636	0.00968	32.11092	0.23190
9	4.4	1.72001	0.00812	0.89714	0.00388	0.01065	0.00015	0.01080	0.00018	0.00023	0.00003	1.84126	0.01476	32.37054	0.30306
10	5.6	1.98469	0.00727	1.04858	0.00378	0.01233	0.00020	0.01922	0.00018	0.00029	0.00003	1.81130	0.01357	31.84843	0.28439
11	6.5	3.49835	0.03090	0.81744	0.00902	0.01086	0.00013	0.00834	0.00018	0.00716	0.00009	1.66495	0.05367	29.29592	0.94795
12	7.0	1.69258	0.01521	0.88070	0.00594	0.01037	0.00015	0.00868	0.00018	0.00025	0.00002	1.83751	0.02219	32.30532	0.41854
13	7.7	1.23284	0.00987	0.63760	0.00257	0.00768	0.00011	0.00868	0.00018	0.00011	0.00003	1.88261	0.02099	33.09095	0.40072
14	8.5	1.32019	0.00771	0.68473	0.00216	0.00819	0.00008	0.01026	0.00018	0.00017	0.00003	1.85462	0.01689	32.60343	0.33572

	MSWD	% ³⁹ Ar	Initial Ratio
Plateau age	32.1	0.38	1.6
Isochron age	32.5	0.39	1.3
			86
			na
			277.4±9.4

Step no	Power (%)	⁴⁰ Ar	±σ ₄₀	³⁹ Ar	±σ ₃₉	³⁸ Ar	±σ ₃₈	³⁷ Ar	±σ ₃₇	³⁶ Ar	±σ ₃₆	⁴⁰ Ar/ ³⁹ Ar	±σ	Age (Ma)	±σ (Ma)
SLP17-10 (La Placa) whole rock, grain size 0.3-0.5 mm, 0.0382 mg, J=0.009703865±0.0000485193															
1	0.3	0.03155	0.0023	0.00028	0.0003	0.00003	0.00002	-0.00043	0.0008	0.00005	0.00002	61.73442	75.8479	846.87	830.3885
2	0.6	1.00104	0.0121	0.11786	0.0012	0.00195	0.0001	0.08319	0.0005	0.00265	0.00003	1.77302	0.1344	30.78	2.3181
3	0.9	3.21498	0.0189	0.99477	0.0061	0.01312	0.0001	0.72982	0.0005	0.00483	0.0001	1.78178	0.0328	30.93	0.5855
4	1.2	3.75210	0.0131	1.59263	0.0085	0.01932	0.0001	1.16933	0.0005	0.00284	0.0001	1.82443	0.0161	31.66	0.3178
5	1.5	3.78826	0.0108	1.95757	0.0121	0.02224	0.0002	1.32950	0.0005	0.00028	0.00003	1.89207	0.0139	32.82	0.2888
6	2.1	5.01006	0.0217	2.57457	0.0081	0.02939	0.0003	1.61104	0.0005	0.00050	0.00003	1.88822	0.0110	32.76	0.2497
7	2.7	4.69989	0.0222	2.46109	0.0099	0.02744	0.0002	1.54289	0.0005	0.00026	0.00003	1.87861	0.0122	32.59	0.2643
8	3.3	4.15327	0.0275	2.15567	0.0096	0.02463	0.0002	1.41606	0.0005	0.00023	0.00003	1.89485	0.0157	32.87	0.3153
9	4.4	3.00030	0.0183	1.56726	0.0075	0.01786	0.0002	1.02738	0.0005	0.00030	0.00003	1.85668	0.0160	32.21	0.3185
10	5.7	2.48388	0.0137	1.27463	0.0053	0.01510	0.0002	0.92318	0.0005	0.00046	0.00003	1.84087	0.0154	31.94	0.3082
11	7.7	2.77713	0.0193	1.41388	0.0078	0.01679	0.0002	1.06198	0.0005	0.00054	0.00003	1.84940	0.0185	32.09	0.3552

	MSWD	% ³⁹Ar	Initial Ratio
Plateau age	32.67	0.39	1.01
Isochron age	32.51	0.48	2.4
			66.5
			na
			290±9.6

Step no	Power (%)	⁴⁰ Ar	±σ ₄₀	³⁹ Ar	±σ ₃₉	³⁸ Ar	±σ ₃₈	³⁷ Ar	±σ ₃₇	³⁶ Ar	±σ ₃₆	⁴⁰ Ar/ ³⁹ Ar	±σ	Age (Ma)	±σ (Ma)
<i>SLP17-14 (La Placa) whole rock, grain size 0.3-0.5 mm, 0.0565mg, J=0.009639335 ± 4.81967E-05</i>															
1	0.3	0.02598	0.00073	0.00029	0.00003	0.00002	0.00002	0.00072	0.00011	0.00013	0.00002	-40.37230	-23.92136	-889.27221	681.04773
2	0.6	0.20920	0.00154	0.00680	0.00009	0.00026	0.00003	0.02014	0.00011	0.00075	0.00001	-2.19379	-0.66779	-38.55931	11.86549
3	0.9	0.23900	0.00233	0.01835	0.00030	0.00036	0.00003	0.05965	0.00011	0.00067	0.00002	2.12006	0.38822	36.49646	6.61845
4	1.2	0.49609	0.00198	0.04977	0.00039	0.00087	0.00004	0.18338	0.00011	0.00135	0.00005	1.88467	0.30904	32.48046	5.28072
5	1.5	1.22369	0.00389	0.35070	0.00191	0.00443	0.00006	1.27254	0.00011	0.00181	0.00005	1.94940	0.04610	33.58567	0.80437
6	2.1	1.01341	0.00470	0.42019	0.00214	0.00520	0.00010	1.44765	0.00011	0.00070	0.00004	1.91272	0.03289	32.95943	0.58494
7	2.7	1.27038	0.00730	0.65120	0.00324	0.00777	0.00010	1.99388	0.00139	0.00038	0.00003	1.77739	0.02117	30.64726	0.39249
8	3.3	1.34030	0.00625	0.71380	0.00366	0.00850	0.00009	2.50620	0.00139	0.00032	0.00003	1.74279	0.01654	30.05553	0.31967
9	4.4	1.24149	0.00965	0.66867	0.00566	0.00793	0.00013	2.35863	0.00139	0.00028	0.00004	1.73089	0.02819	29.85202	0.50440
10	5.7	1.27088	0.00694	0.69090	0.00435	0.00832	0.00014	2.78995	0.00139	0.00026	0.00002	1.72824	0.01698	29.80674	0.32587
11	7.7	1.78673	0.00873	0.96382	0.00607	0.01148	0.00023	3.95772	0.00139	0.00031	0.00003	1.75844	0.01784	30.32314	0.34009

			MSWD	% ³⁹ Ar	Initial Ratio
Plateau age	30.13	0.42	1.02	81.5	na
Isochron age	30.47	0.99	3.5	na	311±17

Step no	Power (%)	⁴⁰ Ar	±σ ₄₀	³⁹ Ar	±σ ₃₉	³⁸ Ar	±σ ₃₈	³⁷ Ar	±σ ₃₇	³⁶ Ar	±σ ₃₆	⁴⁰ Ar*/ ³⁹ Ar	±σ	Age (Ma)	±σ (Ma)
SLP17-20 (La Placa) whole rock, grain size 0.3-0.5 mm, 0.0396mg, J=0.00960707± 4.80354E-05															
1	0.3	0.24840	0.00288	0.03033	0.00038	0.00064	0.00004	0.01713	0.00084	0.00073	0.00003	0.97190	0.33630	16.76678	5.77541
2	0.6	2.95426	0.01134	0.56369	0.00244	0.00959	0.00013	0.17396	0.00084	0.00757	0.00011	1.23057	0.06226	21.20301	1.07161
3	0.9	4.60960	0.03390	1.28911	0.00837	0.02091	0.00028	0.50612	0.00084	0.00941	0.00006	1.39559	0.03124	24.02744	0.54748
4	1.2	5.21865	0.03792	2.32888	0.01178	0.03682	0.00042	1.44776	0.00084	0.00654	0.00006	1.40189	0.01943	24.13520	0.35323
5	1.5	4.65527	0.02151	2.04198	0.01039	0.03270	0.00026	1.90737	0.00084	0.00512	0.00008	1.53084	0.01767	26.33920	0.32891
6	2.1	3.20456	0.01473	1.42063	0.00596	0.02313	0.00021	1.94331	0.00084	0.00334	0.00006	1.55321	0.01780	26.72119	0.33168
7	2.7	5.98247	0.05629	2.14008	0.01690	0.03594	0.00035	3.54177	0.00084	0.00901	0.00011	1.53858	0.03284	26.47137	0.57611
8	3.3	3.35366	0.01395	1.28933	0.00319	0.02079	0.00026	2.93116	0.00084	0.00437	0.00005	1.58892	0.01664	27.33097	0.31471
9	4.4	2.75433	0.00829	0.95072	0.00308	0.01552	0.00014	3.02233	0.00084	0.00408	0.00007	1.61682	0.02460	27.80721	0.44200
10	5.7	1.57253	0.00621	0.57229	0.00192	0.00904	0.00015	2.09047	0.00084	0.00206	0.00004	1.67109	0.02517	28.73319	0.45244

	MSWD	% ³⁹ Ar	Initial Ratio
Plateau age	-	-	na
Isochron age	28	2	20

Step no	Power (%)	⁴⁰ Ar	±σ ₄₀	³⁹ Ar	±σ ₃₉	³⁸ Ar	±σ ₃₈	³⁷ Ar	±σ ₃₇	³⁶ Ar	±σ ₃₆	⁴⁰ Ar/ ³⁹ Ar	±σ	Age (Ma)	±σ (Ma)
SLP17-15 (Panalillo) sanidine, grain size 0.1-0.3 mm, 0.031mg, J=0.009805594 ± 4.9028E-05															
1	0.6	0.06139	0.00547	0.00076	0.00182	0.00011	0.00081	0.00196	0.01193	0.00024	0.00055	-12.12323	-217.39499	-228.31733	4364.56779
2	0.9	0.44042	0.00591	0.00358	0.00182	0.00016	0.00081	-0.00234	0.01194	0.00128	0.00055	16.49613	46.67272	270.48744	710.68724
3	1.2	1.83239	0.01375	0.07646	0.00191	0.00291	0.00081	0.02519	0.01194	0.00645	0.00055	-1.22180	-2.15707	-21.74418	38.62148
4	1.5	0.59943	0.00678	0.06489	0.00187	0.00180	0.00081	0.01472	0.01195	0.00226	0.00055	-1.17485	-2.53015	-20.90370	45.28017
5	2.0	0.81205	0.00753	0.13470	0.00192	0.00388	0.00081	0.03002	0.01195	0.00282	0.00055	-0.21980	-1.22387	-3.89247	21.69710
6	2.6	1.30151	0.00758	0.42205	0.00301	0.00786	0.00082	0.05572	0.01196	0.00242	0.00055	1.37028	0.38920	24.07886	6.79479
7	3.3	0.99706	0.00822	0.47237	0.00274	0.00773	0.00082	0.06322	0.01197	0.00138	0.00055	1.23840	0.34778	21.77538	6.07935
8	4.2	0.91676	0.00696	0.44748	0.00338	0.00794	0.00082	0.07602	0.01197	0.00122	0.00055	1.23683	0.36770	21.74801	6.42753
9	5.2	0.72760	0.00504	0.37326	0.00134	0.00437	0.00012	0.03228	0.00142	0.00028	0.00009	1.72864	0.07036	30.32339	1.23308
10	6.5	1.06992	0.00566	0.52859	0.00291	0.00652	0.00018	0.03859	0.00142	0.00062	0.00008	1.67153	0.04897	29.32976	0.86455
11	8.0	1.37878	0.00863	0.68703	0.00509	0.00838	0.00016	0.07806	0.00142	0.00094	0.00009	1.59675	0.04115	28.02777	0.73012
12	10.0	2.17588	0.01757	1.04632	0.00559	0.01265	0.00016	0.09548	0.00142	0.00149	0.00008	1.65464	0.03035	29.03570	0.54759
13	12.0	2.73692	0.03948	0.93503	0.01304	0.01219	0.00014	0.07494	0.00142	0.00436	0.00010	1.53346	0.05718	26.92508	1.00537
14	14.0	2.06103	0.00493	0.97727	0.00232	0.01172	0.00020	0.05451	0.00142	0.00109	0.00009	1.77596	0.02707	31.14635	0.49543

	MSWD	% ³⁹Ar	Initial Ratio
Plateau age	28.65	0.37	1.4
Isochron age	30.6	2	3.2
			83.4
			na
			256±61

Step no	Power (%)	⁴⁰ Ar	±σ ₄₀	³⁹ Ar	±σ ₃₉	³⁸ Ar	±σ ₃₈	³⁷ Ar	±σ ₃₇	³⁶ Ar	±σ ₃₆	⁴⁰ Ar*/ ³⁹ Ar	±σ	Age (Ma)	±σ (Ma)
SLP17-16 (Panalillo) sanidine, grain size 0.1-0.3 mm, 0.0269mg, J=0.009820493 ± 4.91025E-05															
1	0.5	0.02426	0.00907	0.00232	NaN	0.00003	0.00002	0.00000	0.00015	0.00007	NaN	2.09582	3.91234	36.75451	67.91703
2	0.9	2.40244	0.02241	0.01263	0.00028	0.00167	0.00005	0.00176	0.00015	0.00800	0.00010	0.99850	2.95614	17.60420	51.86521
3	1.2	2.65762	0.02510	0.02576	0.00053	0.00198	0.00009	0.00228	0.00015	0.00909	0.00011	-2.23650	-1.59661	-40.06552	28.92304
4	1.5	2.15042	0.02180	0.06898	0.00051	0.00213	0.00008	0.00913	0.00015	0.00719	0.00010	0.04354	0.53577	0.77132	9.48806
5	2.0	4.93083	0.03821	0.30654	0.00289	0.00634	0.00011	0.01350	0.00015	0.01545	0.00021	1.03626	0.24343	18.26656	4.27033
6	2.7	7.10684	0.02717	0.52825	0.00222	0.01021	0.00005	0.03844	0.00015	0.02155	0.00010	1.27106	0.07659	22.37987	1.34482
7	3.4	8.17871	0.04307	1.01427	0.00792	0.01590	0.00019	0.05275	0.00029	0.02419	0.00014	0.94426	0.05992	16.65228	1.05517
8	4.4	10.59831	0.05862	1.53740	0.00695	0.02271	0.00023	0.07512	0.00029	0.02864	0.00028	1.33181	0.06680	23.44260	1.17394
9	5.6	4.31153	0.01491	0.66515	0.00301	0.00994	0.00014	0.03266	0.00029	0.01134	0.00008	1.39135	0.04373	24.48364	0.77391
10	7.0	3.63643	0.00890	0.55492	0.00163	0.00798	0.00008	0.03921	0.00029	0.00969	0.00007	1.33983	0.04251	23.58288	0.75254
11	8.5	3.23238	0.01429	0.47113	0.00242	0.00687	0.00011	0.01833	0.00029	0.00872	0.00011	1.33803	0.07737	23.55146	1.35808
12	10.0	4.81527	0.01324	0.77867	0.00293	0.01140	0.00014	0.04136	0.00029	0.01244	0.00010	1.41455	0.04297	24.88907	0.76102
13	12.0	5.87268	0.01458	1.06564	0.00418	0.01474	0.00015	0.04920	0.00029	0.01441	0.00011	1.47455	0.03467	25.93719	0.61899
14	14.0	3.31117	0.00987	0.86639	0.00197	0.01115	0.00016	0.02512	0.00029	0.00642	0.00007	1.60830	0.02779	28.27137	0.50464

	MSWD	% ³⁹Ar	Initial Ratio
Plateau age	24.15	0.82	0.58
Isochron age	29.6	3.5	7.5
			51.5
			na
			282±12

Step no	Power (%)	⁴⁰ Ar	±σ ₄₀	³⁹ Ar	±σ ₃₉	³⁸ Ar	±σ ₃₈	³⁷ Ar	±σ ₃₇	³⁶ Ar	±σ ₃₆	⁴⁰ Ar/ ³⁹ Ar	±σ	Age (Ma)	±σ (Ma)
SLP17-19 (Panalillo) sanidine, grain size 0.1-0.3 mm, 0.0764mg, J=0.009835392 ± 4.9177E-05															
1	1.2	1.98129	0.00733	0.04116	0.00096	0.00160	0.00009	0.00143	0.00068	0.00644	0.00006	1.41969	0.49982	25.01641	8.74755
2	1.5	0.68167	0.00710	0.02308	0.00095	0.00070	0.00006	0.00076	0.00068	0.00216	0.00005	1.58525	0.67412	27.91129	11.77865
3	2.0	2.71789	0.00749	0.07134	0.00098	0.00253	0.00004	0.00255	0.00068	0.00882	0.00006	1.18508	0.28900	20.90624	5.06988
4	2.5	2.25194	0.00774	0.10096	0.00110	0.00246	0.00008	0.00501	0.00068	0.00709	0.00005	1.34010	0.15706	23.62308	2.75311
5	3.0	1.37325	0.00720	0.24516	0.00105	0.00361	0.00007	0.00838	0.00068	0.00339	0.00004	1.47472	0.05491	25.97915	0.96896
6	3.7	1.18737	0.00727	0.25232	0.00117	0.00336	0.00007	0.00838	0.00068	0.00246	0.00005	1.79663	0.07164	31.60064	1.25895
7	4.5	0.90857	0.00715	0.35482	0.00118	0.00411	0.00008	0.01197	0.00068	0.00097	0.00004	1.74645	0.03806	30.72557	0.68125
8	5.5	1.02138	0.00694	0.39584	0.00125	0.00472	0.00012	0.01018	0.00068	0.00113	0.00005	1.72942	0.03932	30.42838	0.70235
9	6.5	1.42872	0.00805	0.60905	0.00186	0.00712	0.00014	0.01803	0.00068	0.00105	0.00004	1.83304	0.02341	32.23540	0.43826
10	7.7	1.82092	0.00700	0.87933	0.00209	0.01030	0.00015	0.02432	0.00068	0.00090	0.00004	1.76374	0.01565	31.02703	0.31335
11	9.0	3.59243	0.01041	1.62078	0.00331	0.01907	0.00010	0.04856	0.00068	0.00256	0.00005	1.74528	0.01251	30.70515	0.26609
12	10.5	2.31103	0.00754	1.09677	0.00194	0.01299	0.00012	0.03174	0.00068	0.00122	0.00005	1.77436	0.01465	31.21232	0.29862
13	12.5	3.88175	0.00807	1.68664	0.00269	0.01936	0.00024	0.07554	0.00068	0.00310	0.00007	1.75258	0.01417	30.83248	0.29055
14	14.5	3.09208	0.00845	1.49707	0.00343	0.01744	0.00022	0.05468	0.00068	0.00147	0.00005	1.77299	0.01152	31.18842	0.25355

	MSWD	% ³⁹Ar	Initial Ratio
Plateau age	31.05	0.37	1.7
Isochron age	30.99	0.49	3
			94.6
			na
			292.3±4.9

Step no	Power (%)	⁴⁰ Ar	±σ ₄₀	³⁹ Ar	±σ ₃₉	³⁸ Ar	±σ ₃₈	³⁷ Ar	±σ ₃₇	³⁶ Ar	±σ ₃₆	⁴⁰ Ar/ ³⁹ Ar	±σ	Age (Ma)	±σ (Ma)
SLP17-23 (Panalillo) sanidine, grain size 0.1-0.3 mm, 0.0325mg, J=0.009850291 ± 7.987E-05															
1	0.2	0.00043	0.00082	-0.00008	0.00009	0.00003	0.00002	0.00007	0.00012	0.00001	0.00002	22.24100	95.01483	357.38182	1385.04533
2	0.5	0.12195	0.00096	0.00603	0.00013	0.00013	0.00003	0.00040	0.00012	0.00035	0.00002	3.02345	1.24353	52.94425	21.46345
3	0.9	0.40041	0.00183	0.05776	0.00074	0.00092	0.00005	0.00160	0.00012	0.00109	0.00004	1.31317	0.22334	23.18635	3.92272
4	1.2	0.31489	0.00207	0.07448	0.00065	0.00102	0.00004	0.00106	0.00012	0.00069	0.00003	1.47309	0.13754	25.98969	2.41833
5	1.5	0.83210	0.00676	0.32756	0.00301	0.00398	0.00010	0.00808	0.00012	0.00097	0.00002	1.66041	0.03425	29.26794	0.64341
6	2.0	0.30119	0.00266	0.11910	0.00055	0.00158	0.00003	0.00292	0.00012	0.00029	0.00003	1.81017	0.08718	31.88453	1.54349
7	2.7	0.97335	0.00700	0.44556	0.00388	0.00520	0.00007	0.01128	0.00012	0.00070	0.00002	1.71247	0.02733	30.17796	0.53569
8	3.4	1.51208	0.01509	0.77432	0.00889	0.00887	0.00020	0.01546	0.00012	0.00061	0.00003	1.71627	0.03057	30.24434	0.58697
9	4.4	1.64429	0.02375	0.80701	0.01190	0.00947	0.00025	0.01470	0.00012	0.00094	0.00003	1.68843	0.04049	29.75780	0.74706
10	5.6	0.95267	0.00380	0.37110	0.00097	0.00438	0.00007	0.00854	0.00012	0.00120	0.00003	1.60553	0.02910	28.30819	0.55771
11	7.0	0.30115	0.00215	0.15658	0.00095	0.00183	0.00003	0.00326	0.00012	0.00014	0.00003	1.66279	0.06602	29.30959	1.17810
12	8.5	2.46656	0.02733	1.12168	0.01341	0.01312	0.00030	0.02132	0.00012	0.00207	0.00003	1.64751	0.03257	29.04241	0.61560
13	9.0	0.98904	0.00374	0.41378	0.00176	0.00515	0.00009	0.00910	0.00012	0.00104	0.00004	1.64341	0.03282	28.97065	0.61952

			MSWD	% ³⁹Ar	Initial Ratio
Plateau age	29.4	0.5	1.8	100	na
Isochron age	30.4	0.74	0.99	na	272±15

Step no	Power (%)	⁴⁰ Ar	±σ ₄₀	³⁹ Ar	±σ ₃₉	³⁸ Ar	±σ ₃₈	³⁷ Ar	±σ ₃₇	³⁶ Ar	±σ ₃₆	⁴⁰ Ar*/ ³⁹ Ar	±σ	Age (Ma)	±σ (Ma)
SLP17-13 (Cabras) whole rock, grain size 0.3-0.5 mm, 0.0559mg, J=0.0096716± 4.8358E-05															
1	0.3	0.07913	0.00180	0.00075	0.00071	0.00004	0.00003	0.00019	0.00144	0.00027	0.00002	-2.60297	-10.38043	-45.99895	185.79856
2	0.6	1.28527	0.00769	0.08301	0.00102	0.00192	0.00008	0.12907	0.00144	0.00405	0.00006	0.92590	0.24093	16.08350	4.16731
3	0.9	1.18678	0.00381	0.30154	0.00153	0.00454	0.00010	0.44766	0.00145	0.00279	0.00006	1.17033	0.06272	20.30554	1.08684
4	1.2	1.68241	0.01130	0.83760	0.00555	0.00968	0.00011	1.21463	0.00145	0.00196	0.00006	1.31005	0.02724	22.71454	0.48272
5	1.5	1.37243	0.01110	0.96489	0.00733	0.01137	0.00022	1.47190	0.00145	0.00048	0.00003	1.27335	0.01825	22.08204	0.33313
6	2.1	1.67691	0.00596	1.23491	0.00334	0.01501	0.00020	2.62228	0.00145	0.00039	0.00004	1.26441	0.01191	21.92801	0.23236
7	2.7	1.69995	0.00506	1.23201	0.00443	0.01527	0.00019	3.23687	0.00145	0.00040	0.00004	1.28194	0.01207	22.23018	0.23558
8	3.3	0.94035	0.00357	0.66871	0.00272	0.00874	0.00011	2.19228	0.00145	0.00030	0.00003	1.27196	0.01670	22.05819	0.30803
9	4.4	1.75785	0.00671	1.25177	0.00216	0.01603	0.00012	4.59718	0.00145	0.00045	0.00003	1.29615	0.00996	22.47498	0.20479
10	5.7	1.32664	0.00913	1.01058	0.00442	0.01294	0.00019	3.59353	0.00084	0.00034	0.00004	1.21363	0.01632	21.05253	0.30034
11	7.7	1.47146	0.00911	1.11524	0.00786	0.01442	0.00011	4.13878	0.00084	0.00044	0.00003	1.20135	0.01475	20.84070	0.27468

	MSWD	% ³⁹Ar	Initial Ratio
Plateau age	22.21	0.29	1.6
Isochron age	21.93	0.59	5.9
			75.6
			na
			293±15

Tabla A.8.- Información de química mineral. Abreviaciones B= análisis en borde; C= análisis en el centro.

Composición representativa de feldespatos alcalinos de las rocas félsicas del CVSSM

Muestra	SLP1711	SLP1711	SLP1711	SLP1711	SLP1711	SLP1711	SLP1728	SLP1728	SLP1728	SLP1728	SLP1728	SLP1728
Análisis	B	B	C	C	B	B	B	B	C	C	B	B
Unidad	SSM	SSM	SSM	SSM	SSM	SSM	Zapote	Zapote	Zapote	Zapote	Zapote	Zapote
Roca (TAS)	Riolita	Riolita	Riolita	Riolita	Riolita	Riolita	Riolita	Riolita	Riolita	Riolita	Riolita	Riolita
SiO ₂	65.304	65.088	66.015	64.88	65.354	66.273	66.146	66.913	66.136	66.184	67.31	66.297
TiO ₂	0.04	-0.028	0.003	0.029	0.001	0.058	-0.013	-0.033	0.012	0.067	0.008	-0.045
Al ₂ O ₃	18.817	19.041	19.128	18.92	18.963	19.004	18.468	19.033	18.606	18.846	18.672	18.679
Fe ₂ O ₃ ^t	0.018	0.021	0.011	0.029	0.021	0.036	0.038	0.003	0.06	0.04	-0.002	0.063
MnO	-0.036	0.016	-0.015	0.003	0.002	0.026	0.008	0.002	-0.01	-0.012	0.008	0.003
MgO	0.024	0.016	0.019	-0.019	-0.037	-0.004	-0.031	-0.008	0.016	-0.007	-0.002	-0.018
CaO	0.241	0.205	0.197	0.208	0.231	0.245	0.293	0.293	0.365	0.339	0.302	0.298
Na ₂ O	3.658	3.549	3.509	3.551	3.515	4.139	4.426	4.239	4.178	4.178	4.637	4.211
K ₂ O	11.391	11.455	11.466	11.415	11.575	10.903	10.546	10.589	10.49	10.743	10.151	10.806
P ₂ O ₅	0.007	-0.017	0.017	-0.014	-0.003	-0.01	-0.017	-0.007	0.003	-0.01	-0.044	-0.02
Total	99.464	99.346	100.35	99.002	99.622	100.67	99.864	101.024	99.856	100.368	101.04	100.274
%Ab	32.41	31.69	31.44	31.77	31.22	36.15	38.40	37.29	37.03	36.54	40.38	36.66
%An	1.18	1.01	0.98	1.03	1.13	1.18	1.40	1.42	1.79	1.64	1.45	1.43
%Or	66.41	67.30	67.59	67.20	67.65	62.66	60.20	61.29	61.18	61.82	58.16	61.90

Composición representativa de clinopiroxenos de las rocas intermedias del CVSSM

Muestra	SLP1720	SLP1720	SLP1720	SLP1720	SLP1720	SLP1720	SLP1710	SLP1710	SLP1710
Análisis	B	B	C	C	B	B	B	C	B
Unidad	La Placa	La Placa	La Placa	La Placa	La Placa	La Placa	La Placa	La Placa	La Placa
Roca (TAS)	Andesita	Andesita	Andesita	Andesita	Andesita	Andesita	Andesita	Andesita	Andesita
SiO ₂	52.553	49.773	49.312	49.374	50.791	51.336	51.139	51.236	51.272
TiO ₂	0.491	0.955	1.111	1.027	0.791	0.913	0.437	0.297	0.451
Al ₂ O ₃	2.285	3.464	3.998	3.823	3.549	1.37	2.824	2.265	2.484
Fe ₂ O ₃ ^t	11.023	11.81	12.031	11.665	12.378	11.613	6.384	6.701	6.233
MnO	0.238	0.271	0.29	0.3	0.331	0.28	0.178	0.186	0.165
MgO	14.276	13.533	13.472	13.477	13.461	15.936	17.541	18.217	18.048
CaO	17.926	19.142	18.933	18.809	18.972	17.32	19.6	18.721	19.683
Na ₂ O	0.246	0.388	0.371	0.327	0.363	0.166	0.317	0.281	0.206
K ₂ O	0.449	-0.009	0.001	-0.011	0.006	0.037	-0.013	-0.008	0.004
P ₂ O ₅	0.023	0.016	-0.01	0.00	0.00	-0.02	0.017	0.017	0.056
Total	99.51	99.343	99.509	98.791	100.642	98.951	98.424	97.913	98.602
%Wo	38.5	40.4	40.1	40.2	39.9	35.5	39.92	37.89	39.55
%En	42.7	39.8	39.7	40.0	39.4	45.5	49.71	51.30	50.47
%Fs	18.8	19.8	20.2	19.8	20.7	18.9	10.37	10.81	9.98

Composición representativa de clinopiroxenos de las rocas básicas del CVSSM

Muestra	SLP1713	SLP1713	SLP1713	SLP1713	SLP1713	SLP1713
Análisis	B	B	B	B	C	C
Unidad	Cabras	Cabras	Cabras	Cabras	Cabras	Cabras
Roca (TAS)	Basalto	Basalto	Basalto	Basalto	Basalto	Basalto
SiO ₂	50.974	51.593	51.596	51.757	51.997	51.322
TiO ₂	0.981	1.064	1.181	1.001	0.896	0.936
Al ₂ O ₃	1.579	1.598	1.336	1.729	1.509	1.575
Fe ₂ O ₃ ^t	10.782	10.95	11.732	9.769	10.515	11.448
MnO	0.245	0.269	0.292	0.267	0.3	0.302
MgO	16.264	15.957	15.807	15.95	16.338	16.241
CaO	18.416	17.851	17.572	18.897	17.799	17.522
Na ₂ O	0.209	0.188	0.234	0.252	0.215	0.251
K ₂ O	-0.009	-0.009	-0.012	-0.017	0.007	-0.012
P ₂ O ₅	0.02	0.013	-0.033	-0.016	0.02	0.026
Total	99.461	99.474	99.705	99.589	99.596	99.611
%Wo	37.13	36.61	35.94	38.66	36.38	35.59
%En	45.63	45.54	44.98	45.41	46.47	45.90
%Fs	17.25	17.85	19.08	15.93	17.15	18.52

Composición representativa de ortopiroxenos de las rocas intermedias del CVSSM

Muestra	SLP1720	SLP1720	SLP1720	SLP1720	SLP1720	SLP1720	SLP1710	SLP1710	SLP1710	SLP1710
Análisis	B	C	C	C	B	B	B	C	C	B
Unidad	La Placa	La Placa	La Placa	La Placa	La Placa	La Placa	La Placa	La Placa	La Placa	La Placa
Roca (TAS)	Andesita	Andesita	Andesita	Andesita	Andesita	Andesita	Andesita	Andesita	Andesita	Andesita
SiO ₂	53.346	53.21	53.527	54.086	52.994	52.868	52.93	54.039	53.918	54.449
TiO ₂	0.488	0.425	0.325	0.331	0.368	0.397	0.404	0.486	0.197	0.232
Al ₂ O ₃	3.625	2.664	1.678	2.38	2.994	2.254	2.761	2.917	2.713	2.588
Fe ₂ O ₃ ^t	11.641	11.513	11.605	10.137	10.858	14.717	20.453	19.366	9.831	10.569
MnO	0.21	0.178	0.21	0.197	0.198	0.283	0.023	-0.012	0.212	0.19
MgO	28.397	29.712	29.743	30.998	29.802	27.648	5.059	5.573	30.657	30.669
CaO	2.874	1.698	1.896	1.323	1.797	1.493	1.218	1.486	1.672	1.727
Na ₂ O	0.061	0.032	0.072	0.074	0.009	0.018	0.143	0.158	0.024	0.022
K ₂ O	-0.009	-0.007	0.006	-0.001	-0.009	0.01	1.418	1.195	0.01	-0.002
P ₂ O ₅	0.024	0.028	-0.003	0.014	0.017	0.024	0.072	0.078	0.028	-0.003
	100.657	99.453	99.059	99.539	99.028	99.712	84.481	85.286	99.262	100.441
%Wo	5.57	3.26	3.61	2.52	3.47	2.89	5.05	6.13	3.21	3.27
%En	76.61	79.34	78.91	82.20	79.99	74.56	29.17	31.98	81.84	80.90
%Fs	17.82	17.40	17.48	15.28	16.54	22.55	65.79	61.89	14.95	15.82

Composición representativa de clinopiroxenos de las rocas básicas del CVSSM

Muestra	SLP1713	SLP1713	SLP1713	SLP1713	SLP1713	SLP1713	SLP1713
Análisis	B	B	C	C	C	B	B
Unidad	Cabras	Cabras	Cabras	Cabras	Cabras	Cabras	Cabras
Roca (TAS)	Basalto	Basalto	Basalto	Basalto	Basalto	Basalto	Basalto
SiO ₂	49.94	52.61	51.17	52.789	51.884	52.926	51.447
TiO ₂	0.544	0.478	0.468	0.426	0.415	0.532	0.462
Al ₂ O ₃	0.845	0.707	0.702	0.682	0.491	0.556	0.566
Fe ₂ O ₃ ^t	20.099	20.267	17.654	17.578	19.732	21.004	22.247
MnO	0.471	0.443	0.449	0.406	0.511	0.472	0.525
MgO	17.87	21.263	22.047	21.782	22.162	21.847	20.644
CaO	7.451	4.201	5.674	5.911	3.613	2.248	3.124
Na ₂ O	0.187	0.044	0.125	0.086	0.049	0.06	0.007
K ₂ O	-0.009	-0.013	-0.006	-0.017	-0.019	-0.017	-0.006
P ₂ O ₅	0.033	0.013	-0.047	-0.02	0.051	-0.017	0.02
Total	97.431	100.013	98.236	99.623	98.889	99.611	99.036
%Wo	15.43	8.43	11.26	11.78	7.20	4.56	6.31
%En	51.51	59.35	60.87	60.41	61.48	61.65	58.01
%Fs	33.06	32.23	27.87	27.81	31.31	33.79	35.68

Tabla A9.- Geotermómetros en equilibrio. Ecuaciones tomadas de Putirka (2008)

<p>Feldespato-liquido en equilibrio</p>	$\frac{10^4}{T(K)} = 17.3 - 1.03 \ln \left[\frac{X_{Ab}^{afs}}{X_{Na0.5}^{liq} X_{Al0.5}^{liq} (X_{SiO_2}^{liq})^3} \right] - 200(X_{Ab}^{afs})$ $- 2.42(X_{Na0.5}^{liq}) - 28.8(X_{Al0.5}^{liq}) + 13500(X_{CaO}^{liq} - 0.0037)^2$ $- 550(X_{K0.05}^{liq} - 0.056)(X_{Na0.5}^{liq} - 0.089) - 0.078P(kbar)$
<p>Ortopiroxeno-líquido en equilibrio</p>	$\frac{10^4}{T(K)} = 4.07 - 0.329[P(GPa)] + 0.12(H_2O^{liq})$ $+ 0.567 \ln \left[\frac{X_{Fm2Si2O6}^{opx}}{(X_{SiO_2}^{liq})^2 (X_{FeO}^{liq} + X_{MnO}^{liq} + X_{MgO}^{liq})^2} \right]$ $- 3.06[X_{MgO}^{liq}] - 6.17[X_{K0.05}^{liq}] + 1.89[Mg\#^{liq}] + 2.57[X_{Fe}^{opx}]$
<p>Clinopiroxeno- líquido en equilibrio</p>	$T(^{\circ}C) = \frac{93100 + 544P(kbar)}{61.1 + 36.6(X_{Ti}^{cpx}) + 10.9(X_{Fe}^{cpx}) - 0.95(X_{Al}^{cpx} + X_{Cr}^{cpx} - X_{Na}^{cpx} - X_K^{cpx}) + 0.395[\ln(a_{En}^{cpx})]^2}$
<p>Líquidos en equilibrio con ol+plg+cpx</p>	$T(^{\circ}C) = -583 + 3141[X_{SiO_2}^{liq}] + 15779[X_{Al_2O_3}^{liq}] + 1338.6[X_{MgO}^{liq}]$ $- 31440[X_{SiO_2}^{liq} \cdot X_{Al_2O_3}^{liq}] + 77.67[P(GPa)]$