

Consolidated Chemical Provinces on Mars

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Interpretations A. Rani^{1,2}, A. Basu Sarbadhikari^{1,*}, D. R. Hood³, O. Gasnault⁴, S. Nambiar¹ and S. Karunatillake⁵ ¹Physical Research Laboratory, Planetary Sciences Division, Ahmedabad, Gujarat 380009, India ²Indian Institute of Technology, Gandhinagar, Earth Sciences, Gujarat 382355, India ³Baylor University, Department of Geosciences, Waco, Texas 76798, USA ⁴Institut de Recherche en Astrophysique et Planétologie, Université de Toulouse, CNRS, CNES, 31028 Toulouse, France ⁵Louisiana State University, Geology and Geophysics, Baton Rouge, Louisiana 70808, USA *Correspondence to: A. Basu Sarbadhikari, amitbs@prl.res.in **Key Points:** We have consolidated three key multivariate methods, yielding the first unified set of chemical provinces of Mars Igneous geochemistry pervades volatile-rich regions, consistent with water-limited isochemical weathering throughout Mars's geologic history Secular chemical trends suggest complex processes of mantle melting for highlands, which differs from a possibly garnet-rich lowland source

Consolidated Chemical Provinces on Mars: Implications for Geologic

Abstract

Chemical provinces were defined on Mars a decade ago using orbital nuclear spectroscopy of K, Th, Fe, Si, Ca, Cl, and H₂O. However, past multivariate analyses yielded three sets of provinces, suggesting methodologic variability. Province-stability to the inclusion of Al and S is also unknown, presenting additional uncertainties for geologic insight. Here we consolidate key multivariate methods to define the first cross-validated provinces. In southern highlands, the highly incompatible K and Th show non-uniform distribution with higher values in mid Noachian and Hesperian than late Noachian – early Hesperian volcanic terrains. Silica- and Al-depletion trends from Noachian to Amazonian indicate highly differentiated mantle with variable degree of melting. Late Hesperian lowlands are highly depleted in Al and enriched in K and Th, consistent with volcanic resurfacing from a low-degree partially melted, garnet-rich mantle. Furthermore, older volatile-rich regions such as Medusae Fossae Formation exhibit igneous geochemistry, consistent with water-limited isochemical weathering throughout Mars's history.

Plain Language Summary

Topographically Mars can be divided into two large regions: southern highlands and northern lowlands. However, the compositional evolution of these two landforms and their source characteristics remains unclear. Therefore, using enhanced satellite nuclear spectroscopic data, we develop a set of consolidated geochemical provinces of Mars using three multivariate analysis techniques. We identified the correlation of distinct geochemical provinces with mapped geologic units, which demonstrates the effect of bedrock composition over bulk soil composition. Our study depicts discrete igneous provinces within the southern highlands, which follow a secular chemical pattern, indicating a highly differentiated mantle

source with an evolving formation pressure and degree of melting. The northern lowlands, on the other hand, are uncorrelated with the chemical trend of southern highland provinces, suggesting a distinct evolution. Our findings further reveal that regardless of volatile enrichment, igneous crustal composition dominates over chemical alteration signatures indicating a limited role of liquid water on the martian landscape.

1. Introduction

A *trilogy* of multivariate cluster analysis methods applied to the chemical maps derived from Mars Odyssey gamma-ray and neutron spectroscopy (GRS) initially identified chemical provinces and supported interpretations of regional-scale geology (Karunatillake et al., 2009; Gasnault et al., 2010; Taylor et al., 2010). However, the lack of an intermethodological synthesis to characterize geological processes has hindered subsequent works and stymied advances with other datasets (cf., Rogers & Hamilton 2015). Furthermore, key chemical maps used in subsequent papers (Baratoux et al., 2014; Ojha et al., 2018; Plesa et al., 2018) were unavailable or only in a preliminary state for the original trilogy: Aluminium (Al), sensitive to both igneous evolution and secondary mineralogy (cf., Nimmo & Tanaka, 2005; Ehlmann et al., 2011; McLennan, 2012); and Sulfur (S), sensitive to cycling processes within the martian critical zone across the atmosphere and crust (Halevy et al., 2007; King & McLennan, 2010).

Here we use nine elemental mass fraction maps from the GRS: key rock-forming elements (Fe, Si, Ca, and Al) with large ion lithophiles (K and Th) and volatiles (stoichiometrically equivalent H₂O, Cl, and S) to resolve the aforementioned challenges. We synthesize and advance prior multivariate methods (Karunatillake et al., 2009; Gasnault et al., 2010; Taylor et al., 2010) to derive the first consolidated chemical provinces of Mars. We

consider three of the provinces as case studies on geologic insight, revealing diverse mantle sources and magmatic processes in volcanic areas, characterizing possible global ashfall units, and evaluating the relative roles of igneous versus aqueous environments.

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2. Data and Methods

We use chemical maps derived from GRS spectra to delineate the chemically distinct provinces. The average decimetre sampling depth and coarse spatial resolution of GRS overcome the effects of fine dust mantles making GRS elemental analysis ideal for primary and secondary geological process studies on broad scales. GRS measures the energy spectrum of gamma photons emitted from the martian surface and characteristic spectral peaks allow us to measure the abundance and distribution of most major rock-forming elements, along with some minor elements present on the martian surface. There are two main processes on the martian surface that give rise to gamma-photon emission. Gamma photons can be emitted from the decay of natural radioactive elements (K and Th) or emissions induced by cosmic particle interactions with the martian atmosphere and nonradioactive elements of the martian surface. However, due to increasing H (reported as stoichiometric H₂O mass fractions) towards the poles causing mass dilution for radioactive elements and compositional modelling issues for others due to neutron moderation, derived chemical maps are restricted to the mid-to-low latitudes, roughly within ±50°. Furthermore, chemical maps are generated at a resolution of $5^{\circ} \times 5^{\circ}$ due to broad footprint of the GRS instrument (Boynton et al., 2007). We use the latest refined datasets of GRS, including Al and S used in the literature (cf., Hood et al., 2016, 2019; Ojha et al., 2018, 2021), and compare them with GRS data used a decade ago (Boynton et al., 2007; Karunatillake et al., 2009; Gasnault et al., 2010; Taylor et al., 2010), including the archived data at the NASA Planetary Database System (https://pds-geosciences.wustl.edu/ody/ody-m-grs-5-elements
<a href="https://pds-geosciences.wustl.

2.1. Multivariate Cluster Analysis

We apply three multivariate cluster analyses to discern a set of internally uniform chemical provinces that reflect the compositional variability of the martian crust. The three methods (Supporting Information S1.2 provides more details) consist of non-hierarchical clustering with principal component analysis (NHC-PCA: Taylor et al., 2010), hierarchical clustering with principal component analysis (HC-PCA: Gasnault et al., 2010), and modified student's t-test to distinguish spatially overlapping Gaussian tail clusters (t-GTC: Karunatillake et al., 2009).

The three methods optimize clustering differently but still present spatially overlapping trends suggestive of distinct provinces. NHC-PCA optimizes Euclidian distances between cluster centers, where the number of clusters is specified a priori. In comparison, HC-PCA builds a hierarchy of clusters using a top-down clustering approach based on a divisive algorithm (Kaufman & Rousseeuw, 2005), where the complete martian landscape mapped by GRS belongs to one cluster and is split iteratively, moving down the hierarchy to highlight the differences between the sub-clusters. The modified t-GTC method identifies spatially contiguous areas where two or more elements deviate from their respective average mid-to-low latitude crustal abundances by one standard deviation or greater. The t-GTCs and

HC-PCA-derived provinces reduce spatial autocorrelation biases by using unclassified (boundary) pixels to separate provinces from each other. Complementarily, the NHC-PCA method delineates border-sharing provinces.

We combine the intermediate provinces from the NHC-PCA, HC-PCA, and t-GTC methods into a consolidated map using a spatial contouring approach. A contour map of provinces is made by categorizing the regions into three levels: provinces derived from all three-methods overlapping, provinces with any two-methods overlapping, and provinces with no overlap (Figure S1). We demarcate the boundaries of final provinces resulting from at least two-methods overlapping, reducing the spatial uncertainty in any method (Supporting Information S2 provides more details). Provinces that deviate minimally from the average martian crust (AMC) composition (Supporting Information S2.1 and S2.2) are omitted so that the consolidated chemical province map highlights crustally distinct regions. In addition, we use mapped geology (Tanaka et al., 2014) for context on chemical trends.

3. Results

All three methods lead to large chemically uniform provinces, with minimal intraprovince compositional variance, reinforcing the subdued variability in chemical maps of
Mars. The intermediate chemical province sets resulting from the three underlying methods
(Supporting Information S2 provides more details) are labeled numerically (Province-1 to
Province-6; Figures 1a-c and Tables S1 and S2). Province-2 from NHC-PCA and HC-PCA
methods lacks the minimum criterion of at least two methods overlapping (Figure S1;
Supporting Information S2 provides more details), and Province-4 is compositionally
identical to the AMC within statistical uncertainty. Therefore, Province-2 and 4 are deprioritized in the consolidated chemical province map (Figure 1d). Moreover, provinces that

overlap spatially across the methods are compositionally similar even if driven by different multivariate methods (**Tables S2**, **S3**). This further justifies our contour-consolidation method yielding the merged chemical province map (**Figure 1** and **Tables 1** and **S3**). Overall, overlapping areas labeled as 'Province-1' in **Figures 1a-c** have been combined into 'Province-A', 'Province-3' into 'Province-B', 'Province-5' into 'Province-C', and 'Province-6' into 'Province-D', resulting in four consolidated provinces (**Figure 1d** and **Tables 1** and **S1**). Compositional consistency across the numeric intermediate provinces enables this synthesis (**Supporting Information S2**). While geographically widely separated areas may constitute a single province, such regions show some compositional heterogeneity.

3.1. Distinct Geochemical Provinces

Four geographically distributed regions in the southern highlands constitute Province-A (Figure 1d). This province broadly represents mid and late Noachian highlands (> 30%) to early Hesperian aged volcanic units (> 20%) (Figures 2a and S2), containing many of the large Hesperian volcanic provinces. This province is enriched in Al and depleted in K, Th, and volatiles (H₂O, S, Cl) relative to the AMC (Figure 2b and Tables 1, S3, and S4; Supporting Information S2.2). The K/Th and S/Cl ratios are above average, while the Fe/Si ratio is consistent with the AMC.

Province-B comprises Amazonian and Amazonian-Hesperian aged volcanic units (> 30%) with Amazonian to Hesperian transitional geological units (~ 18%) (Figures 2a and S2). The distribution of K and Th is consistent with the AMC (Taylor et al., 2006); however, the K/Th ratio is slightly low compared to the AMC (Figure 2b and Tables 1, S3 and S4; Supporting Information S2.2). This province is moderately depleted in Si, Ca, and Fe and highly depleted in Al, while the volatiles (especially Cl and S) are highly enriched relative to

the AMC. The S/Cl ratio is depleted from preferential Cl enrichment. Compared to the AMC, the Fe/Si ratio is below average.

Province-C consists of two geographic regions: Sinus Meridiani (SM) and the Medusae Fossae Formation (MFF), dominated by mid-Noachian highlands (~ 40 %) (Figures 1d, 2a, and S2 and Table S1). These regions are enriched in volatiles (H₂O, Cl, and S) (cf. Diez et al. 2008; Feldman et al., 2011; Maurice et al., 2011), while Fe, Ca are slightly enriched relative to the AMC (Figure 2b and Tables 1, S3 and S4; Supporting Information S2.2). The ratio of S/Cl and K/Th is consistent with the AMC. However, Fe enrichments make the Fe/Si ratio higher relative to the AMC.

Two regions of broadly late Hesperian lowlands form Province-D (> 60%) (Figures 2a and S2). It is confined to Southern Acidalia Planitia and Utopia Planitia of the Northern lowland near the dichotomy boundary (Figure 1d and Table S1). Since both these regions are found within the northern lowlands and the northern extent of each is obscured by the chemical mapping boundary (Supporting Information S1), they may constitute a larger, contiguous region. The broader extent of Province-D, which includes the western and eastern part of this province, spatially overlaps with the TES-derived Surface Type 2 mineralogy in the martian northern lowlands (Bandfield et al., 2000; Rogers et al., 2007; Rogers & Hamilton 2015). This province is highly depleted in Al, enriched in Si, Fe, K, and Th relative to the AMC (Figure 2b and Tables 1, S3 and S4), contrasting with other provinces. The K/Th and S/Cl ratio is consistent with the AMC, whereas Fe/Si is higher.

4. Discussion

4.1. Extent of aqueous alteration and K/Th trends

We first evaluate the extent of aqueous alteration evident at regional scales to The Chemical Index of decimeter depths. Alteration (CIA=Al₂O₃/(Al₂O₃+CaO+Na₂O+K₂O)×100) in standard molar abundances of oxides corresponding to the provinces A-D is < 50, which indicates no signature of weathering (Figure 2c) at these broad scales. The K/Th ratio is similar across all the provinces, and with the AMC within statistical uncertainty (Figure 2b). Specifically, K/Th ratio is effectively invariant with the abundance of the volatiles (H₂O, SO₃, and Cl) even across the maximum in Province-C to a minimum in Province-A (Figure 3a). The most incompatible mapped elements, K and Th (partition coefficient <<1; Borg & Draper, 2003) do not fractionate in the solid phases during magmatic processes. Collectively, our study shows that even the volatilerich regions are compositionally consistent with water-limited isochemical weathering throughout Mars's geologic history (Hurowitz & McLennan, 2007; Tosca et al., 2008). Consequently, the provinces are broadly representative of well-preserved igneous processes at least to decimeter depth scales and averaged over a few hundred kilometres horizontally.

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4.2. Secular chemical trends on Mars

Earlier work by Taylor et al. (2010) grouped all volcanic landscapes into a single province irrespective of age. However, our study supports the findings (Karunatillake et al., 2009; Gasnault et al., 2010) that volcanic regions form separate provinces by age (Figure 2a), as shown by Baratoux et al. (2011). Moreover, the minimally weathered secular volcanic composition of provinces A-C in the southern highlands of Mars shows distinct geochemical trends. Unlike southern highlands, Province-D, the late Hesperian lowland unit, deviates from the secular chemical trend (Figure 3b-d), indicating distinct lithology. The provinces A-C from mid-Noachian to Amazonian show Al-, Fe-, Si- and Ca-depletion (Figure 3b-d). As

hypothesized by Balta & McSween (2013), higher SiO₂ in older regions of Mars may indicate dehydration melting of the mantle with hundreds of mg/kg water, similar to that of Province-A and -C of our study, in contrast to the relatively silica-depleted younger Province-B. Unlike Earth, considering a stagnant-lid regime of Mars (Golombek & Phillips, 2009; Grott et al., 2013), the decrease of SiO₂ and concomitant increase in olivine component in the parent magma from the oldest to youngest volcanic terrains of southern highlands could be explained by the growth of lithospheric thickness, which would increase the formation pressure, supporting the results of Baratoux et al. (2013). This contrasts with the topmost micron-layer observation by Thermal Emission Spectrometer (TES), suggestive of weathering reflected by the non-detection of olivine in the Hesperian highland terrains, at the spatial resolution of 1 pixel per degree (Rogers & Hamilton, 2015). GRS-scale footprints in this study consistently reveal the igneous crustal composition albeit weathering processes was prevalent during martian surface evolution. In addition, high Ca/Al for Province-B and -D could indicate a path for sampling a high-pressure, garnet-rich source (Treiman, 2005; Dasgupta & Hirschmann, 2007; Balta & McSween, 2013). The abundances of K and Th are relatively low in late Noachian - early Hesperian Province-A as compared to Hesperian-Amazonian Province-B and mid-Noachian Province-C in the highland volcanic terrains (Figure 3e). Earlier studies advocated the early Noachian crust building phenomenon for the enrichment of these large ion lithophile elements (LILEs: K and Th) in Noachian than Hesperian terrains (Rogers & Hamilton, 2015). However, our study shows that the youngest volcanic province also contains a high relative abundance of those LILEs, which contradicts the hypothesis of LILE fractionation during the early Noachian crust formation. Moreover, the non-systematic trend of the abundances of LILEs and other secular chemical trends may necessitate complex melting processes in a highly differentiated mantle source on a regional

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to global scale (cf. Basu Sarbadhikari et al., 2016; 2017; Udry et al., 2020) that existed from the early-stage evolution of Mars.

Compared to prior works, the emergence of distinct regions as well as the exclusion of previously suggested provinces (**Figure 1d**) reflects our methodological advances along with the use of Al and S. Given the collective distinctness among the consolidated chemical provinces, we consider three regions for detailed case studies: Lunae Planum, Medusae Fossae Formation, and Southern Acidalia Planitia (**Figure 1d**).

4.3. Lunae Planum (LP) in Province-A

LP region is a geologically distinct plateau of early Hesperian-aged highland material (Tanaka et al., 2014), which lies between Chryse Planitia and Olympus Mons in Province-A (Figure 1d). Geochemically, the LP is depleted in K, Th, and H₂O relative to the AMC. LP follows the general K and Th crustal correlation on Mars and substantially overlaps with the K and Th distribution in Hesperia and Thaumasia Planum (Figures 3e and S3). Consequently, LP's K and Th depletion likely reflects volcanic resurfacing, not aqueous weathering (Taylor et al., 2006). H₂O depletion in the LP is also comparable in magnitude to that in the vicinity of Greater Thaumasia (Hood et al., 2016). Given minimal chemical weathering across the provinces (Section 4.1), the H₂O depletion is likely volcanic and could suggest a dry mantle source as predicted for Hesperian volcanism (cf., Balta & McSween, 2013). However, that mantle evolution model also predicts high Th abundances, not seen in our study. Overall, the LP likely follows the same volcanic evolution trends as identified in the Greater Thaumasia area, i.e., low H₂O abundance and temporally decreasing K and Th, maybe a part of the same evolving volcanic province (Hood et al., 2016). Furthermore, higher

K and Th abundance within LP relative to Greater Thaumasia could place it as the oldest member of an underlying volcanic series.

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4.4. Medusae Fossae Formation (MFF) in Province-C

The discrete wind-sculpted outcrops of the mid-Noachian MFF extend along the martian equator (Figure 1d) and their origin remains uncertain (c.f., Kerber & Head, 2010; Zimbelman & Scheidt, 2012; Wilson et al., 2018). Geochemically, the MFF region is highly enriched in Fe, Ca, and volatiles (SO₃, Cl, and H₂O) relative to the AMC (**Figure S4**) (Keller et al., 2006). However, the abundance of K, Th, Al, Si, and the value of K/Th is consistent with the martian crust. As highly incompatible elements in igneous processes, the ratio of Cl/K would inherit the initial mantle abundance (Taylor et al., 2010). Although the ratio is globally maintained, the MFF region shows significantly different Cl/K than the rest of Mars. Chlorine is highly abundant, and K is generally lower than that in AMC (Figures 3e and S4), consistent with Cl enrichment in airborne dust and volcanic aerosol in the MFF (Newsom & Hagerty, 1997; Goetz et al., 2005). Meanwhile, high S and Cl content, moderately high H₂O (Figure S5), fine-grained nature, preserved primary mineralogy, and extensive erosional features suggest that MFF is the source of the global dust reservoirs of Mars (Ojha et al., 2018). Thus, our work supports the provenance of martian dust from mostly mechanical weathering and transport of a pyroclastic ashfall unit, which possibly arose from SW-oriented paleo-winds during Tharsis volcanic eruptions (Keller et al., 2006; Diez et al., 2009). Our results don't support the hypothesis of water-ice existence in polar layer-like deposits to explain the volatile enrichment in the equatorial MFF (Wilson et al., 2018). Summarily, the voluminous pyroclasts of MFF are most likely the product of low-moderate degree partial

melting of a distinct volatile-rich mantle source during mid-Noachian age in comparison to the late Noachian - Amazonian volcanic provinces of the martian highlands.

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4.5. Southern Acidalia Planitia (SAP) in Province-D

SAP is a late Hesperian plain area (Tanaka et al., 2014) in the northern lowlands that geographically lies between Tharsis and Arabia Terra and is part of Province-D (Figure 1d). Geochemically, SAP is enriched in K, Th, Fe, Si and depleted in Al relative to the rest of the AMC (Figure S6). Despite K and Th enrichment, the K/Th ratio of SAP is consistent with the martian crust (Figures 3e and S6), which suggests primary rock composition (Karunatillake et al., 2006; Taylor et al., 2006), diverging from the extensive alteration posited by Bouley et al. (2020). Al/Si versus Mg/Si plots of SAP (Figure S7a) show a slope different from the martian crust but resembling that of younger martian meteorites (after, McSween et al., 2009). The Al depletion in SAP best correlates with that of the mean basaltic shergottite composition (Nimmo & Tanaka, 2005), and given the K, Th, and K/Th ratio signatures, it is more likely to be related to mantle composition than chemical weathering (Taylor et al., 2006) at least at the scales of our observations. Further, mineralogical observations by Compact Reconnaissance Imaging Spectrometer for Mars (CRISM) reveal mafic mineral signatures in outcrops, crater ejecta, and central peaks of the craters in SAP (Pan et al., 2017) resembling Hesperian-aged volcanism (Salvatore et al., 2010). Also, the non-detection of Al-bearing sulfates (Figure S7b) that would result from low-pH aqueous alteration (McLennan, 2012) supports an igneous origin for the apparent depletion of Al. The collective K and Th enrichment and Al-depletion can best be explained by a low degree of partial melting at high pressure derived from a garnet-rich mantle source for SAP.

5. Summary and Conclusion

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The first consolidated chemical provinces of Mars reveal several geochemical signatures of broad geologic import. Our findings demonstrate secular chemical trends among the southern highland provinces A-C may indicate complex processes of an evolving degree of partial melting within a highly differentiated mantle source. Distinct lithology of the late Hesperian lowland unit, Province-D, does not follow the secular chemical trends like highland provinces. This could provide a mechanism for sampling a high-pressure, garnetrich source for Province-D. We also find that even older volatile-rich regions preserve igneous geochemistry, indicating how throughout Mars's geologic history, water-limited isochemical weathering dominated regionally to decimeter depths scales. Specifically, as a distinct plateau of early Hesperian-aged highlands, the Lunae Planum region shows compositional links – and possibly shared provenance - with Province-A, much like Hesperia Planum, Syrtis Major, and part of Greater Thaumasia. The similarity of Province-C's K/Th trend with the average crust and enrichment of volatiles (especially S and Cl) supports mostly regional volcanic materials (e.g., ashfall pyroclastic deposits), especially for the Medusae Fossae Formation, despite volatiles' enrichment. In contrast to the provinces A-C, Southern Acidalia Planitia of Province-D in the northern lowlands is lithologically distinct with the highest abundance of K and Th of the provinces. However, the similarity of K/Th and depletion of Al relative to the average crust argues against sedimentary signatures of putative paleo-ocean(s). The northern lowlands may mostly represent the mafic crust of Mars, which might have originated by a low degree of partial melting at high pressure as suggested by high abundances of incompatible elements (K, Th) and Al- depletion rather than the conventional thought that the northern lowlands are highly fractionated. These findings also demonstrate the overall utility of the integrative multivariate methodology we have developed for martian geochemical data, motivating possible applications to other planetary bodies (e.g., Mercury, Moon, and Vesta) with comparable datasets.

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- data (doi: 10.17632/3jd9whd78m.1). Archival GRS-derived chemical maps are from NASA
- 353 Planetary Database System (PDS, https://pds-
- 354 geosciences.wustl.edu/missions/odyssey/grs cgs.html). Mars geology map taken from
- USGS-310 archive (https://pubs.usgs.gov/sim/3292/), and topographic data from MOLA-
- 356 HRSC blended DEM data by Fergason et al. (2018).

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Figure Caption

- Figure 1: Results obtained from multivariate analysis by NHC-PCA (a), HC-PCA (b), and t-
- 361 GTC (c) methods, overlaid on the Mars Orbiter Laser Altimeter and High-Resolution Stereo
- 362 Camera (MOLA-HRSC) data map (Smith et al., 2001) to delineate the geochemical
- provinces. Provinces-2 and -4 do not appear in the t-GTC method (c) as they are
- compositionally similar to the average crustal composition (Tables S2 and S3). (d) The
- consolidated geochemical provinces on Mars are derived by integrating provinces from three
- different multivariate techniques and overlaid onto mapped geology (Tanaka et al., 2014;

different colors indicate different geologic units with major units in legend). The white dashed outline areas delineate our regional (LP, SAP, and MFF) case studies.

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Figure 2: (a) The relative areal fractions of secular, transitional, and volcanic units within each chemical province corresponding to the consolidated geochemical map. The prefix "N" indicates Noachian, "H" indicates Hesperian, and "A" indicates Amazonian, and suffix "v" indicates the associated volcanic units (after, Tanaka et al., 2014). A detailed version of Figure 2a is given in Figure S2, both as per Tanaka et al. (2014) geology map. (b) Modified box-and-whiskers diagram to compare the distribution of elements for all provinces to the global geochemistry of the AMC. The top of the upper box represents 75th percentile/25th percentile ratio, the bottom of the lower box represents the 25th/75th percentile ratio, and the ratio of the medians by the boundary between them. The whiskers here represent the uncertainty on the ratio of medians using the Median Absolute Deviation (MAD) (Karunatillake al., 2011). (c) The Chemical Index of Alteration (CIA=100*[Al₂O₃/(Al₂O₃+CaO+Na₂O+K₂O)]) diagram is used to assess the extent of chemical weathering in the provinces A-D. For CIA, we are using standard molar abundances of oxides. Non-GRS Na₂O is calculated using Baratoux et al. (2014).

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Figure 3: Diagrams of K/Th normalized to the bulk silicate Mars (~ 5300) versus GRS derived total volatile concentration (a), Fe versus Si (b), Al versus Fe (c), and Ca versus Si (d) of the provinces A-D are represented by colored boxes (southern highland provinces A-C) and ellipse (northern lowland Province-D). The modest variation in K/Th is consistent with the bulk Mars, implies to low or no alteration at decimeter depths in the consolidated provinces. The regional uncertainty is shown at the right corner of each plot, represented by

the average of standard error of mean (1 σ) from all provinces (**Table 1**). From Noachian to Amazonian, all the three plots (**b-d**) exhibit a compositional trend, marked by a dashed arrow, except Province-D (**Figure S2**). The prefix "N" indicates Noachian, "H" indicates Hesperian, and "A" indicates Amazonian, and the suffix "v" indicates associated volcanic units, "h" indicates highland, "m" indicates middle and "lHl" indicates late Hesperian lowland. (**e**) The scatter values of K (wt. %) versus Th (mg/kg) in Southern Acidalia Planitia (SAP), Lunae Planum (LP), Medusae Fossae Formation (MFF), consolidated provinces A-D, and AMC. SAP, LP, and MFF generally follow the same trend of K/Th as the AMC.

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Figure :	1.
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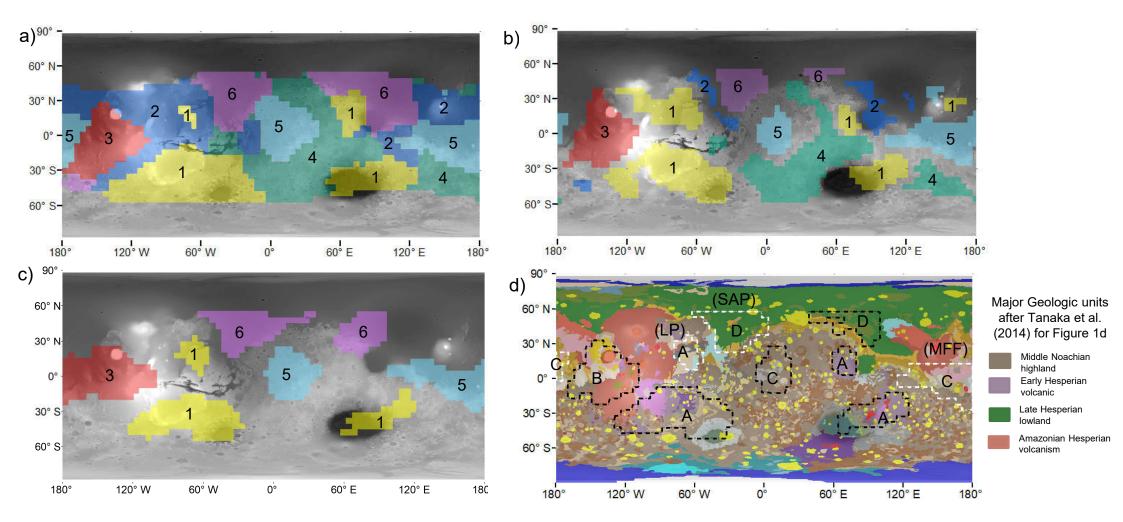


Figure 2	2.
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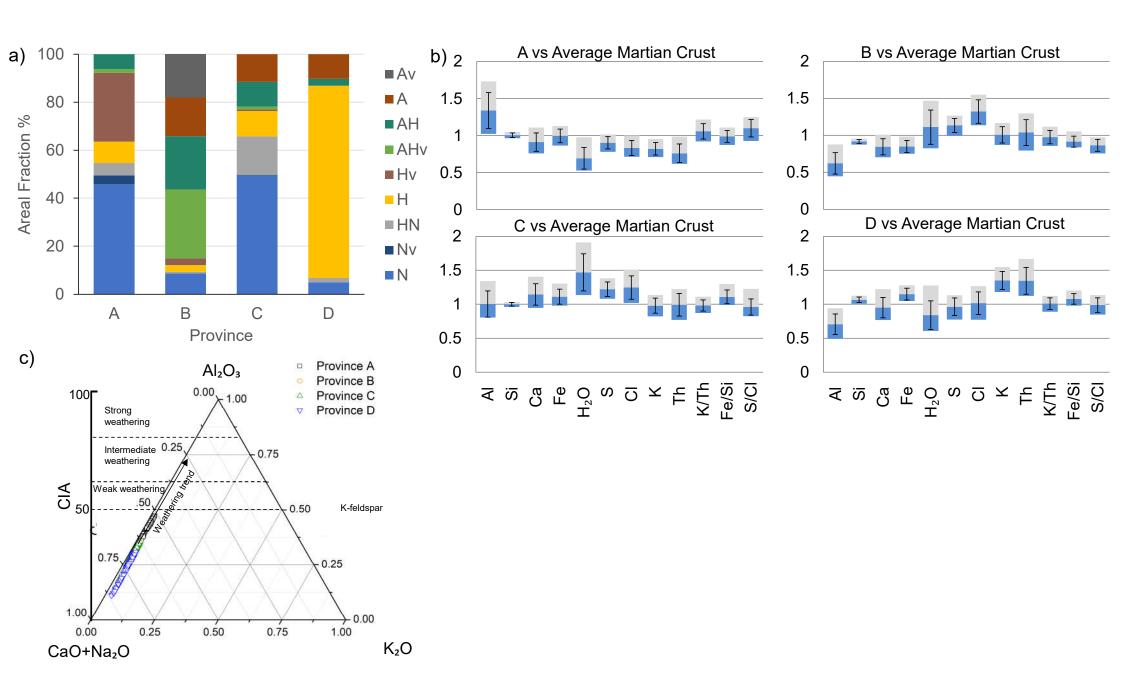


Figure	3.
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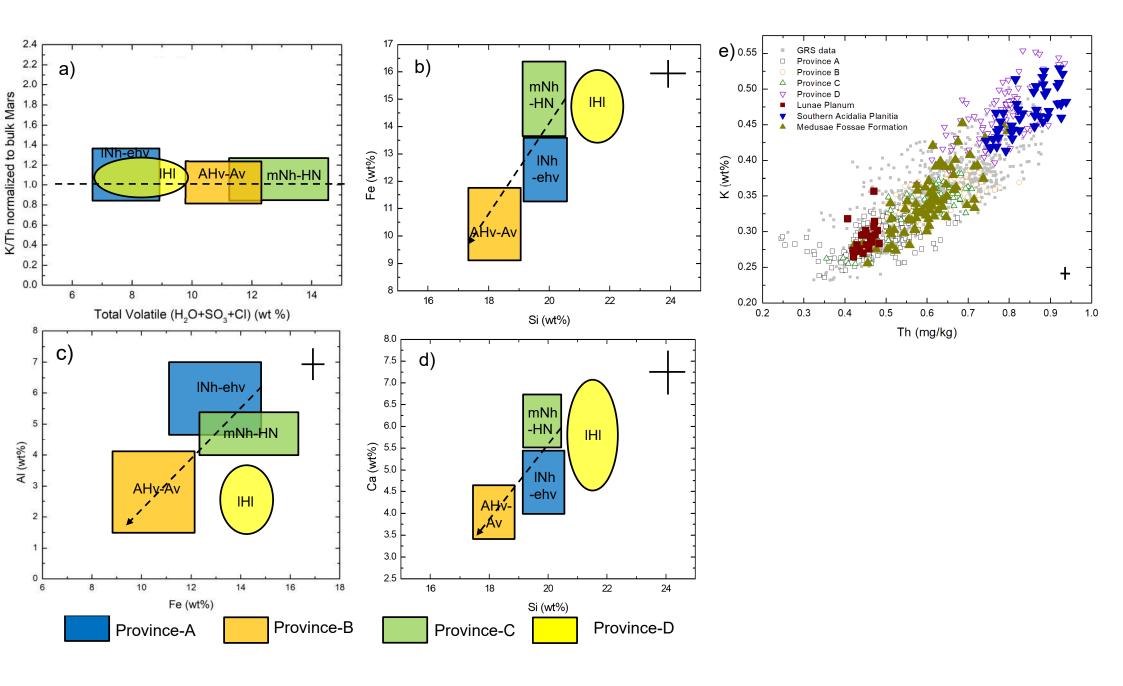


Table 1: Arithmetic mean stoichiometric oxide mass fraction as a percentage (wt%, if not mentioned otherwise), elemental ratio, and its corresponding root means square uncertainty (σ) are calculated from mean measurement uncertainties for the grid points composing a geochemical province. Whereas, K/Th ratio uncertainty is calculated from $[(K/Th)*((\sigma_K/K)^2+((\sigma_{Th}/Th)^2)^{1/2})]^{1/2}$ (GRS derived) of the consolidated chemical provinces.

	Province-A	Province-B	Province-C	Province-D	Mid-latitudinal Average Martian Crust (AMC)
K ₂ O	0.35±0.01	0.42±0.01	0.40±0.01	0.56±0.02	0.43±0.01
FeO	16.1±0.9	14.0±0.8	18.2±1.1	18.4±1.2	16.3±1.0
SiO ₂	42.7±2.3	39.1±2.1	42.4±2.6	44.9±3.0	42.5±2.5
CaO	6.6±1.2	5.9±1.1	7.9±1.4	6.9±1.7	7.0±1.3
Th mg/kg	0.48 ± 0.04	0.62±0.05	0.59±0.05	0.80 ± 0.06	0.61±0.05
Al ₂ O ₃	10.6±2.1	5.5±1.8	8.6±2.2	6.2±2.6	8.5±2.2
H ₂ O	2.9±0.4	4.2±0.4	5.5±0.5	3.5±0.6	3.9±0.5
Cl	0.41±0.03	0.61±0.03	0.57±0.04	0.46 ± 0.04	0.47±0.03
SO ₃	4.9±0.5	6.0±0.5	6.5±0.6	5.2±0.7	5.4±0.6
K/Th	6060±580	5600±430	5680±520	5800±440	5760±470