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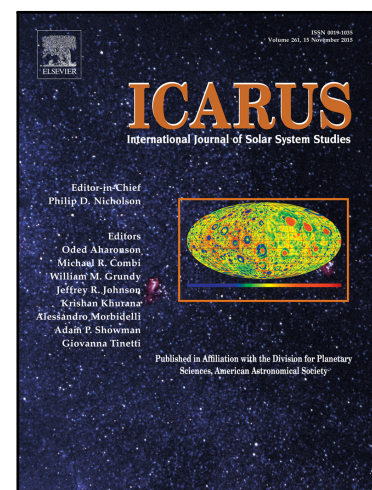
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Lithologic variation within bright material on Vesta revealed by linear spectral unmixing

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Abstract

Vesta's surface is mostly composed of pyroxene-rich lithologies compatible with howardite, eucrite and diogenite (HED) meteorites (e.g., McCord et al., 1970; Feierberg and Drake, 1980). Data provided by the Visible and Infrared (VIR) spectrometer, onboard the NASA Dawn spacecraft, revealed that all Vesta reflectance spectra show absorption bands at $\sim 0.9 \mu\text{m}$ and $\sim 1.9 \mu\text{m}$,

28 which are typical of iron-bearing pyroxenes (De Sanctis et al., 2012a). Other
 29 minerals may be present in spectrally significant concentrations; these in-
 30 clude olivine and opaque phases like those found in carbonaceous chondrites.
 31 These additional components modify the dominant pyroxene absorptions.
 32 We apply linear spectral unmixing on bright material (BM) units of Vesta to
 33 identify HEDs and non-HED phases. We explore the limits of applicability
 34 of linear spectral unmixing, testing it on laboratory mixtures. We find that
 35 the linear method is applicable at the VIR pixel resolution and it is useful
 36 when the surface is composed of pyroxene-rich lithologies containing moder-
 37 ate quantities of carbonaceous chondrite, olivine, and plagioclase. We found
 38 three main groups of BM units: eucrite-rich, diogenite-rich, and olivine-rich.
 39 For the non-HED spectral endmember, we choose either olivine or a fea-
 40 tureless component. Our work confirms that Vesta's surface contains a high
 41 content of pyroxenes mixed with a lower concentration of other phases. In
 42 many cases, the non-HED endmember that gives the best fit is the featureless
 43 phase, which causes a reduction in the strength of both bands. The anticor-
 44 relation between albedo and featureless endmember indicates that this phase
 45 is associated with low-albedo, CC-like opaque material. Large amounts of
 46 olivine have been detected in Bellicia, Arruntia and BU14 BM units. Other
 47 sites present low olivine content ($< 30\%$) mostly with a high concentration
 48 of diogenite.

49 HIGHLIGHTS

- 50 • Characterization of Vesta's bright material (BM) units by linear un-
 51 mixing analysis;

- Detection of different lithologies within BM regions;
- Detection of phases other than pyroxenes from VIR data on Vesta surface;
- Calculation of mixing coefficients for the detected lithologies for each in BM unit;

1. Introduction

1.1. Early knowledge of Vesta

The first spectral study of Vesta dates back to Bobrovnikoff (1929). McCord et al. (1970) compared Vesta spectra with the meteorite Nuevo Laredo, inferring a relation between Vesta and pyroxenes. Pyroxenes are characterized by the presence of two crystal field absorptions, centered approximately at 0.9 and 1.9 μm (McCord et al., 1970), due to ferrous iron (Fe^{+2}) in octahedral sites (e.g., Burns, 1993). Consolmagno and Drake (1977) suggested a possible link between Vesta and eucrite meteorites, and Feierberg and Drake (1980) proposed that Vesta is a mixture of howardite and eucrite. Studies by Lupishko et al. (1988) show a clear inverse correlation between the polarization and brightness of Vesta, correlated with the west-east dichotomy. Hubble Space Telescope (HST) observations between 1994 and 1996 confirmed this inference by highlighting the geological differences on the asteroid (Binzel et al., 1997). Moreover the identification of the prominent impact basin at the south pole (now named Rheasilvia) corroborates the hypothesis that Vesta is the parent body of the howardite, eucrite and diogenite (HED) meteorites (Thomas et al., 1997). In addition, the disk-integrated mid-infrared

spectra of Vesta have shown the presence of minor constituents as olivine, feldspar, and chromite (Donaldson Hanna and Sprague, 2009). Studies by Li et al. (2010) revealed that the vestan surface can be divided into several geological units (more specifically, regions of lithologic variation), in particular, eucrite-rich and diogenite-rich units, as well as slightly weathered and freshly exposed units. Shestopalov et al. (2010) concluded that the global diversity of Vesta might be caused by variations of non-basaltic source sediments enriched by dark rocks, spinel group minerals, or even chondritic-like material. Moreover, they showed that certain vestan units contain olivine abundances of several volume percent.

1.2. *The Dawn-Era and Vesta's lithologies*

Dawn entered orbit around Vesta in July 2011 (Russell and Raymond, 2011), and enabled many discoveries during about one year of observations. For the first time, Dawn acquired data of Vesta's surface at high spatial resolution, allowing for production of complete geological and lithological maps (De Sanctis et al., 2012a; Williams et al., 2014). Dawn's Visible and Infrared (VIR) imaging spectrometer (De Sanctis et al., 2011) provided high spectral resolution data covering the majority of Vesta's surface at varying spatial scales. The spatial resolution of the VIR maps ranges from ~ 800 m/pixel in the Survey phase to ~ 180 m/pixel in the HAMO and HAMO-2 phases (see Table 1 of Zambon et al. (2015)).

Vesta is the parent body of the HED meteorites. HEDs encompass a large variety of igneous rocks, similar to basalts, cumulate gabbros, orthopyroxenites and igneous brecciated mixtures (Mittlefehldt et al., 1998). The diogenites are coarse-grained cumulates that originated in a plutonic layer deep in the

100 crust (Mittlefehldt et al., 1998; Beck and McSween, 2010; McSween et al.,
 101 2011, 2013). The mineralogy of diogenites is dominated by orthopyroxenes
 102 (from 87% to 99%); all diogenites contain < 5% chromite and some contain
 103 olivine, typically at contents < 10% (McSween et al., 2011). Eucrites occur
 104 as basaltic or cumulate rocks. They are dominated by Ca-poor pyroxenes
 105 and plagioclase, with minor amounts of metal, troilite, chromite, ilmenite,
 106 and silica (Mayne et al., 2010; McSween et al., 2011). Eucrites are believed
 107 to have crystallized as lavas on the surface or within relatively shallow dikes
 108 and plutons (McSween et al., 2011). Basaltic eucrites contain Fe-rich pyrox-
 109 enes, whereas cumulate eucrites are predominantly unbrecciated and their
 110 chemistry is similar to basaltic eucrites, but richer in Mg (Mittlefehldt et al.,
 111 1998; McSween et al., 2011). Howardites are brecciated achondrites, prin-
 112 cipally mixtures of eucrite and diogenite clasts, and reflectance spectra of
 113 howardites have pyroxene band center positions intermediate between those
 114 of eucrites and diogenites (Mittlefehldt et al., 1998; McSween et al., 2011).
 115 Impact mixing of eucrite and diogenite has produced the polymict breccias
 116 and howardites.
 117 Eucrite, diogenite and howardite lithologies are present on Vesta's surface,
 118 as revealed by VIR data (De Sanctis et al., 2012a; Ammannito et al., 2013a)
 119 augmented by Dawn Framing Camera images (Reddy et al., 2012b) and
 120 Gamma Ray And Neutron Detector data (Lawrence et al., 2013; Prettyman
 121 et al., 2011, 2013, 2014). Recently, many papers on the analysis of Vesta's
 122 surface composition using Dawn data have been published (e.g., Ruesch et al.
 123 (2014); Ammannito et al. (2013a); De Sanctis et al. (2013b); Le Corre et al.
 124 (2013); Reddy et al. (2013); Thangjam et al. (2013)). The crust of Vesta is

dominated by howardite enriched in eucrite (De Sanctis et al., 2012a, 2013b). A few outcrops of diogenite are present in localized areas in the south polar region, corresponding to the rim of the large impact basin Rheasilvia. Large areas of diogenite-enriched howardite have also been identified in the northern hemisphere (longitude 0° - 90° E) and are interpreted to consist of Rheasilvia ejecta (De Sanctis et al., 2012a; Ammannito et al., 2013a). Ammannito et al. (2013a) suggested a high content of olivine (50-80 vol.%) in the area of Bellicia (lat 40° N, lon 40° E) and Arruntia craters (lat 40° N, lon 70° E), highlighting the presence of lithologies different from that of typical HEDs. Others eleven sites containing much smaller amounts of olivine (located up to 39° S latitude) have also been identified by Ruesch et al. (2014), and six other new olivine-rich regions were proposed by Palomba et al. (2015), almost all located at latitudes below 28° N. Dark material units on Vesta were discussed by Jaumann et al. (2012); McCord et al. (2012); Reddy et al. (2012a), and Palomba et al. (2014). Hydrated mineral phases were shown to be correlated with this low-reflectance material (De Sanctis et al., 2012b), confirming that the dark material can be attributed to the presence of a carbonaceous chondrite (CC)-like component delivered by impacts.

1.3. Bright material units: A slice of fresh material

The spectral analysis of the bright material units reveals that they are generally characterized by greater band depths than their surroundings. Most bright units have a howardite rich-eucrite, composition, with some exceptions such as the bright unit called "BU15" in Zambon et al. (2014).

BM units could represent pristine material that has been recently exposed on the surface, and thus contains less carbonaceous-chondrite-like contami-

150 nation introduced by impact mixing than other areas of Vesta (see Zambon
 151 et al. (2014) for more detail). To be consistent with Zambon et al. (2014), we
 152 follow the same classification of the BM units, as described by Mittlefehldt
 153 et al. (2012). The abbreviation CWM stands for crater wall material, SM
 154 means slope material, and RM refers to radial material. The spectral anal-
 155 ysis employed in Zambon et al. (2014) to derive Vesta's lithologies is based
 156 on selected spectral parameters, namely band center (BC), band depth (BD)
 157 and Band Area Ratio (BAR).

158 The goal of our work is to use linear spectral unmixing to automatically
 159 identify the main lithologies of specific regions and determine their semi-
 160 quantitative mixing coefficients, including identification and mapping of min-
 161 eralogical phases other than the spectrally dominant pyroxenes. The mixing
 162 coefficients depend on various effects, such as, the grain size, the abundance
 163 of the mineralogical phases present in the scene, and the scale at which the
 164 components are mixed (i.e., macroscopic vs. intimate; Combe et al. (2008)).
 165 Since the grain size on Vesta, at the VIR spatial scale, is quite homogeneous
 166 ($< 25 - 45 \mu\text{m}$) (Hiroi et al., 1994; Palomba et al., 2014; Zambon et al.,
 167 2014), we can assume that the grain size does not substantially affect our
 168 derived mixing coefficients. Thus the unmixing results principally depend on
 169 the abundance of the mineralogical phases and on the type of mixing.

170 Unmixing methods are useful for understanding the composition of a surface
 171 (e.g. Pieters and Englert, 1993; Keshava and Mustard, 2002; Bioucas-Dias
 172 et al., 2012). Mixing models can be either linear or non-linear. In the linear
 173 case, the spectrum of a region can be considered the area-weighted average of
 174 the endmembers present (Singer and McCord, 1979; Hapke, 1993; Bioucas-

175 Dias et al., 2012). The endmembers can be extracted from the scene, or
 176 can be reference spectra of plausible analogue phases. The endmembers are
 177 generally assumed to represent the different components, which are the fun-
 178 damental constituents of the scene, so that each pixel in the scene can be
 179 modeled as a linear combination of the endmembers (Bioucas-Dias et al.,
 180 2012).

181 The case of non-linear mixing arises when light is scattered by multiple mate-
 182 rials in the scene. Scattered light interacts with more than one endmember,
 183 and the resulting spectrum is a non-linear combination of the components.
 184 Models for non-linear mixing require comprehensive information on the ex-
 185 pected minerals, such as scattering coefficients, particle sizes, and optical
 186 constants (e.g., Hapke (1981); Shkuratov et al. (1999)). Here we use linear
 187 unmixing as a first step in order to derive additional information on the com-
 188 position of Vesta's surface.

189 Many authors have linear unmixing algorithms for analysis of remote sensing
 190 data for a variety of bodies (e.g., Adams et al. (1986); Fox III et al. (1990);
 191 Ramsey and Christensen (1992); Blewett et al. (1995); Combe et al. (2008);
 192 Dalton (2007); McCord et al. (2012)). Adams et al. (1986) already applied
 193 linear unmixing methods to Viking lander data of Mars, while Dalton (2007)
 194 applied linear unmixing to Europa data using a set of laboratory spectra as
 195 endmembers, and Combe et al. (2008) used the Multiple-Endmember Lin-
 196 ear Spectral Unmixing Model (MELSUM) to analyze Mars data from the
 197 Infrared Mineralogical Mapping Spectrometer (OMEGA) onboard MarsEx-
 198 press. A first use of linear unmixing methods on Vesta was presented by
 199 McCord et al. (2012), who found that the entire surface of Vesta as observed

by VIR could be modeled using the weighted sum spectral endmembers, 'bright' and 'dark' (McCord et al., 2012). In this paper, we apply the same linear unmixing method employed by Tosi et al. (2015) and Zambon et al. (2015) to extended areas of Vesta.

Here we considered high spatial resolution data from the Dawn mission at Vesta, and we selected a set of plausible laboratory spectra analogues as endmembers. We apply the linear unmixing method to the bright material (BM) units of Vesta, previously analyzed by Zambon et al. (2014). BM units are often associated with impact craters. Impacts expose fresh, unweathered material from beneath the surface, which could be indicative of the original vestan crust composition (Zambon et al., 2014). The previous spectral analysis revealed the main lithologies (eucrite, diogenite and howardite) and the spectral characteristics of these units. Here we study the same BM units with the goal of quantifying the mixing coefficients of the different lithologies. We also aim to identify the presence of other mineralogical phases in addition to the spectrally dominant, HED components. In principle, this work can be extended also to larger region on Vesta, determining the global distribution of Vesta's non-HED lithologies.

2. Dataset description and VIR spectra characteristics

The Dawn mission at Vesta consisted of four phases, based on the altitude of the spacecraft: Survey (2735 km altitude), HAMO (High Altitude Mapping Orbit, 695 km), LAMO (Low Altitude Mapping Orbit, 210 km) and HAMO-2, an extension of the mission similar to the HAMO phase (Russell and Raymond, 2011). Here we used HAMO and HAMO-2 VIR data,

224 which cover large part of Vesta's surface with a high spatial resolution (~ 180
 225 m/pixel). The VIR instrument has two distinct detectors, or "channels": the
 226 visible, covering wavelengths from $0.25\ \mu\text{m}$ to $1.07\ \mu\text{m}$, and the infrared, with
 227 sensitivity from $1.02\ \mu\text{m}$ to $5.10\ \mu\text{m}$ (De Sanctis et al., 2011). Each channel
 228 has 432 spectral bands, so the average spectral sampling is $1.8\ \text{nm}/\text{band}$ for
 229 the visible channel and $9.8\ \text{nm}/\text{band}$ for the infrared channel (De Sanctis
 230 et al., 2011). Each VIR image cube has 256 spatial samples with a variable
 231 number of lines determined by the length of a scan, and 432 spectral bands.
 232 For details about the VIR data calibration, refer to a document archived
 233 at the NASA Planetary Data System Small Bodies Node: http://sbn.psi.edu/archive/dawn/vir/DWNVVIR_I1B/DOCUMENT/VIR_CALIBRATION/VIR_CALIBRATION_V2_4.PDF.
 234 The VIR visible and infrared data are each separately geo-
 235 referenced, then the two VIS and IR subsets are merged together to form
 236 a single spectrum covering the entire spectral range from 0.25 to $5.10\ \mu\text{m}$.
 237 The last 19 spectral channels of the visible range, which usually are affected
 238 by higher instrumental noise, are discarded.
 239
 240 In this work described here, we consider VIR data calibrated in units of re-
 241 flectance factor (I/F) from $0.6\ \mu\text{m}$ to $2.5\ \mu\text{m}$, focusing our analysis on the
 242 two pyroxene bands. We considered the average spectra of BM units used for
 243 spectral characterization by Zambon et al. (2014). The wavelength position
 244 of the centers of the two bands gives information on the HED lithologies.
 245 The displacement of the pyroxene band centers is sensitive to temperature
 246 variations (e.g., Burbine et al. (2009), Reddy et al. (2012c)). However, given
 247 the usual range of daytime temperatures recorded on Vesta for BM units (be-
 248 tween 252 and 265 K, Tosi et al. (2014)), this effect is essentially negligible

249 in VIR data (Longobardo et al., 2014).

250 The strength (depth) of the pyroxene absorption bands is a function of the
 251 abundances of the minerals, the particle size, and the presence of opaque
 252 phases (e.g., Clark (1999)). Band depths are also affected by temperature
 253 variations and the phase angle of the observations (Longobardo et al., 2014;
 254 Reddy et al., 2012c). The relatively small range of temperatures found on
 255 Vesta does not affect the band depth shape: Dalton et al. (2011) demon-
 256 strated that a temperature range of 150K is necessary to observe substantial
 257 band-depth effects (whereas on Vesta we found variations never larger than
 258 50 K). Band depths are the only spectral parameter among those considered
 259 that vary depending on illumination and viewing geometry. However, the
 260 variation of band depth with phase angle is very small for bright regions,
 261 although variation can become important as albedo decreases (Longobardo
 262 et al., 2014). In this work, observations at phase angles between 28° and
 263 52° and incidence angles lower than 60° have been selected. According to
 264 Longobardo et al. (2014), in this narrow range band depths are only weakly
 265 dependent on phase and hence a photometric correction is not mandatory.
 266 Phase, incidence and emission angles of the BM units are reported in Tables
 267 7, 8 and 9 of the Zambon et al. (2014) supplementary online material (SOM).
 268 In this work, we utilize the "Claudia" coordinate system (supplemental ma-
 269 terial of Russell et al., 2012; Li et al., 2012; Roatsch et al., 2012; Reddy et al.,
 270 2013), which has been used by Zambon et al. (2014) for tabulation of the
 271 BM unit locations.

3. Analytical method

3.1. Linear unmixing

Understanding the composition of a surface using reflectance spectroscopy may be challenging. Analysis of spectral parameters is useful but does not offer a complete characterization of the surface materials. Key additional information can be provided by the application of mixing model algorithms. Based on past analysis (De Sanctis et al., 2012a; Jaumann et al., 2012; McCord et al., 2012; Ammannito et al., 2013a; De Sanctis et al., 2013a), we assume that reflectance spectra of Vesta can be reproduced by a linear combination of at least three endmembers, chosen from a sample of eucrites, diogenites, olivines and a featureless endmember. To describe the primary components of Vesta's surface, we consider only diogenite and eucrite endmembers, and do not include howardite as a separate endmember. Howardites, which are the major component of Vesta terrains (De Sanctis et al., 2012a), are brecciated rocks with variable amounts of both eucrite and diogenite; thus howardites can be spectrally characterized by a mixing of the two components. The mixing equation is:

$$Y_{mix} = a_1 y_1 + a_2 y_2 + a_3 y_3 \quad (1)$$

where y_1 , y_2 , y_3 are the scaled spectra of the endmembers and a_1 , a_2 , a_3 are their respective mixing coefficients. The unmixing algorithm requires all mixing coefficients to be non-negative, and their sum must be equal to unity.

The method we developed is able to automatically find the endmembers and the mixing coefficients, subject to the positivity constraint, which allows exclusion of solutions that are not physically meaningful. We outline the principal steps of our algorithm as follows:

1. Spectral slope removal. Normalization of the spectrum by the continuum slope is done for two reasons. First, the position of the band center is defined as the location of the reflectance minimum inside the band as measured on a continuum-removed spectrum. Second, the process of continuum normalization removes the influence of albedo from the unmixing process. Albedo is closely tied to the photometric conditions of the observation rather than mineralogy of the surface. To remove the slope, we consider the line between the first point ($0.6 \mu\text{m}$) and the last point ($2.5 \mu\text{m}$) of the spectrum, then we divide the spectrum by this line. To be consistent, the slope was removed in the same way from both VIR spectra and the spectral endmembers (See Fig. 1).

FIGURE 1

2. We selected a sample of 14 laboratory spectra from a larger spectral library. In Section 3.2, we describe in detail the endmembers and the criteria for their selection.
3. We calculate the χ^2 values for all the combinations. We select the best fit spectrum corresponding with the minimum χ^2 value, with a confidence limit of 95%. In Table 2, we report all the χ^2 values of the bright units, associated to the best-fit, with the corresponding values of the correlation coefficient (r).

Following these steps, the algorithm is able to determine automatically the best combination of endmembers that describe the scene, along with their mixing coefficients. We accept only best fits with $\chi^2 < 1$.

321 3.2. Endmembers selection

322 Careful choice of endmembers is necessary to obtain consistent results.
 323 In our case, we consider as endmembers a number of laboratory spectra,
 324 accounting for pure minerals and for HED meteorites that are assumed to be
 325 representative of materials that may be present on Vesta's surface.

326 We select the endmembers from the Reflectance Experiment Laboratory
 327 (RELAB) database (<http://www.planetary.brown.edu/relabdocs>). The
 328 endmembers selection includes consideration of the particle size. The supple-
 329 mentary online material from Reddy et al. (2012b) discusses the grain size
 330 differences between eucrite and diogenite. Studies based on the analysis of
 331 HED laboratory samples show that the grain size of diogenites in howardite
 332 breccias ranges from 500 μm to 1.5 millimeters whereas the average grain size
 333 of eucritic material is $\leq 70 \mu\text{m}$ (Beck et al., 2011; Reddy et al., 2012b SOM).
 334 Previous work based on ground-based observation and remote sensing data
 335 suggest that for the spatial resolution available, most of Vesta's surface can
 336 be best represented by regolith with particle sizes $< 25\mu\text{m}$ (Hiroi et al., 1994;
 337 Palomba et al., 2014; Zambon et al., 2014). This limits our choice to labo-
 338 ratory spectra acquired for samples in the $< 25\mu\text{m}$ size fraction, if available.
 339 Since we are working with continuum-removed spectra, absolute reflectance
 340 does not enter in our analysis. Therefore, we cannot distinguish between a
 341 low-reflectance featureless phase and a high-reflectance featureless phase: we
 342 can just observe a decrease in band depth. To account for this effect, we
 343 introduce a featureless endmember represented by a straight horizontal line.
 344 Nonetheless, examination of the mean albedo values derived in Zambon et al.
 345 (2014) for all the bright units can be useful in identifying the presence of low-

346 vs. high-reflectance featureless endmembers.

347 We selected 14 endmember spectra including six eucrite, five diogenite
 348 and two olivine samples from the RELAB database, plus the straight line
 349 representing a featureless endmember. This limit is imposed in order to
 350 allow computations to be completed in reasonable time. Since eucrite-rich
 351 howardite is the dominant component on Vesta's surface (De Sanctis et al.,
 352 2013b) and because eucrites show a wide range of petrographic and spectral
 353 characteristics, we include a larger number of them in our endmembers set.
 354 The endmembers selection was done by considering the spectral parameters,
 355 the spectral shape and their petrographic differences. We selected spectra
 356 with band centers and band depths that cover the possible range of values
 357 (Fig. 2), and those with different spectral characteristics, to take into ac-
 358 count all of the possible spectral variations (see Fig. 3). Moreover for eucrite
 359 we considered both cumulate and basaltic samples, as well as polymict and
 360 monomict material. We also tested the distribution frequency of each spec-
 361 trum from a preliminary combination of two endmembers chosen among the
 362 whole available spectral library, performing a linear unmixing test. With just
 363 two endmembers, the entire library of laboratory spectra can be used in the
 364 analysis because of the reduced computational time. We calculate the spec-
 365 tral parameters with the same methods described in Section 4 of Zambon et
 366 al. (2014).

367 In Fig. 4, we show the frequency of the endmembers found in the case of
 368 a linear combination of two endmembers. Eucrite appears to be the most
 369 abundant component (68% of the total components), followed by diogenite
 370 (25%) and olivine (7%). Taking the above discussion into consideration, we

select for our analysis: 1) the eucrites Bereba (6e), Bouvante (7e), Cachari (8e), Padvarninkai (20e), Serra de Mage (22e), Y74450 (24e); 2) the diogenites ALH77256 (2d), GRO95555 (5d), (6d) Johnstown, (8d) Tatahouine and Y75032 (10d); and 3) two olivine samples, Jackson County (1o) and Green olivine (2o) (see Table 1). In Fig. 3 we plot the selected endmembers.

FIGURE 2, 3, 4

TABLE1

4. Evaluation and limits of the linear method

Linear spectral unmixing can be a satisfactory approach for evaluating the composition of a surface (e.g. Combe et al. (2008); Bioucas-Dias et al. (2012)), even if a surface is a non-linear (intimate) mixture of several mineralogical phases. The linear assumption implies an error in the estimation of the abundances, but permits only a semi-quantitative estimation of the different components. In this regard, we performed two types of tests: 1) We selected different laboratory mixtures of possible Vesta analogues and we applied linear unmixing to the spectra of such mixtures. This exercise provides an idea of the uncertainty in the determination of the abundances. 2) We evaluate the stability of the solution for the VIR data by considering the ten cases showing smaller values of χ^2 . If for these χ^2 values we find the same endmembers and similar mixing coefficients, then the solution is stable and the results are reliable.

393 4.1. Test on laboratory mixtures

394 To perform this test, we use spectra for sets of mixtures available in the
 395 RELAB database, and those used in Cloutis et al. (2013) and Serventi et al.
 396 (2013). In Table 1 of the supplementary online material (SOM), we list the
 397 spectral endmembers of the mixtures, indicating the origin, the particle size
 398 and the texture. We selected mixtures with particle sizes similar to those
 399 inferred for Vesta. In particular, we consider mixtures of two and three
 400 laboratory spectra:

- 401 1. Olivine and orthopyroxene mixtures;
- 402 2. Orthopyroxene and plagioclase mixtures;
- 403 3. Millbillillie eucrite and Murchison carbonaceous chondrite;
- 404 4. Plagioclase, low and high calcium pyroxenes;
- 405 5. Olivine, low and high calcium pyroxenes;

406 In Section 1 of the SOM and in Fig. 5, all the results obtained for the
 407 laboratory mixtures are displayed.

408 We notice that the error in the estimation of the mixing coefficients de-
 409 pends on the number of endmembers in the mixture and on the endmember
 410 itself. Generally, for mixtures of two pyroxenes with a third phase (e.g.,
 411 plagioclase or olivine), we found differences between the true values and
 412 the modeled mixing coefficient up to 11% for the non-pyroxene endmember.
 413 In particular, for the case in which the third component is plagioclase, we
 414 have an underestimation of this phase of as much as 11% if its abundance
 415 is $> 50\%$. In mixtures with plagioclase contents $< 40\%$, the error is lower
 416 (Table 2 and Fig. 1 of the SOM). If olivine is one of the endmembers, the

417 difference between the measured abundances and the modeled mixing coef-
 418 ficients is about +10%, when the olivine abundance is $\sim 70\%$. For lower
 419 proportions of olivine in the mixture, the differences between measured and
 420 modeled values are decreased (Table 3 and Fig. 2 of SOM and Fig. 5).
 421 For mixtures of just two minerals, the results are similar for low calcium
 422 pyroxenes and olivine. The estimated error increases with greater amounts
 423 of olivine. In the case of an olivine content of 90%, we found the underes-
 424 timation of olivine to be 15% (Table 4 and Fig. 3 of the SOM). For lesser
 425 proportions of olivine, the error is below 8%. For plagioclase mixed with
 426 orthopyroxene at ratios ranging from 90:10 to 20:80, we find a consistent
 427 underestimation of plagioclase from 12 to 19% (Table 5 and Fig. 4 of the
 428 SOM).
 429 Two-component mixtures of Murchison CC with Millbillillie eucrite have a
 430 completely different trend. For Murchison abundances from 5% to 50%, the
 431 mixing model consistently overestimated the CC component. The error in
 432 CC abundance ranges from 11 to 26% despite the low χ^2 values (Table 6 and
 433 Fig. 5 of the SOM). We consider good model results to be those for which
 434 the difference between measured and modeled values is 5% (absolute) or less,
 435 and we deem results to be acceptable when the difference is $< 10\%$. Cases
 436 for which the difference between measured and the modeled values is over
 437 10% were rejected. Our tests on laboratory mixtures reveal that some mix-
 438 tures can be modeled successfully with the linear method, while some other
 439 mixtures may not. Mixtures of different pyroxenes and mixture of pyroxenes
 440 with olivine allow a good estimation of the mixing coefficients with the linear
 441 methods with respect to the other types of mixtures. In the case of Vesta, we

can spectrally model its surface as a mixture of at most three endmembers: pyroxenes, olivine, and a featureless component.

As mentioned previously, our analysis does not consider the absolute reflectance, thus we cannot distinguish high- and low-reflectance featureless phases, but rather the presence of a generic featureless component that weakens and modifies the bands. In the case of plagioclase, we find that the error on plagioclase mixing coefficient decreases with increasing number of endmembers. Unfortunately laboratory data for mixtures of two pyroxenes with CC are not available.

FIGURE 5

4.2. Stability of the method

To test the stability of the method we consider the results of the ten models showing the smaller χ^2 values. The method can be considered stable if, for small χ^2 variations, the results do not vary substantially. If small χ^2 variations lead to different endmembers, then the model results cannot be considered to be reliable. In Fig. 6 we plot the best-fit spectrum for the ten Aelia-SM cases with the lowest χ^2 . Variations in χ^2 of the order of 0.0001 led to variations of 1% in the abundances estimation, while the endmembers remain unchanged, indicating robust results. We performed this test on all the VIR spectra under consideration, and we obtained similar results. We thus can conclude that our method yields stable solutions for the choice of the endmembers and the estimation of the mixing coefficients.

FIGURE 6

5. Bright material units: Lithological variation as revealed by linear unmixing

A first statistical analysis of the BM units on Vesta indicates that the algorithm always chooses three endmembers, and two of these are generally eucrite and diogenite, with comparable frequency ($\sim 33\%$) (Fig. 7). The third endmember is the featureless one with a frequency of 24.8%, while olivine is present in few BM units with a frequency of 8.6% (Fig. 7).

Plots in Fig. 8 show a dependence between the modeled mixing coefficients for eucrite and diogenite found here and the Zambon et al. (2014) band centers for the BM units. We find that the content of eucrite increases with increasing values of band centers, while the content of diogenite increases with decreasing values of band centers. We observe a better correlation between the band II center and eucrite and diogenite mixing coefficient, than for the band I center. Generally, the band I center has a lower variability with respect to the band II center (see De Sanctis et al. (2012a); Ammannito et al. (2013a); Zambon et al. (2015)). The poorer correlation between the band I center and the eucrite and diogenite mixing coefficients is mainly due to VIR instrumental problems in the vicinity of band I, and on the bridging of the visible and infrared channels. Thus we observe a better correlation between mixing coefficients of eucrite and diogenite and the band II center than for the band I center.

In Fig. 9, we show the linear unmixing application for some representative

case of BM units. All the results have been reported in Table 2 and in section 2 of the SOM.

Aelia is a small crater (4.34 km) located at 14.26°S 140.6°E. It is characterized by mixed dark and bright material in its ejecta and by a high diogenite content in its bright slope material (Palomba et al., 2014; Zambon et al., 2014). Linear unmixing results indicate high mixing coefficient of diogenite (67%) for the Aelia-SM and only 9% of eucrite as expected. Aelia-RM which represents the ejecta showing a greater amount of eucrite than diogenite and a particularly large amount of the featureless endmember (33%). The lower reflectance values of this BM unit suggest the presence of a dark phase that is responsible for reducing the band depths (Fig. 9). Another diogenitic unit is Licinia-SM (Zambon et al., 2014), with mixing coefficients of diogenite (53%) higher than eucrite (30%) (Fig. 9). The bright units with shorter band centers are BU5-CWM and BU5-RM (Zambon et al., 2014), for these BM units we find a diogenite content of 65% and 57% respectively (Fig. 7 of the SOM) and BU8-CWM with a diogenite mixing coefficient of 65% (Fig. 8 of the SOM). Consistent with the findings of Zambon et al. (2014), the most eucritic BM units are Eumachia-CMW with an eucrite mixing coefficient of 57% and a corresponding diogenite content of 14% of diogenite (Fig. 9), and Tuccia-RM with 57% of eucrite and 27% of diogenite (Fig. 14 of the SOM).

Olivine on Vesta has been identified in few areas, most notably in the region of Bellicia and Arruntia craters, but smaller olivine amounts have been found also near the Albana and Pomponia craters, also located in the northern hemisphere (Palomba et al., 2015). Local olivine enrichments have been suggested by Ruesch et al. (2014) to be present also in the equatorial

516 region and in the southern hemisphere. Our analysis suggests that olivine is
 517 present in several BM units, with amounts between 12 and 53%. We identify
 518 sites with large olivine mixing coefficients ($> 48\%$) in correspondence with
 519 Arruntia, Bellicia in agreement with Ammannito et al. (2013b), and BU14
 520 indicated as detection 3 reported by Ruesch et al. (2014). In these spectra we
 521 observe a band I asymmetry typical of olivine and a reduction in the depth of
 522 band II, indicating a relatively high amount of olivine, but not high enough
 523 to completely obscure band II (Fig. 7). Among the possible olivine-bearing
 524 sites found by Ruesch et al. (2014), we note that the Licinia-SM BM unit
 525 has a mixing coefficient of olivine as high as 19%. Our algorithm chooses
 526 olivine as a third endmember also for regions different from those observed
 527 by Ammannito et al. (2013b) and Ruesch et al. (2014). Generally we find an
 528 olivine mixing coefficient $< 30\%$ for these sites, except for Calpurnia-CWM
 529 which has an olivine mixing coefficient of 35% (Fig. 10 of the SOM).

530 We cannot distinguish between opaque and neutral/featureless, high re-
 531 flectance components. In Fig. 10 we show the relationship between albedo
 532 of the BM units (Schröder et al., 2013) and the modeled mixing coefficient of
 533 the featureless endmember. In this plot, the albedo tends to decrease with
 534 increasing mixing coefficient of the featureless component. We interpret this
 535 trend as an indication that the algorithm's selection of the flat endmember
 536 can be attributed to the presence of an opaque dark phase in most cases.
 537 However, some exceptions exist. The bright streak called BU-15 SM is the
 538 brightest unit on Vesta, with a higher content of diogenite (47%) than eucrite
 539 (30%). This unit is different from the others because it is characterized by
 540 a reduction of the band depth with respect to the surroundings even though

it has high albedo (Zambon et al., 2014). The model composition of BU-15 SM is eucrite and diogenite, with a featureless endmember mixing coefficient of 23% (Table 2). Thus, BU-15 SM may represent a unique case on Vesta, where the presence of a high-albedo featureless component, such as plagioclase, plays an important role in controlling the spectral properties.

FIGURE 7, 8, 9,10

6. Discussion

The global compositional analyses hitherto performed on Vesta's surface with Dawn/VIR data were based on the spectral parameters of the kind used by De Sanctis et al. (2012a), and Ammannito et al. (2013b). Spectral parameters are very useful for determining the main lithology of a surface, but for a quantitative analysis mixing models are necessary.

In this work we focused our attention on eucrite and diogenite as the dominant components, plus olivine or a featureless material. We analyzed the Vesta spectra for BM units discussed by Zambon et al. (2014). In that paper, the authors give the results of a compositional analysis based on spectral parameters. Unlike previous papers, the present work determines a semi-quantitative estimation of the mixing coefficients of each lithology. Here we do not consider the spectral slope or the reflectance, and thus we cannot establish the specific nature of this featureless endmember. However, by examining the albedo of each BM unit, we can make inferences about the identity of the featureless phase.

We find that in general, the mixing coefficient of the featureless endmember

565 is anticorrelated with albedo, suggesting that this component is a low-albedo
 566 opaque material. In fact, bright areas are often contaminated by dark ma-
 567 terial (Jaumann et al., 2012; Palomba et al., 2014). In Fig. 11, we show the
 568 main results from the linear unmixing model plotted on the compositional
 569 map from Ammannito et al. (2013b). The mixing coefficients of the different
 570 components obtained by the model and the distribution of the BM units are
 571 generally in good agreement with previous analyses (e.g., De Sanctis et al.,
 572 2012a, 2013b; Zambon et al., 2014). First, in all of the BM units, both eucrite
 573 and diogenite spectra are used to fit VIR spectra, reflecting the distribution
 574 of those components in the regolith (De Sanctis et al., 2013a,b).
 575 From the results of our analysis, we can see that several BM units are dom-
 576 inated by eucrite and diogenite components (see Section 8 and Table 2). In
 577 particular, we can classify these units as eucrite-rich units and diogenite-rich
 578 units. A third type is olivine-rich BMs.
 579 The distribution of eucrite-rich and diogenite-rich units is similar to the com-
 580 position of surrounding terrains as inferred by the analysis of spectral param-
 581 eters.
 582 Olivine-rich BM units are associated with recognized olivine-bearing craters
 583 (Ammannito et al., 2013b; Ruesch et al., 2014). Moreover, where more
 584 units are seen in the same crater, they frequently belong to only one family.
 585 The only exceptions are a few diogenite-rich units present on eucritic ter-
 586 rains (Aelia, Canuleia, BU3, BU6), and few craters (Aelia, Canuleia, Oppia,
 587 Justina, BU9) where BMs show both compositions in the same location.
 588 Eucrite-rich BM units may have a mixing coefficient of eucrite as high as 44%
 589 (red symbols in Fig.11). Those BM units are mostly located in the equatorial

region, between 90°E and 300°E longitude, and their distribution is compatible with eucrite-rich regions (Ammannito et al., 2013a). Furthermore, they can be subdivided in two populations: 1) one where the mixing coefficient of diogenite is higher than the mixing coefficient of the featureless endmember (diamond); 2) one where the mixing coefficient of the featureless endmember is greater than the mixing coefficient of diogenite (square). Those subgroups are equally distributed in number and by geography.

In eucrite-rich BM units, the olivine component was never used to fit the spectra. This is consistent with the rare appearance of olivine, as secondary mineral, in eucrite meteorites (Mittlefehldt et al., 1998). There is just one exception, the crater wall material in BU2 (BU2 CWM in Table 2), where the BM is fitted by the Serra de Mage eucrite spectrum (86%), with almost negligible contribution from olivine and the featureless endmember. This is also the only unit where both eucrite and diogenite spectra were not selected. Note that the Serra de Mage eucrite shows spectral characteristics very close to howardite spectra. The spectra of this eucrite is selected only in one other case: the model for an anomalous diogenite rich-units (Canuleia SM) where only HED spectra were used to model the data.

The diogenite-rich BMs are characterized by the association of the highest mixing coefficient of diogenite spectra (yellow symbols in Fig.11), with a lower threshold of 40%, except for the crater wall unit in BU9 (37%). The distribution of diogenite-rich BMs is widespread in longitude and in latitude, extending farther to the south than eucrite-rich units. The distribution of diogenite-rich BMs is also consistent with the results of previous works and the spectral parameter maps, and they are generally correlated with dio-

615 genitic and howarditic terrains (Ammannito et al., 2013a).

616 Only two regions show diogenite-rich bright materials, within eucrite ter-
 617 rains, the Aelia unit and the units around 300°E, where are concentrated
 618 BU3, BU6, and the Canuleia unit. The diogenite-rich BM units can be dif-
 619 ferentiated by the presence of the eucrite (diamonds in Fig. 11) or another
 620 components (square) as the second spectra in the unmixing solutions. Unlike
 621 eucrite-rich BM units, the sub-group with the spectra of eucrites selected as
 622 the second component is the most frequent, whereas few units are character-
 623 ized by a non-HED component selected as the second endemember.

624 Moreover, in various cases ($\sim 8.6\%$) olivine was used instead of the fea-
 625 tureless component (white points within diogenite-rich BMs), even if with
 626 mixing coefficient lower than 30%, with the exception of the crater wall unit
 627 in Calpurnia (35%). Aelia and Canuleia craters are both in eucritic terrains
 628 and they show both eucrite-rich and diogenite-rich BM locations. Aelia has
 629 diogenite-rich type 1 BMs on the wall, whereas the radial material (RM)
 630 is eucrite-rich. This could indicate that the Aelia impact exposed diogenite
 631 blocks from greater depths, whereas fresh eucritic material that originally
 632 resided close to or at the surface was ejected to form the radial material.
 633 In Canuleia, we have both type 1 and 2 BMs with eucrite and diogenite
 634 compositions, but the differences between the two HED components are less
 635 than 64%, indicating an homogeneous distribution of eucrite and diogenite
 636 components, i.e., howarditic.

637 Only one Aelia BM unit, the RM, shows a clear eucrite-rich composition.
 638 Thus, even in this location, diogenite material could have been exposed by the
 639 impact and mixed with eucrite (i.e., formation of howarditic breccia blocks),

640 while some eucrite-rich fresh material was ejected from the crater, and de-
641 posited on the surrounding eucritic crust.

642 In contrast, BU3 and BU6 craters show both diogenite-rich units as RM
643 (BU6 has also a SM, see Table 2). Those units are very close to the diogenite
644 terrains, so perhaps the presence of diogenite-rich material can be attributed
645 to contributions from southern craters. Alternately, it could be that the eu-
646 critic crust in that area is thinner, and thus fresh diogenitic material was
647 more easily excavated and ejected from those craters.

648 Three other craters (Oppia, BU9 crater, and Justina) show the presence of
649 both eucrite- and diogenite-rich BMs. Oppia and BU9 show both units. In
650 the Oppia crater, two BM units are on the wall. The CWM are diogenite-
651 enriched, whereas the corresponding slope material (SM) is characterized by a
652 higher amount of eucrite. In the second crater (BU9), the CWM is diogenite-
653 rich, whereas a RM BM unit is fitted by a high mixing coefficient of eucrite.
654 BM units at crater Justina have been covered by VIR more than one time,
655 the four wall units (both CWM and SM) are eucrite-rich and the two RM
656 are diogenite-rich BM units. A third family is made of olivine-rich bright
657 units, i.e., where olivine spectra are selected as the most abundant compo-
658 nent. In this family, all of the recognized BMs are enriched in olivine, with
659 diogenite > eucrite in BU14 and eucrite > diogenite in Arruntia and Bellicia
660 areas. These proportions are in agreement with those found by analysis of
661 spectral parameters in Zambon et al. (2014). These units are located in the
662 northern hemisphere and correspond to the olivine-bearing sites detected by
663 Ammannito et al. (2013b) and Ruesch et al. (2014). For Bellicia and BU14,
664 we find an olivine mixing coefficient around of $\sim 50\%$, with slightly lower

values found in Arruntia. These values indicate that olivine is a dominant component in these sites (Ammannito et al., 2013b).

High abundances of olivine are generally expected to be present in mantle and plutonic rocks, which could be present in topographically low areas. In this work, we have employed Mg-rich olivine endmembers with very fine particle sizes, compatible with the predicted Vesta dominant regolith size. Therefore, the absolute abundance values could be lower than suggested by our models, in particular considering that olivine is more fayalitic (Fe-rich) in HEDs. Olivine that is richer in iron has a stronger absorption band, and hence a lower abundance of such Fe-rich olivine (relative to Mg-rich olivine) would be needed to model a given spectrum.

Poulet et al. (2015) discussed the retrieval of composition from VIR spectra by using a non-linear method (Shkuratov et al., 1999). It should be noted that the results from a non-linear technique, although in principle better than those from a linear model, can suffer if the starting data has not been properly treated. For the VIR dataset, extreme care must be taken, for example, in bridging the sections VIS and IR, exclusion of spectral channels suffering from recurring artifacts, and removal of the continuum. Poulet et al. (2015) concluded that coarse-grained olivine is likely to be present in all major units of Vesta. In general, they also found that the abundance of olivine could be lower because of the presence of iron-rich olivine or because of an increase in the grain size of olivine with respect to the other phases. An alternative hypothesis that could explain higher surface abundances of olivine in such northern locations is an exogenic origin - e.g., material delivered by an olivine-rich impactor, as suggested by Le Corre et al. (2015). Moreover,

we find that in those terrains the third selected endmember is frequently a featureless component.

As discussed above, the mixing coefficient of the featureless endmember is anti-correlated with the albedo, suggesting that this component is associated with low-albedo CC-like opaque material. In fact, bright areas are often contaminated by dark material (Jaumann et al., 2012; Palomba et al., 2014). We do find one exception where a BM unit with a large mixing coefficient for the featureless endmember, BU15-SM, has the highest albedo (Zambon et al., 2014). This unit is matched by a diogenite-rich bright unit extending into the Rheasilvia basin, and Zambon et al. (2014) predicted the possible presence of a bright component.

The major bright, iron-poor, mineral in HEDs and in basic igneous composition is plagioclase, which can have a large abundance in magmatic rocks, both effusive and cumulitic, intrusive rocks. Plagioclase can exhibit a characteristic absorption at $1.25\mu\text{m}$ caused by impurity iron at abundances corresponding to $\text{FeO} < 1\%$ (e.g. Burns, 1993; Cheek et al., 2011; Serventi et al., 2013). However, the $1.25\mu\text{m}$ feature cannot be recognized in mixtures with mafic iron-bearing silicates unless the plagioclase is present in very large amounts. If plagioclase is indeed revealed by our analysis, it should be noted that in HEDs it is generally present in eucrites and cumulitic eucrites, but not in diogenites.

Another important aspect showed by our results is the selection of olivine spectra with low mixing coefficient for the diogenite-rich BM units, in place of the featureless endmember. From a spectroscopic viewpoint, this implies that reductions of the intensities of band I and band II are not correlated,

otherwise the model should have selected the featureless spectrum. In addition, a weak asymmetry around the $1\text{-}\mu\text{m}$ absorption is revealed, so the selection of only eucrite and diogenite spectra alone is not able to properly fit VIR data in that specific spectral region. The presence of this low amount of olivine in the diogenite-rich BM units indicates that iron could be present in one or more phases with a band around $1\text{ }\mu\text{m}$ but with a very weak, or absent, absorption around $2\text{ }\mu\text{m}$. In general, these spectral characteristics are consistent with the presence of olivine, high-Ca pyroxene, plagioclase or a glassy component, in basic igneous compositions.

Iron-bearing silicate glasses (e.g., volcanic or impact glasses) have a dominant crystal-field absorption at $\sim 1.1\text{ }\mu\text{m}$, the intensity of which depends on the iron abundance. Thus a glassy component cannot be excluded in units that are formed by impacts. In fact, impact melts have been identified in some HEDs (see e.g., Metzler and Stöffler, 1995; Singerling et al., 2013). High-Ca pyroxenes are not abundant in diogenites, even if there are samples where they are found in abundance higher than 10% (e.g., Mittlefehldt et al., 1998). High-Ca pyroxene can also be frequently found as exsolution lamellae in orthopyroxene crystals, influencing the spectral characteristics of the host low-Ca hosting pyroxene (Gaffey et al., 2002).

Olivine is difficult to detect, especially in small amounts. The shape and the position of the $1\text{ }\mu\text{m}$ band are influenced by the presence of olivine if it is present in high amounts in mixtures with pyroxene (e.g. in particular the orthopyroxene-olivine mixtures studied by Cloutis et al., 1986; Beck et al., 2013).

Beck et al. (2013) argued that olivine could be recognized only at contents

740 > 30% in HEDs, because under this threshold, the olivine is masked in
 741 mixtures with orthopyroxene. (Beck et al., 2013) predicted that olivine in
 742 diogenite and harzburgite terrains could not be detected when present in
 743 amounts below this threshold. Recent studies by Horgan et al. (2014) and
 744 Shestopalov et al. (2015) suggest two different methods for detection of small
 745 amounts of olivine in mixtures with pyroxene. Horgan et al. (2014) used dif-
 746 ferent spectral indices to identify the presence of olivine in the spectra of
 747 Mars. Shestopalov et al. (2015) observed that the uncertainty (variance) of
 748 the band center position in spectra of laboratory olivine-pyroxene mixtures
 749 depends on the shift of the band center at 950 nm. They found that mixtures
 750 with different olivine contents can be identified by their position in a plot of
 751 the variance vs. band center at 950 nm.
 752 On the other hand, Poulet et al. (2015) concluded that best fits to all the stud-
 753 ied Vesta terrains are achieved when a component of forsteritic olivine (Fo70)
 754 is included. They also emphasized that the particle size has an effect on the
 755 olivine abundance obtained by a non-linear unmixing method. Olivine-rich
 756 melts were recently found in some howardites (Beck et al., 2013), so we can-
 757 not rule out that those BM units could have low olivine mixing coefficient.

758 FIGURE 11

759

760 7. Conclusions

- 761 1. To model VIR data for Vesta, we applied a linear spectral unmixing
- 762 model. Our analysis does not take into account the spectral slope and
- 763 the albedo but only the spectral shape.

2. Tests on laboratory mixtures of olivine and low- and high-calcium pyroxene indicate that olivine mixing coefficients are underestimated with an accuracy within 10% for olivine amounts $> 50\%$. For mixtures of plagioclase with low- and high-calcium pyroxenes, the plagioclase amount is underestimated within 11% for plagioclase contents $> 40\%$. Analysis of mixtures of Millbillillie eucrite with carbonaceous chondrite (CC) material indicates that the CC abundance was overestimated by up to 26%.
3. Linear unmixing applied to Vesta spectra is able to detect the principal HED and non-HED lithologies present on the asteroid.
4. We can divide the bright material units into three main groups: eucrite-rich, diogenite-rich, and olivine-rich, depending on the most abundant endmember used to fit the spectrum. The non-HED endmember can be either olivine or a featureless component. This confirms that Vesta is composed of a high content of pyroxenes mixed with a lower concentration of other mineralogical phases.
5. The third endmember is often the featureless phase, which is associated with a reduction in the strength of both pyroxene bands. The mixing coefficient of the featureless endmember is anticorrelated with albedo, suggesting that the featureless phase can be associated with low-albedo, CC-like opaque material.
6. Large amounts of olivine are detected in the Bellicia, Arruntia and BU14 BM units. Where lower olivine contents ($< 30\%$) are found, the sites are generally associated with high concentrations of diogenite.

Figure and Table captions

Figure 1: Example of VIR spectrum before (top) and after (bottom) the removal of the slope.

Figure 2: Spectral parameter plots for the diogenite and eucrite laboratory samples. The gray circles indicate the samples selected for use as endmembers in this study. The numbers refer to the endmember names in Table 1.

Figure 3: Spectra of the endmembers selected. The text labels refer to the sample names in Table 1 (6e Bereba, 7e Bouvante, 8e Cachari, 20e Padvarninkai, 22e Serra de Mage, 24e Y74450, 2d ALH77256, 5d GRO95555, 6d Johnstown, 8d Tatahouine, 10d Y75032, 1o Jackson cty, 2o Green olivine).

Figure 4: Frequency of the endmembers for the linear unmixing with two endmembers.

Figure 5: Linear unmixing applied to a mixture of olivine with low and high calcium pyroxene. The band centers of the modeled spectra are shifted slightly with respect to those of the laboratory spectra. In cases where the χ^2 values differ from 0, there is an imperfect match between the measured and modeled spectra.

Figure 6: The results of linear unmixing applied to the Aelia BM unit for the 10 cases with the lowest χ^2 . These results indicate that small variations in χ^2 do not correlate with changes in the endmembers or with large variations in the mixing coefficients of the endmembers, attesting to the stability of the results. Black lines are the measured spectra, the gray lines are the model mixture spectra.

Figure 7: Frequency of the endmembers selected for the linear unmixing

with three endmembers.

Figure 8: Plots of band centers vs. eucrite mixing coefficients show that eucrite content increases for increasing band center values. Right: Similar plots for band centers and diogenite mixing coefficients indicates that diogenite content decreases with increasing band center values. Band center values for the BM units are those calculated in Zambon et al. (2014). The coefficient of determination (i.e., the squared correlation coefficient R^2) is also provided.

Figure 9: Examples of linear spectral unmixing applied to BM units. Black lines are the measured spectra, the gray lines are the model mixture spectra. Table 1 summarizes the model endmember mixing coefficients for all the BM units. Related plots are presented in the SOM.

Figure 10: The albedo of each BM location plotted against the modeled mixing coefficient of the featureless endmember. As albedo decreases, the mixing coefficient of the featureless endmember increases.

Figure 11: Results obtained for the linear unmixing applied to BM units overlain on a mineralogical map of Vesta. Red color indicates eucrite-rich units, yellow color designates the diogenite-rich units, and the white symbols are the olivine-rich bright materials. Diamonds indicate BM units for which the eucrite or diogenite mixing coefficient is larger than that of the third endmember. Squares correspond to BM in which the eucrite or diogenite amount is lower than that of the third component. The solid white points are BM units with olivine as the third endmember (See also Table 2). The legend at the top refers to the different cases found in our analysis. For example, "D>III" indicates that the mixing coefficient of the diogenite

is greater than those of the third component. "E>III" represents the cases in which the mixing coefficient of the eucrite is greater than that of the third endmember, and so on

Table 1: Sample name, type, band centers, and band depths for all the endmembers in our spectral library. Each spectrum is assigned an identification number that contains "e" for eucrite, "d" for diogenite, and "o" for olivine.

Table 2: Summary of the results obtained from application of linear spectral unmixing on BM units on Vesta. Endmembers are designated as eucrite ("E"), diogenite ("D"), or olivine ("Ol"), while "Line" refer to the featureless component. Different text color and box color are indicative of variations in composition: Red text refers to eucrite-rich BM units where the featureless endmember (line) is the second most abundant. Yellow text indicates diogenite-rich endmember in which the second most abundant endmember is either the featureless one or olivine. Green text represents cases with olivine as the non-HED spectrum. Red boxes correspond to eucrite-rich BM units with diogenite as the second most abundant endmember. Yellow boxes are diogenite-rich BM units with eucrite as the second most abundant endmember. Green boxes represent the olivine-rich BM units. In the last column are reported the values of the correlation coefficient r .

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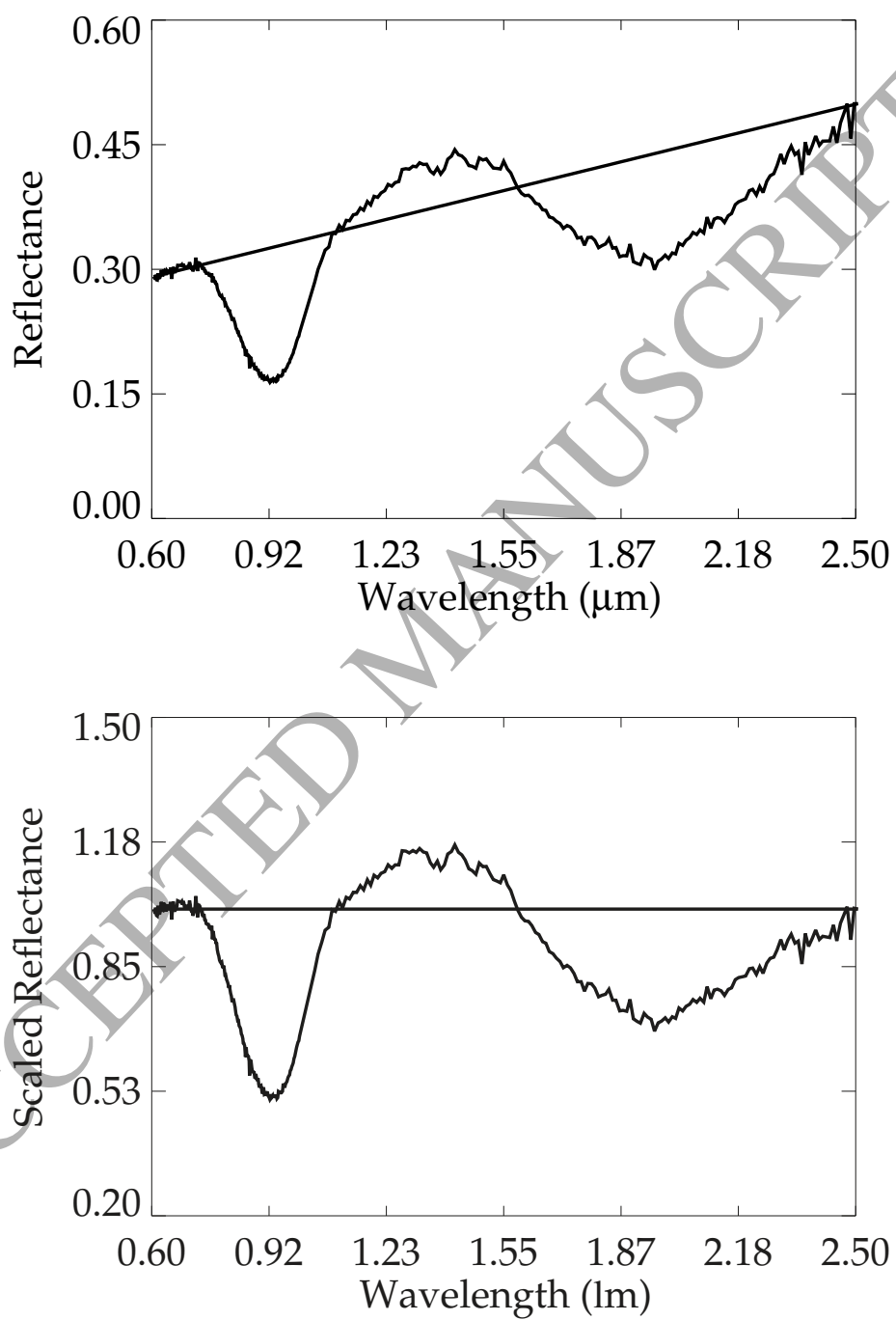


Figure 1

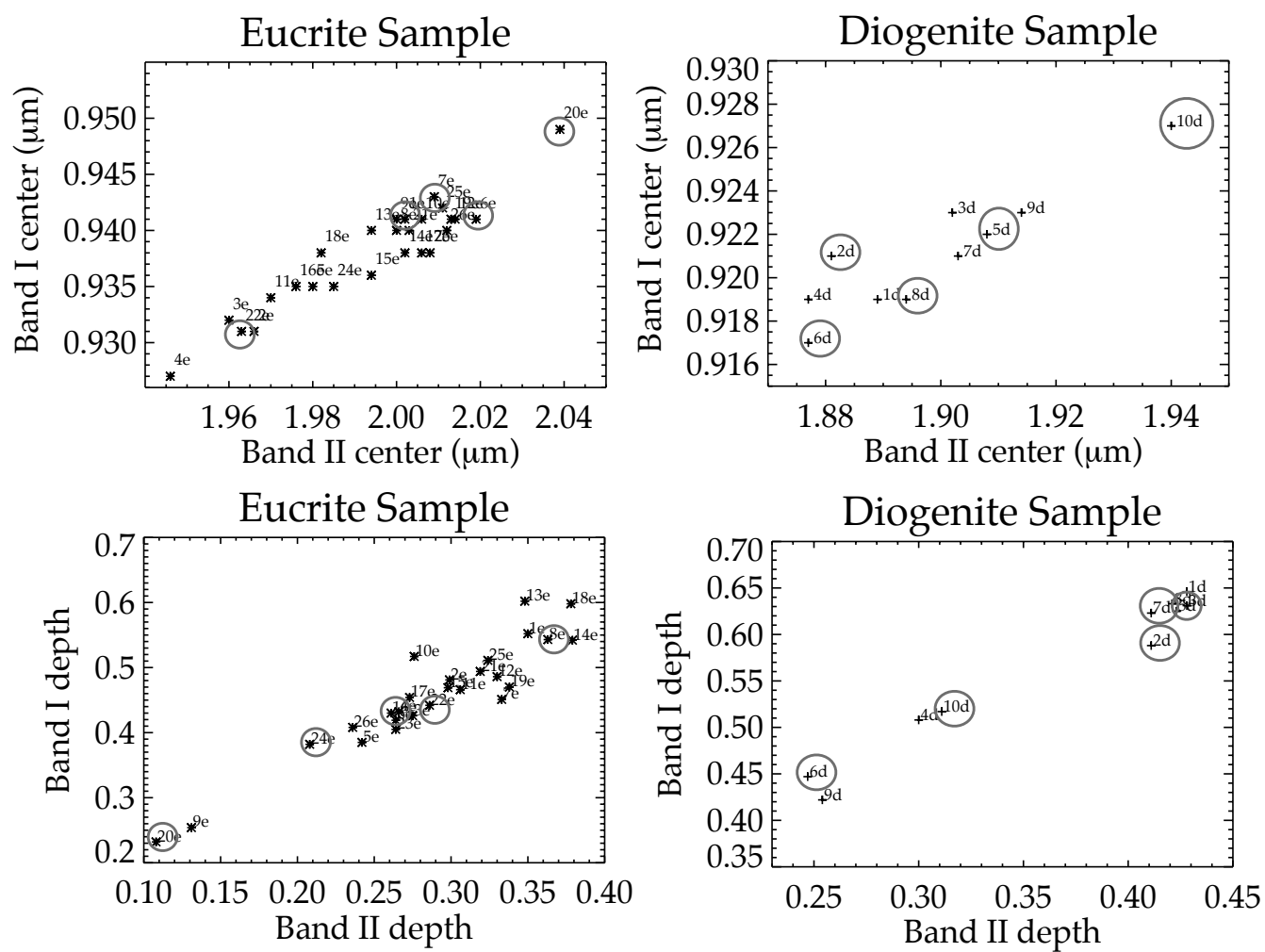


Figure 2

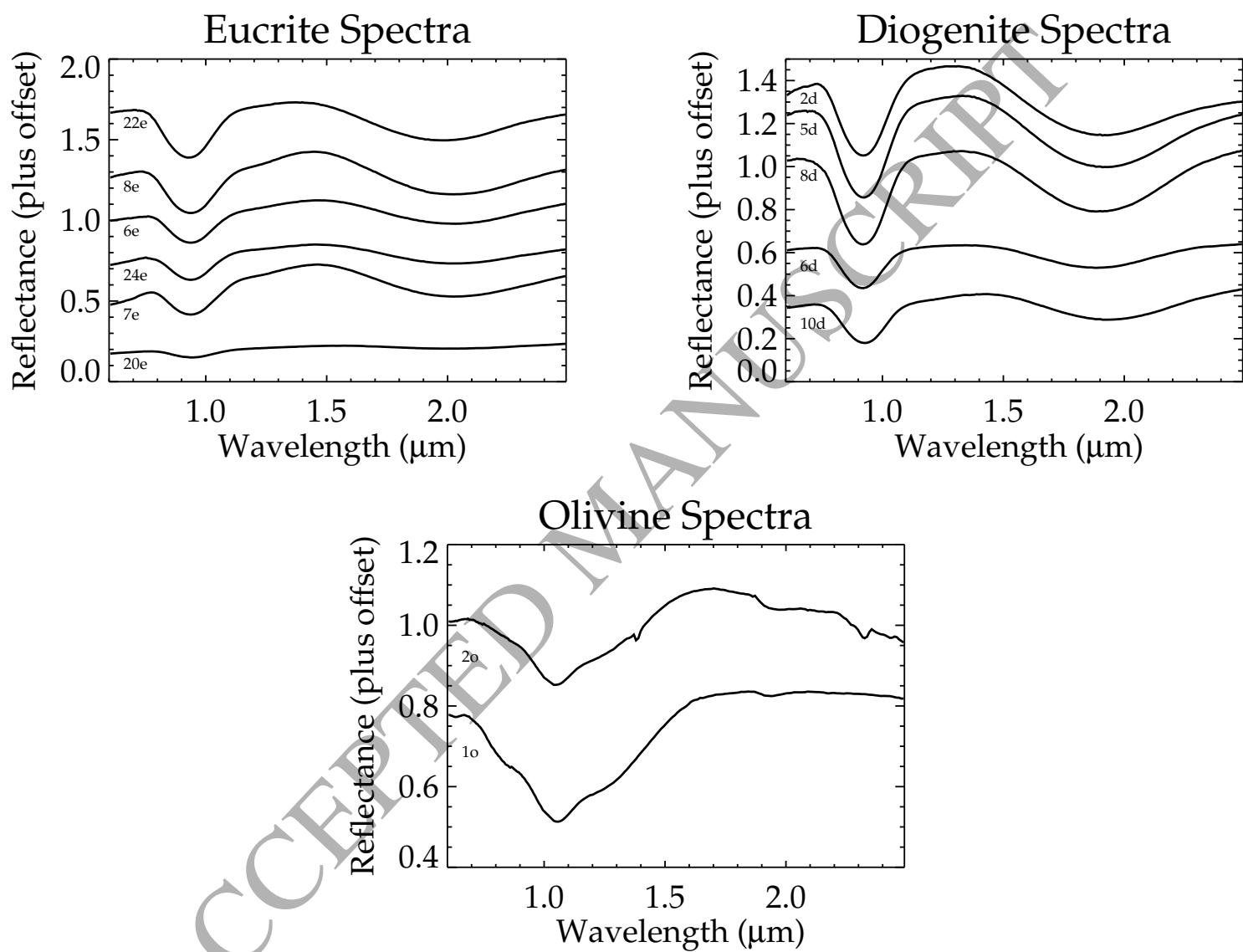


Figure 3

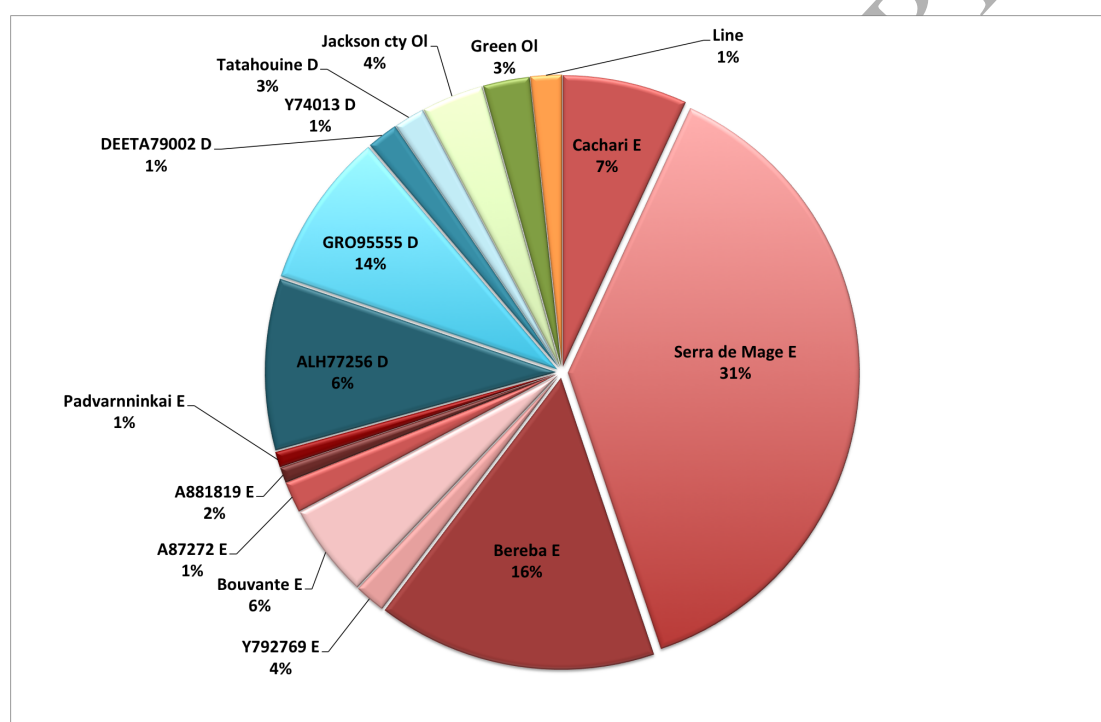


Figure 4

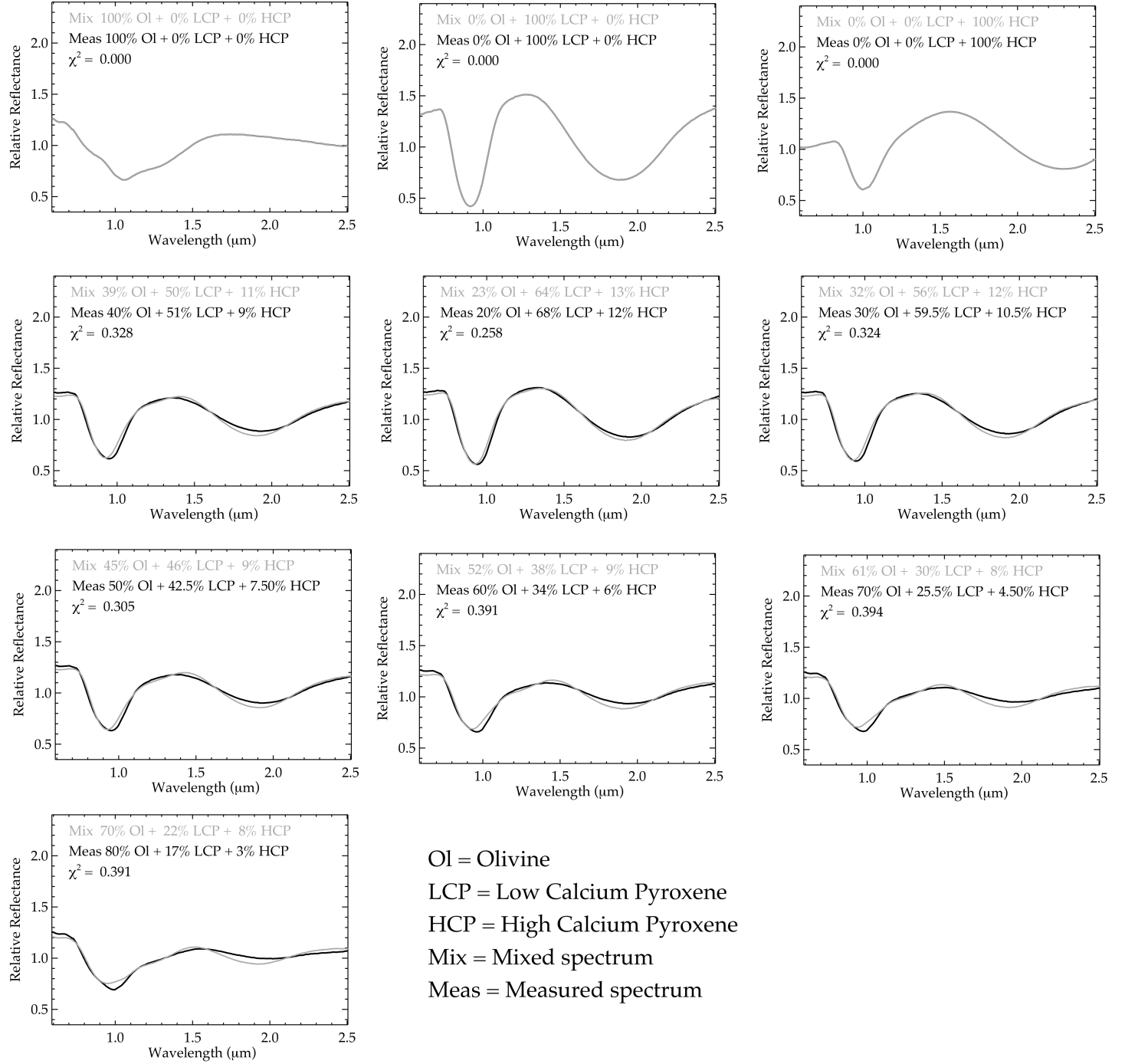


Figure 5

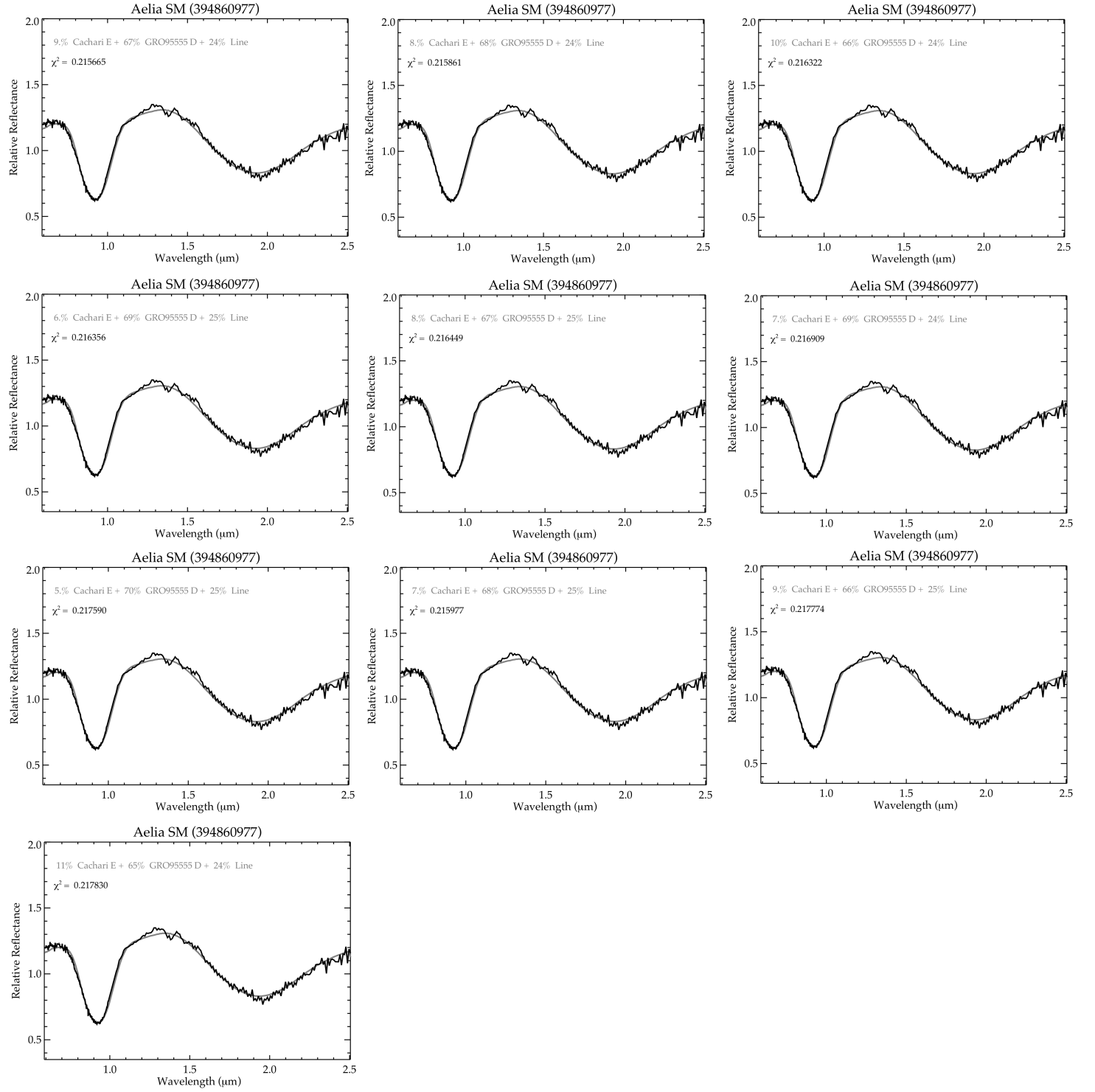


Figure 6

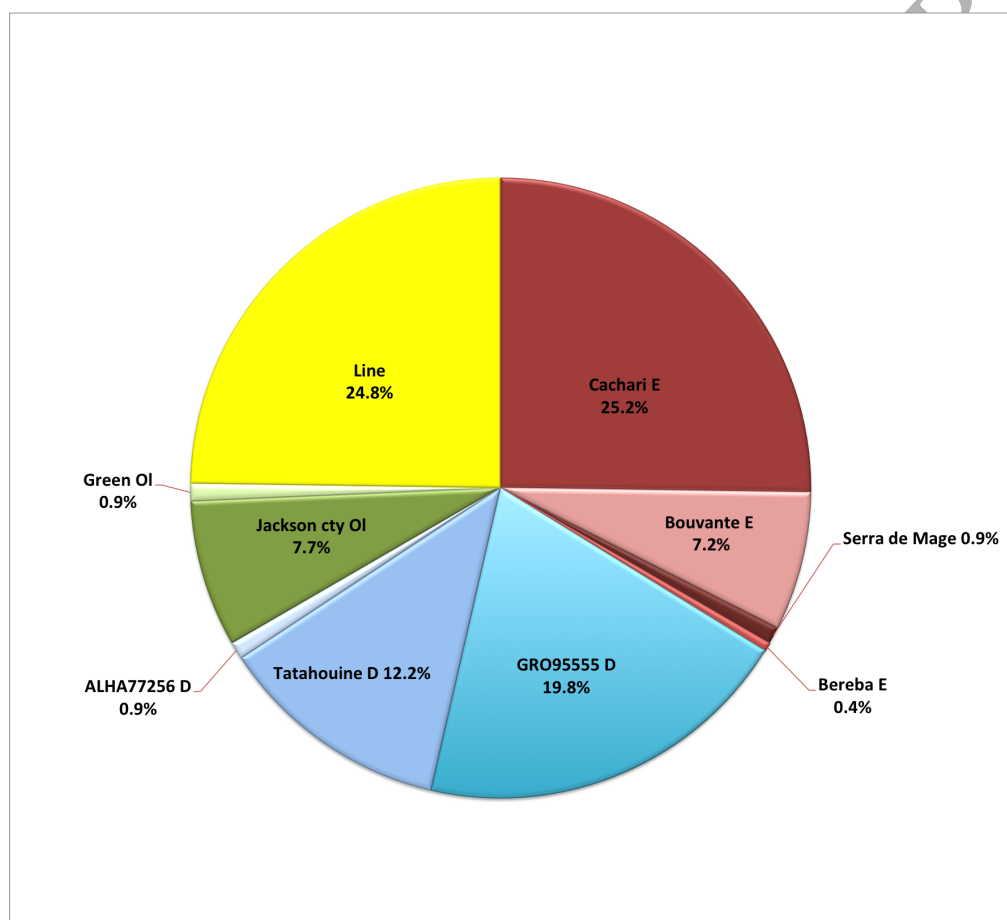


Figure 7

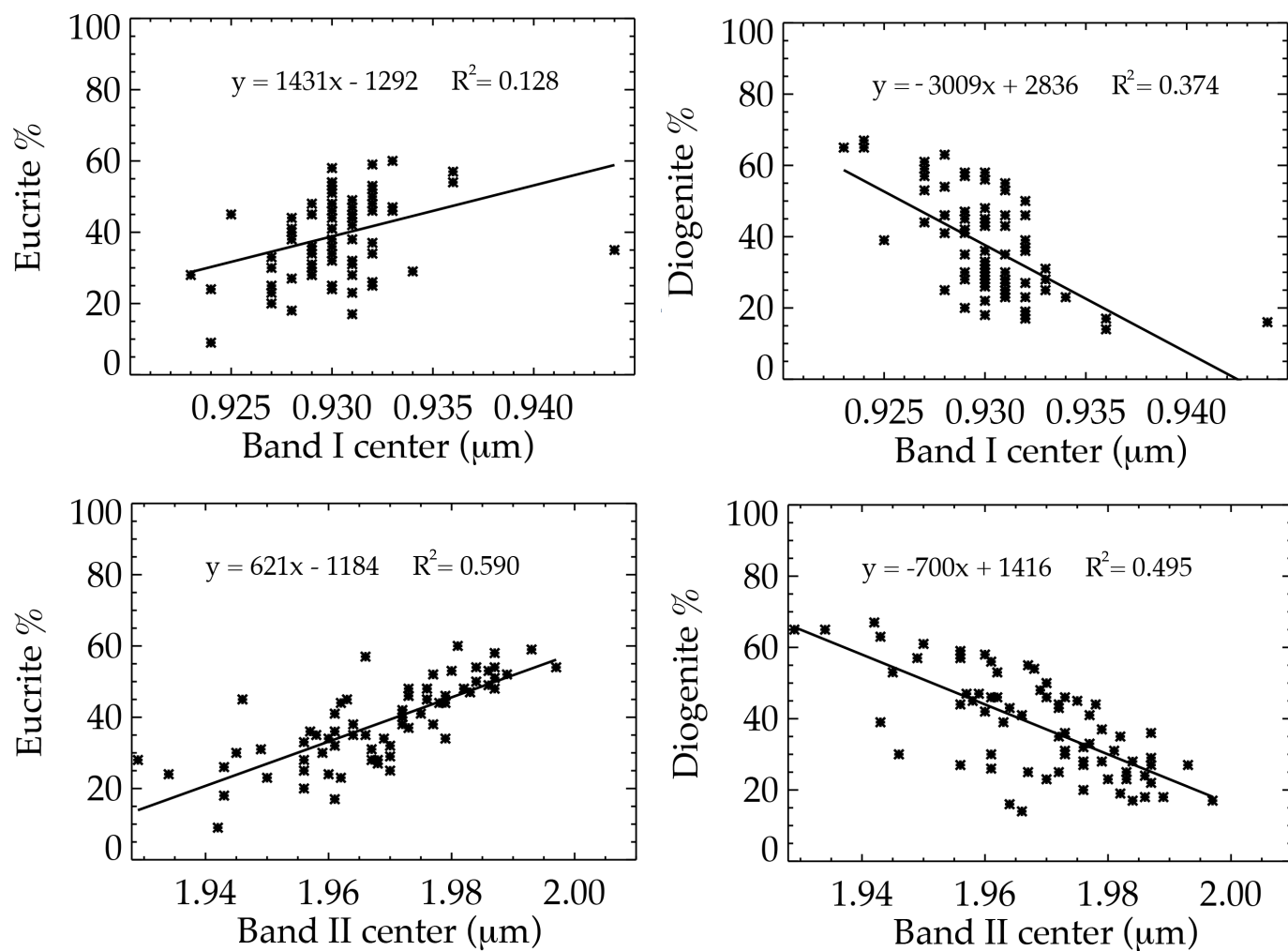


Figure 8

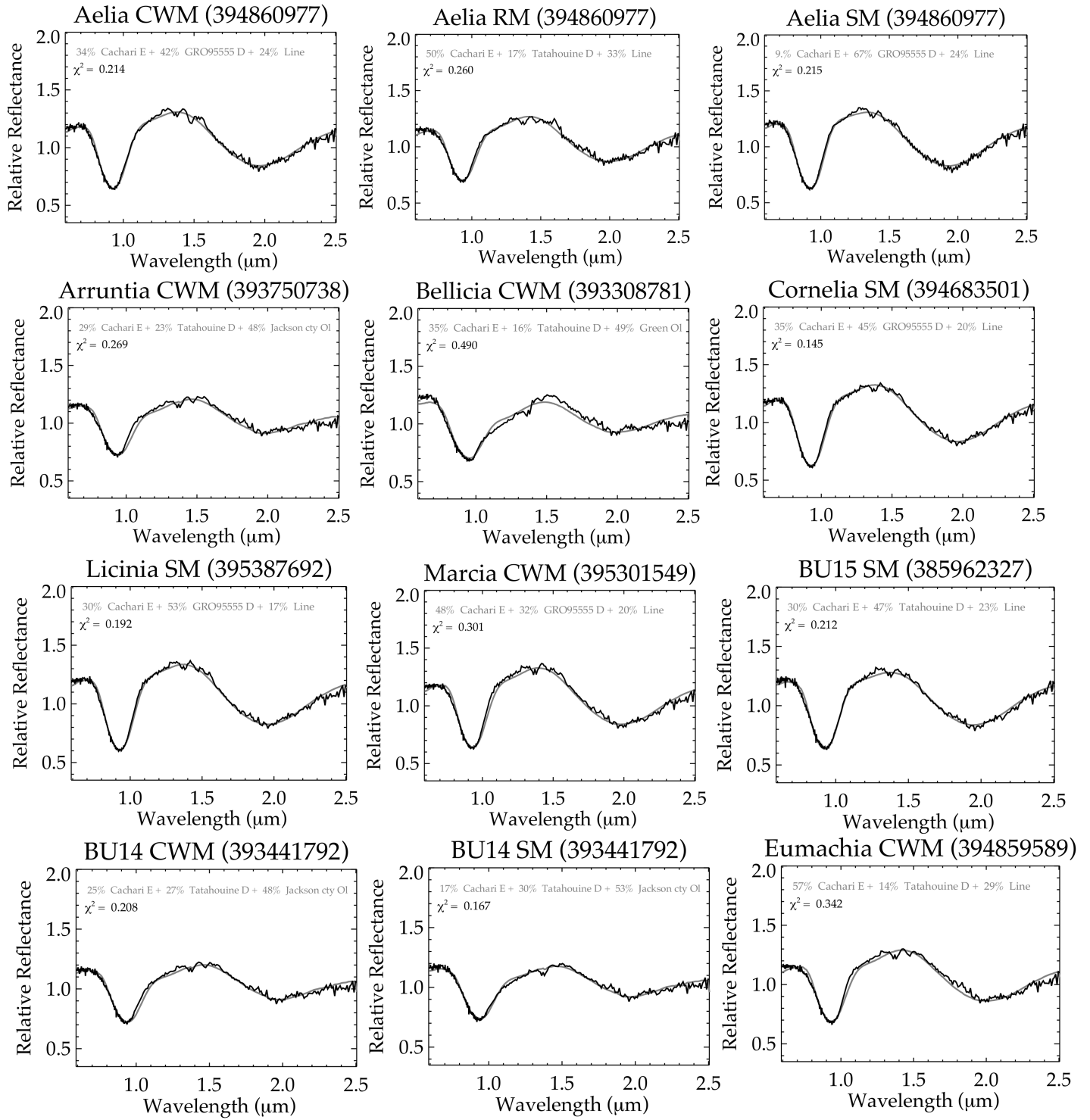


Figure 9

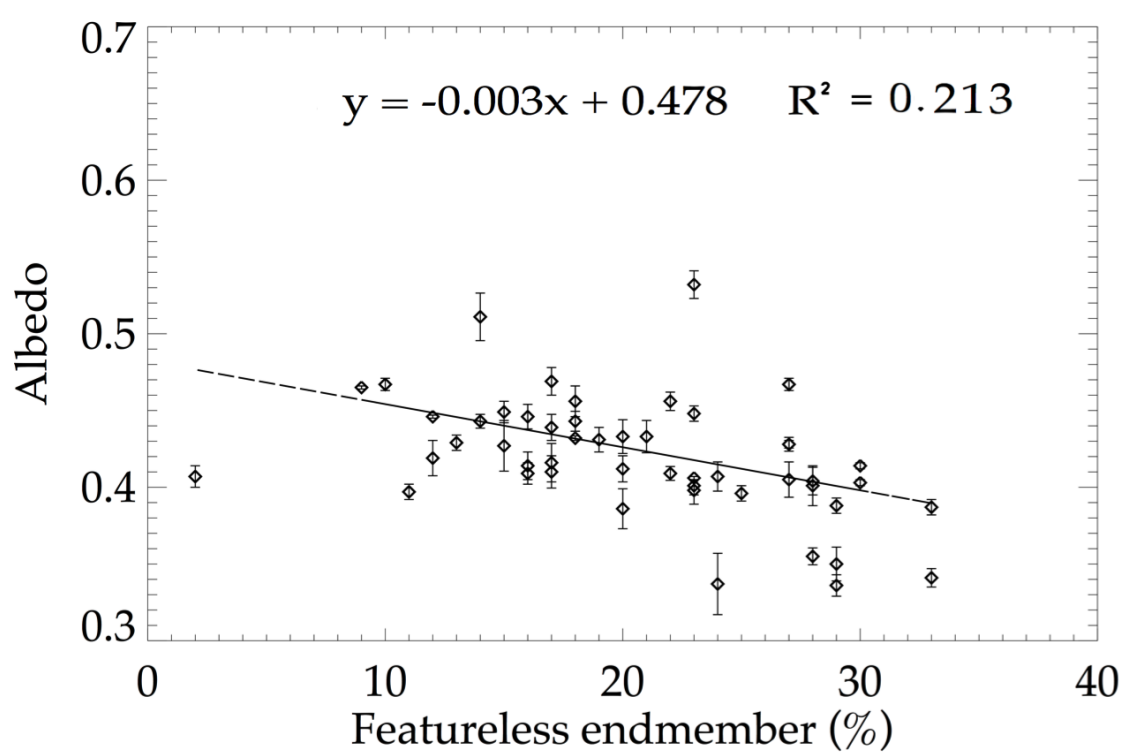


Figure 10

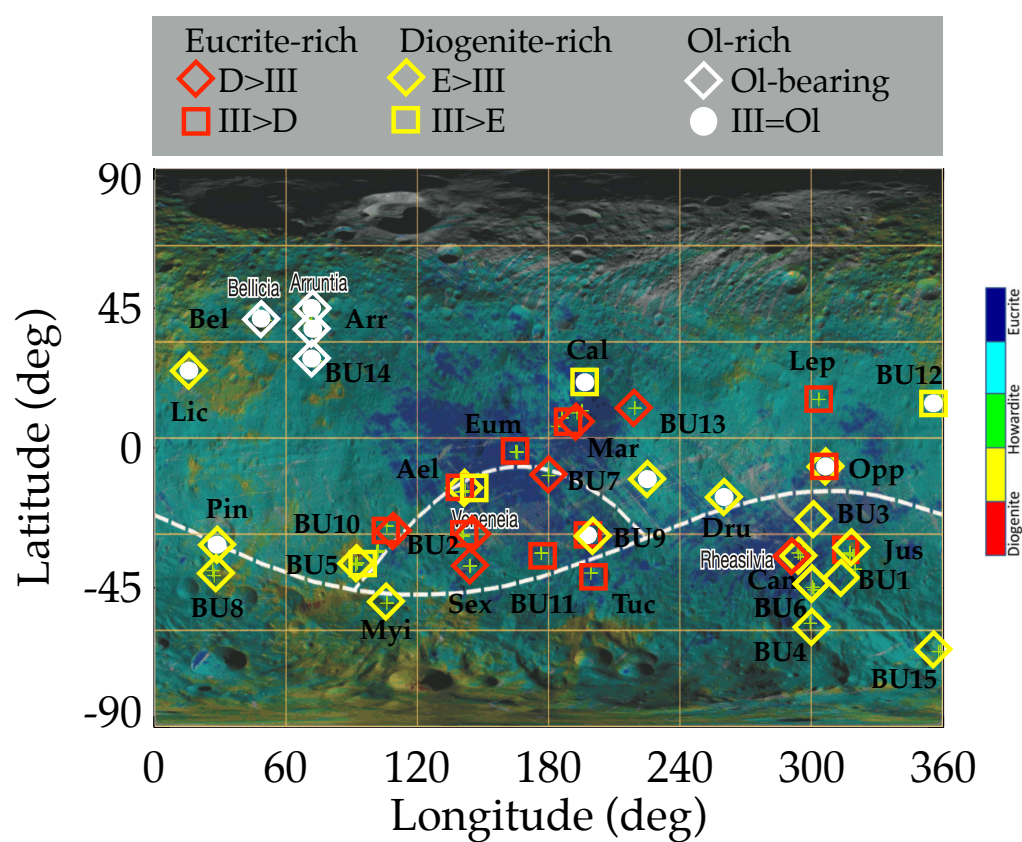


Figure 11

Endmembers						
#	name	type	BCI	BCII	BDI	BDII
1e	A-87272	Eucrite-mmict	0.941	2.002	0.552	0.350
2e	A-881819	Eucrite	0.931	1.966	0.481	0.299
3e	ALH78132	Eucrite-pmict	0.932	1.960	0.426	0.275
4e	ALH85001	Eucrite-Mg rich	0.927	1.946	0.432	0.266
5e	ALHA76005	Eucrite-pmict	0.935	1.980	0.385	0.242
6e	<u>Bereba</u>	Eucrite-mmict	0.941	2.019	0.419	0.264
7e	<u>Bouvante</u>	Eucrite-mmict	0.945	2.009	0.451	0.333
8e	<u>Cachari</u>	Eucrite-mmict	0.940	2.000	0.543	0.363
9e	EET 87542	Eucrite-br	0.941	2.000	0.254	0.131
10e	EET 90020	Eucrite-unbr	0.941	2.006	0.517	0.276
11e	EETA79005	Eucrite-pmict	0.934	1.970	0.466	0.306
12e	GRO 95533	Eucrite-br	0.941	2.014	0.486	0.330
13e	Ibitira	Eucrite-mmict	0.940	1.994	0.602	0.348
14e	Jonzac	Eucrite-mmict	0.938	2.002	0.542	0.379
15e	Juvinas	Eucrite-mmict	0.936	1.994	0.469	0.298
16e	LEW 87004	Eucrite-pmict	0.935	1.976	0.430	0.261
17e	Millbillillie	Eucrite-mmict	0.938	2.006	0.454	0.273
18e	Moore County	Eucrite-cm	0.938	1.982	0.598	0.378
19e	PCA 82502	Eucrite-unbr	0.941	2.013	0.470	0.338
20e	<u>Padvarninkai</u>	Eucrite-mmict	0.949	2.039	0.232	0.108
21e	Pasamonte	Eucrite-pmict	0.940	2.003	0.494	0.319
22e	<u>SerradeMage</u>	Eucrite-cm	0.931	1.963	0.442	0.286
23e	Stannern	Eucrite-mmict	0.938	2.008	0.405	0.264
24e	<u>Y74450</u>	Eucrite-pmict	0.935	1.985	0.382	0.208
25e	Y792510	Eucrite-mmict	0.942	2.011	0.511	0.324
26e	Y792769	Eucrite-pmict	0.940	2.012	0.408	0.236
1d	A881526	Diogenite	0.919	1.889	0.646	0.428
2d	<u>ALH77256</u>	Diogenite	0.921	1.881	0.588	0.411
3d	Aioun el Atrouss	Diogenite-pm	0.923	1.902	0.631	0.428
4d	EETA79002	Diogenite	0.919	1.877	0.508	0.300
5d	<u>GRO95555</u>	Diogenite-an	0.922	1.908	0.625	0.428
6d	<u>Johnstown</u>	Diogenite	0.917	1.877	0.447	0.247
7d	LAP 91900	Diogenite	0.921	1.903	0.623	0.411
8d	<u>Tatahouine</u>	Diogenite	0.919	1.894	0.633	0.421
9d	Y74013	Diogenite	0.923	1.914	0.422	0.254
10d	<u>Y75032</u>	Diogenite	0.927	1.940	0.517	0.311
1o	<u>Jackson cty</u>	Olivine	1.056	-	0.361	-
2o	<u>Green olivine</u>	Olivine	1.054	-	0.226	-

Table 1

Linear unmixing results applied to the BM units									
BM unit	Cube	Endm 1	Abundance	Endm 2	Abundance	Endm 3	Abundance	χ^2	r
Arruntia CWM	(393750738)	Cachari E	0.29	Tatahouine D	0.23	Jackson cty Ol	0.48	0.269	0.984
Arruntia RM	(393750391)	Bereba E	0.36	Tatahouine D	0.26	Jackson cty Ol	0.38	0.194	0.966
Arruntia SM	(393750738)	Cachari E	0.31	GRO95555 D	0.25	Green Ol	0.44	0.264	0.988
Aelia CWM	(394860977)	Cachari E	0.34	GRO95555 D	0.42	Line	0.24	0.214	0.992
Aelia RM	(394860977)	Cachari E	0.50	Tatahouine D	0.17	Line	0.33	0.260	0.987
Aelia SM	(394860977)	Cachari E	0.09	GRO95555 D	0.67	Line	0.24	0.215	0.993
Bellicia CWM	(393308781)	Cachari E	0.35	Tatahouine D	0.16	Green Ol	0.49	0.490	0.981
Calpurnia CWM	(395166561)	Bouvante E	0.26	Tatahouine D	0.39	Jackson cty Ol	0.35	0.270	0.987
Canuleia CWM	(394198821)	Cachari E	0.45	Tatahouine D	0.39	Line	0.16	0.459	0.983
Canuleia RM	(394198821)	Cachari E	0.44	GRO95555 D	0.46	Line	0.10	0.539	0.987
Canuleia SM	(394198821)	Bouvante E	0.28	Serra de Mage E	0.25	GRO95555 D	0.47	0.632	0.984
Canuleia RM	(394198127)	Cachari E	0.35	GRO95555 D	0.41	Line	0.24	0.353	0.988
Canuleia RM	(371905283)	Cachari E	0.60	GRO95555 D	0.31	Line	0.09	0.577	0.985
Cornelia CWM	(371813008)	Bouvante E	0.28	GRO95555 D	0.58	Jackson cty Ol	0.14	0.381	0.989
Cornelia CWM	(394683501)	Cachari E	0.36	GRO95555 D	0.47	Line	0.17	0.157	0.995
Cornelia SM	(394683501)	Cachari E	0.35	GRO95555 D	0.45	Line	0.20	0.145	0.995
Cornelia CWM	(373313780)	Bouvante E	0.31	GRO95555 D	0.57	Jackson cty Ol	0.12	0.485	0.986
Cornelia SM	(373313780)	Bouvante E	0.25	GRO95555 D	0.59	Jackson cty Ol	0.16	0.315	0.991
Drusilla CWM	(393931377)	Bouvante E	0.28	GRO95555 D	0.55	Jackson cty Ol	0.17	0.420	0.987
Drusilla SM	(393931377)	Bouvante E	0.37	Tatahouine D	0.46	Jackson cty Ol	0.17	0.368	0.987
Eumachia CWM	(394859589)	Cachari E	0.57	Tatahouine D	0.14	Line	0.29	0.342	0.985
Eumachia SM	(394859589)	Cachari E	0.54	Tatahouine D	0.17	Line	0.29	0.328	0.985
Justina CWM	(372520864)	Cachari E	0.54	Tatahouine D	0.28	Line	0.18	0.296	0.991
Justina RM	(372520864)	Cachari E	0.38	GRO95555 D	0.43	Line	0.19	0.264	0.989
Justina SM	(372520864)	Cachari E	0.42	GRO95555 D	0.35	Line	0.23	0.305	0.989
Justina CWM	(371904187)	Cachari E	0.58	Tatahouine D	0.27	Line	0.15	0.370	0.983
Justina RM	(371904187)	Cachari E	0.41	GRO95555 D	0.44	Line	0.15	0.323	0.992
Justina SM	(371904187)	Cachari E	0.54	Tatahouine D	0.29	Line	0.17	0.335	0.989
Lepida SM	(395256929)	Cachari E	0.45	ALHA77256 D	0.30	Line	0.25	0.595	0.977
Licinia CWM	(395387692)	Bouvante E	0.18	GRO95555 D	0.63	Jackson cty Ol	0.19	0.273	0.992
Licinia SM	(395387692)	Cachari E	0.30	GRO95555 D	0.53	Line	0.17	0.192	0.994
Marcia CWM	(395301166)	Cachari E	0.53	GRO95555 D	0.23	Line	0.24	0.410	0.984
Marcia SM	(395301166)	Cachari E	0.46	GRO95555 D	0.31	Line	0.23	0.396	0.986
Marcia CWM	(395301549)	Cachari E	0.48	GRO95555 D	0.32	Line	0.20	0.301	0.990
Marcia SM	(395301549)	Cachari E	0.48	GRO95555 D	0.30	Line	0.22	0.353	0.987
Myia RM	(372300571)	Cachari E	0.38	GRO95555 D	0.41	Line	0.21	0.396	0.987
Oppia CWM	(373136361)	Bouvante E	0.38	Tatahouine D	0.43	Jackson cty Ol	0.19	0.364	0.986
Oppia SM	(373136361)	Cachari E	0.45	GRO95555 D	0.27	Line	0.28	0.309	0.979
Pinaria CWM	(371727131)	Cachari E	0.41	ALHA77256 D	0.46	Line	0.13	0.332	0.990
Pinaria SM	(371727131)	Bouvante E	0.27	GRO95555 D	0.54	Jackson cty Ol	0.19	0.308	0.990
Sextilia CWM	(371816828)	Cachari E	0.48	GRO95555 D	0.35	Line	0.17	0.461	0.986
Sextilia SM	(371816828)	Cachari E	0.48	GRO95555 D	0.36	Line	0.16	0.484	0.986
Tuccia RM	(372565807)	Cachari E	0.59	Tatahouine D	0.27	Line	0.14	0.319	0.990

Table 2