Geochemical portrayal of the Pacific Ridge: New isotopic data and statistical techniques

Cédric Hamelina, Laure Dossoa, Barry B. Hananc, Manuel Moreirad, Andrew P. Kositskye and Marion Y. Thomasa

a I.U.E.M., U.B.O., Place Nicolas Copernic, 29280 Plouzané, France
b Centre National de la Recherche Scientifique, UMR 6538, IFREMER, BP70, 29280 Plouzané, France
c Department of Geological Sciences, S.D.S.U., 5500 Campanile Drive, San Diego, CA 92182-1020, USA
d Institut de Physique du Globe de Paris, CNRS UMR 7154, 1 rue Jussieu, 75252 Paris CEDEX 05, France
e Tectonics Observatory, California Institute of Technology, Pasadena, CA 91125, USA

* Corresponding author : Cédric Hamelin, I.P.G.P, 1 rue Jussieu, Bureau 345, 75252 Paris CEDEX 05, France. Tel.: +33 1 83 95 76 72; fax: +33 1 7193 7710, email address : hamelin@ipgp.fr

Abstract:

Samples collected during the PACANTARCTIC 2 cruise fill a sampling gap from 53° to 41° S along the Pacific Antarctic Ridge (PAR). Analysis of Sr, Nd, Pb, Hf, and He isotope compositions of these new samples is shown together with published data from 66°S to 53°S and from the EPR. The recent advance in analytical mass spectrometry techniques generates a spectacular increase in the number of multidimensional isotopic data for oceanic basalts. Working with such multidimensional datasets generates a new approach for the data interpretation, preferably based on statistical analysis techniques.

Principal Component Analysis (PCA) is a powerful mathematical tool to study this type of datasets. The purpose of PCA is to reduce the number of dimensions by keeping only those characteristics that contribute most to its variance. Using this technique, it becomes possible to have a statistical picture of the geochemical variations along the entire Pacific Ridge from 70°S to 10°S. The incomplete sampling of the ridge led previously to the identification of a large-scale division of the south Pacific mantle at the latitude of Easter Island. The PCA method applied here to the completed dataset reveals a different geochemical profile. Along the Pacific Ridge, a large-scale bell-shaped variation with an extremum at about 38°S of latitude is interpreted as a progressive change in the geochemical characteristics of the depleted matrix of the mantle. This Pacific Isotopic Bump (PIB) is also noticeable in the He isotopic ratio along-axis variation. The linear correlation observed between He and heavy radiogenic isotopes, together with the result of the PCA calculation, suggests that the large-scale variation is unrelated to the plume–ridge interactions in the area and should rather be attributed to the partial melting of a marble-cake assemblage.

Research Highlights

► New Sr, Nd, Pb, Hf, and He isotopes data fill a sampling gap along the Pacific Ridge. ► We examine geochemical variation in MORB using a principal component analysis. ► A progressive...
change in the depleted matrix is recognized along the Pacific ridge. In samples devoid of plume influence, He isotopes correlates with Pb isotopes.

Keywords: oceanic basalts; Pacific–Antarctic Ridge; mantle heterogeneity; Principal Component Analysis; Sr Nd Pb Hf isotopes
1. Introduction

Mid-oceanic ridge basalts (MORB) are the result of continuous melting of the ambient upper mantle beneath oceanic ridges. The MORB-source mantle is generally thought to have been depleted ~2 Gy ago by extraction of the continents. Although the range of geochemical variations in oceanic basalts is mostly attributed to the influence of Ocean Island Basalt (OIB), significant geochemical heterogeneity in MORB has been recognized (Hoffman et al., 2003 and reference therein; Rudge et al., 2005). This heterogeneity has been revealed using radiogenic isotope ratios (Sr, Nd, Pb and Hf) in areas devoid of plume influence. Numerous studies have attempted to model the mixing relationship between enriched and depleted domains within the mantle to reproduce the range of isotopic variations observed in oceanic basalts (Albarède, 2001; Meibom and Anderson, 2004; Rudge et al., 2005; Kellogg et al., 2007). The range of Sr, Nd and Pb isotopic compositions depends not only on the end-member compositions but also on the volume of mantle sampled during the melting relative to the length scale of heterogeneities (Kellogg et al., 2007). Therefore, the range of MORB geochemical variations reflects the size, the spatial distribution, and the difference of fusibility of heterogeneities within the upper mantle. In addition to this intrinsic heterogeneity of the MORB mantle, radiogenic isotope studies have led to the definition of distinct broad mantle isotopic domains, such as the archetypal DUPAL anomaly located in the south hemisphere (Dupré and Allègre, 1983; Castillo, 1988). Boundaries between these domains can be (i) extremely sharp as seen at the Antarctic Australian–Antarctic Discordance (ADD), with isotopic ratios (Sr, Nd, Pb and Hf) abruptly changing from Indian to Pacific values (Pyle et al., 1992, Hanan et al., 2004; Meyzen et al., 2007; Cooper et al., 2009) or (ii) more gradual as seen in the transition from Southwest Indian Ridge to South Atlantic MORB (Meyzen et al, 2005).
Another example of these large geochemical provinces is given by the two sub-pacific mantle domains (Vlastélic et al., 1999). The distinctive isotopic properties of these large-scale geochemical domains suggest a long-term isolation of these mantle provinces, each with their own convective histories involving various amounts of melting residues and recycled components. On the basis of an incomplete dataset, a boundary between these two pacific provinces has been located at the latitude of the Easter Island microplate (Vlastélic et al., 1999). But in order to have a complete geochemical view of the southern Pacific Ridge from 66 to 10°S, it was necessary to fill the sampling gap between 53°S and 41°S. This became one of the main objectives of the PACANTARCTIC2 cruise which took place in 2004-05 (Dosso et al. 2005, Klingelhoefer et al., 2006, Moreira et al., 2008; Hamelin et al., 2010). Analyses of Sr, Nd, Pb, Hf and He isotopic compositions of these new samples from the Pacific Antarctic Ridge (PAR) are compiled here together with published data from 66°S to 53°S and from the East Pacific Rise (EPR). Therefore it becomes possible to have a picture of the geochemical variations along the entire Pacific Ridge from 66°S to the 10°S. Adapting a Principal Component Analysis (PCA) to incomplete datasets, we show a detailed portray of the geochemical variability of the Pacific Ridge.

2. Data selection and analytical methods.

2.1. The Pacific Ridge database from 10 to 66°S.

Along the Pacific Ridge, the Chile Triple Junction at 35°S/110°W separates the PAR from the EPR. The two pacific ridges show different geological settings: the full spreading rate increases along the PAR from 54 mm/yr at 65°S to 100 mm/yr at 35°S whereas it decreases along the EPR from 158 mm/yr at 35°S to 146 mm/yr at 10°S. In conjunction, the ridge axis morphology changes from a valley to a dome north of 60°S along the PAR (Ondrèas et al.,
2001), whereas the EPR is characterized by a uniform dome shaped morphology. Three
plume-ridge interactions generate abnormal morphological structures along this section:
Foundation (~38°S), Easter Island (~26°S) and 17°S. They are recognized by a high ridge
cross-sectional area (Klingelhöfer et al., 2006) and the presence of intense off-axis volcanic
activity. Compared to the EPR, the Pacific Antarctic Ridge still remains geochemically poorly
known. A large portion of this plate boundary has been previously surveyed (Lonsdale, 1994;
Cande et al., 1995) but the northern part has only recently been sampled (Dosso et al., 2005;
Moreira et al., 2008; Hamelin et al., 2010).

To generate a coherent database along the pacific ridges, on-axis samples analyzed for one
or more isotopic ratios (Sr-Nd-Pb-Hf-He) are selected from 66°S to 10°S using our new
isotope data (table 1) completed with the petrological database of the Lamont-Doherty Earth
Observatory (http://www.petdb.org). In most cases when the reference values of the standards
are reported, the data are normalized to the values of NBS987, 0.71025 for Sr, of JNd1-1,
0.512104 and La Jolla, 0.511852 for Nd, NBS981, $^{206}$Pb/$^{204}$Pb = 16.9373, $^{207}$Pb/$^{204}$Pb = 15.4925,
$^{208}$Pb/$^{204}$Pb = 36.7054, for Pb and JMC475, 0.282162 for Hf (see http://georem.mpch-
mainz.gwdg.de/ for details and references):.

2.2. Analytical methods.

2.2.1. Double Spike Pb analyses along the PAR

New high-resolution Pb analyses were carried out for samples collected during
PACANTARCTIC1 (PAC1) and PACANTARCTIC2 (PAC2) cruises along PAR segments
from 66°S to 56°S and from 53°S to 41°S respectively (Fig. 1, Table 1). Small chips from the
inner part of the pillow lavas were handpicked to avoid altered surfaces that could be a
potential source of Pb contamination. Powdered samples were leached with 6M HCl at 140°C
for an hour and then rinsed up to 6 times with ultrapure water prior to dissolution. Lead separation was then performed on an anionic exchange resin. Pb analyses were performed at Ifremer (Centre de Brest) on a Finnigan MAT 26x multicollector instrument (MAT261 upgraded by Spectromat), using the double spike technique with the calibrated Southampton-Brest 207/204 spike (Ishizuka et al., 2003). Replicate analyses of the Pb isotope standard NBS981 gave an average of 16.9432±0.0027 and 15.5004±0.0029 and 36.7326±0.0086 for 206Pb/204Pb, 207Pb/204Pb and 208Pb/204Pb, respectively. Pb blanks measured using this procedure were < 100 pg, and thus negligible relative to the amount of sample analyzed.

2.2.2. Hf measurements along the PAR

Hafnium isotopic compositions were analyzed along the PAC1 ridge segments at SDSU on splits from the same samples (Hamelin et al., 2010). Hf was separated using the protocol of Blichert-Toft et al. 1997 with a negligible blank of less than 25 pg. Hf isotope ratios were measured at SDSU using the Nu Plasma. The $^{176}$Hf/$^{177}$Hf was normalized for mass fractionation relative to $^{179}$Hf/$^{177}$Hf=0.7325. The JMC-475 Hf standard $^{176}$Hf/$^{177}$Hf gave 0.282160 ±0.000010 (2σ) during this study. The standard was run alternately with samples to monitor machine performance.

2.2.3. He measurements along the PAR

Helium isotopic compositions have been measured on PAC1 and PAC2 samples at IPGP (Institut de Physique du Globe de Paris). Fresh pieces of glass were cleaned with distilled water, ethanol and acetone using an ultrasonic bath. Some samples were also cleaned with hydrogen peroxide in order to remove some Mg crust. Analytical procedure is identical to previous studies from our laboratory and can be found in Moreira et al. (1995). Samples were crushed under vacuum with analytical blanks of 7±1 nccSTP $^4$He. This corresponds to 0.02 to
0.4% of the samples. Helium concentrations and isotopic composition were measured using the ARESIBOII mass spectrometer (Moreira et al., 2008).

3. New results compared to published MORB data along this ridge section.

These results include samples from 66°S to 56°S (PAC1) and from 53°S to 41°S (PAC2) (Fig. 1). New high resolution Pb analyses as well as Hf and He analyses are presented here in Table 1. The Sr, Nd and Hf analyses on the same samples are found in Vlastélic et al. (2000) and Hamelin et al. (2010).

3.1. Binary isotopic correlations

With 7 isotopic ratios, the number of possible binary isotope diagrams is 21. As an example, we choose here to describe 4 such binary plots, selecting some of the most commonly discussed (Fig. 2A-D). In all diagrams, pacific MORB samples devoid of plume influence define linear correlations that in most cases have been previously described in the literature. This result is in good agreement with the expected coherence in behavior between Rb-Sr, U-Pb, Th-Pb, Sm-Nd and Lu-Hf isotopic systems compared to each other during magmatic processes. When plume-ridge interaction samples are included in our dataset, they define elongated fields overlapping PAR-EPR array except in Figure 2C where He-\(\varepsilon_{Nd}\) isotopic ratios define sub-parallel trends which emerge from the PAR-EPR array and point towards higher \(^{3}\text{He}/^{4}\text{He}\) (R/Ra) ratios. More than 30 years of He isotopes systematic of oceanic basalts worldwide have shown that MORB samples are characterized by a narrow
range of composition compared to OIB samples. However even within this limited range, our data show a linear correlation between $^3\text{He}/^4\text{He}$ (R/Ra) values and $\varepsilon_{\text{Nd}}$ (Fig. 2C). The correlation between He and other isotopic systems will be discussed further below (§4.4.2).

In contrast with the decoupling of Hf and Nd isotope compositions previously documented in MORB (e.g., Patchett and Tatsumoto, 1980; Salters and White, 1998; Chauvel et al., 2001; Debaille et al., 2006), a rather good correlation between these two isotopic systems is observed along our studied area (Fig. 2B). Debaille et al. (2006) have suggested that a distinctive behavior of Hf during disequilibrium melting along their studied ride area (Atlantic ridge 22-35°N) could explain a decoupled behavior of $^{176}\text{Hf}/^{177}\text{Hf}$ with respect to other isotopic ratios. The linear correlations observed in our dataset, show that the hypothesis of a specific behavior of Hf does not apply along the pacific ridges. Recently, good correlations between Hf and Nd isotopic ratios have been reported from other ridge segments such as Mohns Ridge (Blichert-Toft et al., 2005) and the entire mid-Atlantic Ridge (Agranier et al., 2005).

3.2. MORB variability along the south Pacific Ridge.

Latitudinal isotopic variations of south Pacific Ridge basalts are shown in Figure 3. An important geochemical feature from 66 to 10°S is a large scale (spanning approximately 5000km) and coherent variation of all isotopes shown by a bell shape grey line. The extremum of this variation defining the Pacific Isotopic Bump (PIB), is located at the latitude of Foundation (38°S). It reveals a less depleted component, characterized by more radiogenic values for Sr, Pb isotopic ratios, and correlative less radiogenic values for Nd and Hf. Three shorter wavelength variations of the order of 200 to 700km are seen as isotopic anomalies superimposed on the otherwise bell shaped curve of the isotope signature along the ridge. In a way similar to the Pacific Isotopic Bump, these variations are towards more radiogenic Sr and
Pb values and coherently less radiogenic Nd and Hf values. They indicate the influence of enriched materials due to plume-ridge interactions.

Helium isotopes do not fit this Sr-Pb/Hf-Nd coherent behavior. At the latitude of the Foundation-Ridge intersection (38°S), the bell-shaped He curve shows radiogenic $^3\text{He}/^4\text{He}$ ratios. But on the other hand, hot-spot influenced samples form negative “plume anomalies” with very unradiogenic signatures (high $^3\text{He}/^4\text{He}$ ratios, up to 12 R/Ra). The amplitudes of the PIB and the anomalies attributed to the plume effect are of the same order of magnitude for Nd, Hf and Pb. For He and Sr, the PIB has much smaller amplitude than the hotspot anomalies. Additionally, local MORB variability is expressed as spikes, which are likely related to transform faults (Eltanin System).

4. Discussion

4.1 Statistical definition of mantle reference lines

In most binary plots of radiogenic isotopic ratios, representative points of mantle derived material define linear trends. Historically, these trends have been used to define reference lines such as the Northern Hemisphere Reference Line (NHRL) in Pb-Pb plot (Hart, 1984) or the “Mantle Array” in Sr-Nd plot (DePaolo and Wasserburg, 1979) and the Nd-Hf plot (Vervoort and Blichert-Toft, 1999). These reference lines are arbitrary and depend on the abundance of data available at the time of their definition. However, the data colinearity in these plots justifies these convenient choices. The NHRL has been convenient to quantify the Dupal anomaly using the $\Delta 7/4$ and $\Delta 8/4$ (Hart, 1984). In 1999, new MORB data from the south Pacific lead to the definition of the Pacific Reference Line (PRL) which proved convenient to compare two sub-pacific mantle domains, using the $\delta$(Nd-Sr) and $\delta$(Sr-Pb) notations (Vlastélic et al., 1999). With the recent advance in analytical MC-ICPMS
techniques, the number of isotopic data increases dramatically. It becomes possible to look at
these reference lines in multidimensional space from a statistical point of view. Principal
Component Analysis (PCA) is a powerful mathematical tool to study data sets such as a
geochemical database including Sr-Nd-Pb-Hf analyses. The purpose of PCA analysis is to
reduce the number of dimensions in a data set by keeping those characteristics that contribute
most to its variance. This technique has been used in previous mantle heterogeneity studies
(e.g. Agranier et al., 2005, Debaille et al., 2006) The PCA method used with our geochemical
dataset has been initially developed for low-rank matrix approximations (Srebro and Jaakkola,
2003) and was recently adapted for tectonic problems using incomplete geodetic times series
(Kositsky and Avouac, 2010). The main difference with traditional PCA methods is that the
singular value decomposition is replaced by a more sophisticated decomposition, which
appropriately takes into account samples with a missing isotope measurement. This technique
is particularly suitable to geochemical data as it allows computing the principal components
using the whole dataset, increasing therefore the accuracy of the calculation (see
supplementary material for a more detailed discussion about the concept and the limits of our
calculation). Although more recently ICA (Independent Component Analysis) has been
chosen by some authors (Iwamori and Albarède, 2008, Iwamori et al., 2010), the preferred
PCA method used here has the advantage of dealing with missing isotopic data, assuming that
decorrelation is still a good assumption of independence in our dataset.

Because the PCA method is an orthogonal linear transformation, it assumes the linearity of
the data co-variations. In most binary isotopic diagrams, mixing processes are expressed by
hyperboles whose curvatures depend on the elemental concentration ratios of the involved
end-members. But in the case of MORB, pseudo-linear correlations are observed (Fig. 2)
indicating that denominator elements are in approximately constant proportions in the mixing
components. It is very difficult to evaluate the extent of non-linear relationships concealed
within the analytical noise. However, it is worth noting that the geographical variations of the components calculated with our method are approximately the same as those computed in the 3-dimensional space of Pb isotopes, in which relationships are linear. In order to minimize the correlations induced by the $^{204}$Pb analytical noise (it represents only about 1.4% of the total Pb), the computation has been made in the $^{204}$Pb/$^{206}$Pb, $^{207}$Pb/$^{206}$Pb and $^{208}$Pb/$^{206}$Pb space. Considering the problematic of our study, this observation suggests that the curvature can be neglected. One drawback of PCA in general stems from the fact that PCA is a projection method, and sometimes low-dimensional visualization can lead to erroneous interpretations.

4.2. Distribution of the variance among the principal components.

The most striking result of the PCA is that the first principal component accounts for more than 70% of the total variance. This result is remarkable considering the number of geochemical parameters involved in the very large number of samples used in this analysis. It indicates the strong coherence of all isotopic ratios within the depleted mantle domain. These correlations in binary plots of isotopic systems are the result of two antagonistic processes that took place over time: chemical fractionation events leading to the existence of enriched and depleted mantle reservoirs and mechanical mixing of these reservoirs during mantle convection. The chemical fractionation between parent and daughter isotopes is controlled by distribution coefficients. Even if these coefficients can be modified by multiple parameters, they remain consistent from one element to another. Since these processes are approximately linear, the resulting dispersion is located along a line corresponding to PC1 (the first Principal Component).

While the importance of the first component is indisputable, one challenge with PCA is to establish the number of significant components needed to explain the observed data variance. A classical way to illustrate the number of relevant components is to study the residual variance ($\chi^2$) obtained for each component (e.g. Kositsky and Avouac, 2010). This value
drops abruptly after the third component (see supplementary material). At first, we can infer that the information brought by the fourth to the sixth principal components is statistically insignificant and can safely be ignored. Using a PCA computed in the 3-dimensional space of Pb isotopes, the principal components account for 91.7%, 8.2% and 0.1% of the variance for PC1, PC2 and PC3 respectively. This observation suggests that the variance expressed by principal components over the order of 2 is the result of a randomly distributed error. Also the absence of coherent geographical variation for the third component suggests that this variance could either be a very local variation or an artifact of our data compilation (i.e. sampling bias, data and error normalization between different laboratories). This justifies to limit the discussion of the south pacific mantle heterogeneity to the PC1 and PC2 characteristics.

4.3. Mapping geochemical heterogeneities in the mantle using a PCA method.

Using the dataset available at the time, Vlastélic et al. (1999) identified a large geochemical variation in the south Pacific depleted mantle. Based on an incomplete sampling of the ridge, they suggested the existence of a sharp boundary located at the latitude of the Easter Island Microplate (27°S). As an interpretation, they proposed that the Pacific Superswell divided the mantle into two domains each with their own convective histories, producing slight differences in their average isotopic signatures. More recently, the origin of these two domains has been challenged by a model based on plate kinetics (Small and Danyushevsky, 2003). Small and Danyushevsky proposed that geochemical discontinuities result from variations of the asthenosphere consumption, which corresponds to the ratio between the accretion rate and the spreading center migration relative to plumes. Their model predicted that the fast spreading, slowly migrating East Pacific Rise should have higher average melting degrees compared to the slower spreading, rapidly migrating Pacific-Antarctic Rise. In order to identify the two mantle domains along the Pacific Ridge, Vlastélic
et al. (1999) used δ(Sr-Pb) and δ(Nd-Sr), which are defined as the vertical deviations from
references lines in Pb vs Sr and Sr vs Nd isotopic ratio plots respectively. These reference
lines (see §4.1) were drawn intuitively in the greatest variance of their dataset. They can be
directly compared to our first principal component. By construction, whatever the number of
dimensions considered in the PCA, the projection of PC1 in a binary diagram resembles the
relevant reference line (Fig. 2). Because PC2 is orthogonal to the greatest variance, sample
values along this component are correlated to delta notations such as δ(Sr-Pb), δ(Nd-Sr) or
Δ8/4Pb. But compared to delta notations, PC2 has the advantage of being rigorously and
statistically determined in a multidimensional isotopic space. The application of the PCA
method to our data compilation (excluding He isotopes) reveals a geochemical profile of the
Pacific Ridge. The plot of PC1 versus latitude summarizes all the characteristics noted with
the different isotopic systems (Fig. 4). The short scale geochemical variations associated with
hotspots as well as the large-scale variation are clearly visible. 17°S and 25°S (Easter Island)
are well defined by sharp anomalies superimposed on the bell shaped curve. A plot of PC2
versus latitude shows a very different picture (Fig. 4): only the large scale variation is
expressed by this component, the geochemical variations associated with hotspots are
flattened and the corresponding samples are projected along the bell shape curve defining the
Pacific Isotopic Bump. This PIB is also shown on plots of delta versus latitude, illustrating the
equivalence of these parameters and PC2 (Fig. 4). At the Juan Fernandez/Foundation latitude
(36°S), the isotopic variation curve reaches an extreme which corresponds to a less “depleted”
isotopic signature. In contrast with the conclusion of Vlastélic et al. (1999), we proposed that
tgeochemical variations along the Pacific Ridge are not the result of two separated mantle
domains but should rather be seen as a progressive variation of the isotopic composition of the
sub-Pacific depleted mantle. Since the asthenosphere consumption varies abruptly at the Chile
triple junction and is almost constant along the PAR, the plate kinematic model of Small and
Danyushevsky (2003) is also inconsistent with the observed progressive variation of the MORB depleted matrix.

4.4. Mixing relationship within the depleted mantle compared to ridge-hotspot interactions.

4.4.1. Asthenospheric versus hotspots signals as illustrated by Sr-Nd-Pb-Hf.

Except in figures involving He isotopes, geochemical variations related to ridge/hotspot interactions are consistent with variations of samples devoid of plume influence. At first, it is difficult to distinguish the two types of variations in binary isotopic plots (Fig. 2). Nevertheless, a PCA computed with all heavy radiogenic isotopes clearly illustrate a difference: hotspot signatures are exclusively visible with PC1, while the variance related to the large scale variation (PIB) is illustrated by both PC1 and PC2 versus latitude (Fig. 4). In PC1 versus PC2 space, the MORB field extends from the depleted end member of the mantle (DMM) toward a recycled oceanic crust end member with a HIMU affinity (Fig. 5). The samples identified as resulting from a plume-ridge interaction are not part of this “depleted trend”. Rather their data field extends from the depleted trend towards more enriched-type end members such as C and/or EM. This observation supports the idea that the large scale variation in the depleted Pacific mantle is unrelated to ridge/hotspot interactions. This variation is therefore equivalent to the intrinsic variability of MORB recognized in other areas (e.g. Dosso et al., 1999, Donnelly et al., 2004; Debaille et al., 2006; Hémond et al., 2006). It has to be noted that PCA calculations assume a linearity criteria which is not satisfied when dealing with isotopic compositions of the mantle end-members. It is thus not possible to determine more precisely the nature of the different mantle components responsible for the observed trends. Nevertheless, the PCA calculations establish that the progressive
geochemical change of the depleted matrix of the Pacific mantle is not the result of hotspot material being diluted into the depleted mantle. Using a different approach, Meyzen et al. (2007) successfully unscrambled the hotspot and asthenospheric signals. These authors have proposed that the geochemical variations unrelated to ridge/hotspot interactions along the South West Indian ridge and the South Atlantic ridge are related to a broad lower mantle upwelling in this area. Similarly, a broad lower mantle input could be a plausible cause of the Pacific Isotopic Bump associated with the high spreading rates in the vicinity of the Chile Ridge Triple junction.

It is interesting to compare the results of our PCA calculation along the Pacific ridge with the spectral analysis performed along the Atlantic ridge by Agranier et al. (2005). These authors found two contrasting types of spectra along their study area. The first type is associated with ridge-hotspot interactions and is seen in the first principal component. The second type is illustrated by the continuous power decrease with the decreasing wavelength of the second principal component. Agranier et al. (2005) have interpreted this second type of spectra, unrelated to hotspots, as being the result of the continuous size reduction of mantle heterogeneities upon stretching and refolding of the convecting mantle. Despite the differences in geological settings between the Pacific ridge and the Atlantic ridge, our observations are in good agreement with the statistical analysis of Agranier et al. (2005). We view the PIB as the consequence of a progressive change in the relative proportions of the marble-cake components present in the Pacific upper mantle. In this model, hotspot anomalies are superimposed on the intrinsic mantle heterogeneity expressed in MORB.

4.4.2. He isotopes in the depleted mantle compared to hotspot signals.

Unlike other isotopic systems, helium shows a first order discrepancy between the hotspot and the depleted mantle signals in our studied area (Fig. 3 and Fig. 6): the large scale variation
associated to the depleted mantle is characterized by an increase towards more radiogenic (low $^3\text{He}/^4\text{He}$) compositions, whereas the short scale variation associated to the mantle plumes points towards high $^3\text{He}/^4\text{He}$ values as previously noted in Hanan and Graham (1996). The origin of the large range of elevated $^3\text{He}/^4\text{He}$ values in OIB lavas is a long-lived controversial question (Kurz et al., 1982; Allègre et al., 1983; Meibom et al., 2003; Moreira et al., 2004). Historically, the requirement of a high $^3\text{He}/^4\text{He}$ reservoir for the OIB lead to the idea that plumes are tapping a deep, undegassed lower mantle, isolated from the upper mantle convection (O’Nions et al., 1996; Allègre et al., 1997). Numerous alternative models have tried to solve the apparent inconsistency between the high $^3\text{He}/^4\text{He}$ ratio in plumes and the requirement of a source previously processed through partial melting (e.g. Parman et al., 2005; Class and Goldstein, 2005; Purtika, 2008; Albarède, 2008; Davies, 2010).

Compared to plume-influenced samples, systematics of He isotopes in depleted MORB samples have shown a very restricted range of $^3\text{He}/^4\text{He}$ with a peak of distribution around $8\pm1$ (R/Ra) (Allègre et al., 1995). Within the uncontaminated depleted mantle sampled along our studied area, $^3\text{He}/^4\text{He}$ varies from 6 to 9.5 (R/Ra). Samples devoid of plume influence fall along a negative trend toward low $^3\text{He}/^4\text{He}$ values and high $^{206}\text{Pb}/^{204}\text{Pb}$. This variation of He isotopes in our samples devoid of plume influence is clearly related to the PIB identified along the pacific ridge using PCA with Sr, Nd, Pb and Hf isotopes. It is interesting to note that the most enriched samples from this correlation show the lowest $^3\text{He}/^4\text{He}$ values (more radiogenic). In order to reconcile the $^3\text{He}/^4\text{He}$ variations with variations of Sr, Pb, Nd and Hf isotopes, new models take into account the physical specificities of He compared to heavy radiogenic isotopes: its diffusivity in mantle conditions and its capacity to be outgassed from melts at sub-surface pressure (e.g. Hart et al., 2008; Albarède et al., 2008; Gonnermann and Mukhopadhyay, 2009). We propose that the melting of a marble-cake upper mantle, unpolluted by plumes, produces the good correlation observed between He and Pb isotopes in
our depleted samples. The less depleted component of the mantle assemblage is characterized by a high Pb isotopic ratio and a low $^3$He/$^4$He ratio. We suggest that this component was derived from the recycling of an extensively outgassed oceanic crust. When the oceanic crust is subducted back into the mantle, it contains negligible concentration of mantle-derived He (Staudacher and Allègre, 1988). With time, this very low $^3$He/(U+Th) in subducted slabs is expected to produce the end-member with high Pb isotopes value and low $^3$He/$^4$He ratio (Fig. 6).

In our model, the variation of He isotopes along the pacific ridge could be interpreted as: (1) a smaller contribution of the refractory layers (less radiogenic), corresponding to a lower $^3$He/$^4$He ratio at the PIB but because the melting rate is expected to be higher near the Chile Ridge Triple Junction, this hypothesis seems very unlikely, or (2) a higher volume of the recycled end-member in this area. This hypothesis is in agreement with the conclusions derived from the PCA based on heavy radiogenic isotopes (§4.4.1). The intrinsic geochemical variation of the depleted upper mantle comforts the image of a marble cake mantle composed of a refractory component and a recycled oceanic crust component. A broad volume of recycled component associated with the high spreading rates in the vicinity of the Chile Ridge Triple junction is therefore a plausible cause of the Pacific Isotopic Bump. The correlation between He and Pb isotopes revealed by our new data (Fig. 6) suggests that the depleted mantle is the outcome of a mixing of a different nature that the one involved in the plume-ridge interaction.
5. Conclusion

Analyses of this new sampling of the PAR complete the dataset of the Pacific Ridge. Therefore it becomes possible to have a picture of the geochemical variations from 10 to 70°S. Our data show a clear geographical evolution of isotopic characteristics along the Pacific Ridge. In binary isotopic plots, this large-scale variation is expressed by correlations between each isotopic dimension. This observation holds true even for the Nd-Hf system previously reported as decoupled for MORB samples (Debaille et al., 2006). Even more significant in this study is the linear correlation shown in the isotopic He-Nd and $^{3}$He/$^{4}$He - $^{206}$Pb/$^{204}$Pb plots. In these isotopic spaces, the samples affected by plume-ridge interactions depart clearly from the linear correlations displayed by the ridge samples coming from the depleted mantle.

Despite the paucity of combined Sr-Nd-Pb-Hf isotope data on individual samples, the PCA algorithms used here allow us to portray the variation along a huge section of the Pacific Ridge from a statistical point of view. Compared to previous studies of the sub-Pacific mantle, the application of PCA reveals a Pacific Isotopic Bump, which can be seen as a progressive geochemical variation of the depleted upper-mantle matrix rather than a sharp frontier between two mantle domains. Combining PCA results with the information given by He isotopes, we suggest that the Pacific Isotopic Bump is unrelated to plume-ridge interactions. This geochemical variation in the upper mantle reservoir is the result of a marble-cake mantle assemblage composed of a residual mantle component and a recycled oceanic crust component.
Acknowledgements

The Pb isotope work was funded by CNRS/INSU. The Hf isotope work was supported by NSF grants to B.B. Hanan. We thank Joan Miller for technical assistance at SDSU. We acknowledge Rick Carlson for the editorial handling, Francis Albarède and an anonymous reviewer for constructive comments. Cedric Hamelin thanks the Caltech Tectonic Observatory for its hospitality during his stay in Pasadena. We gratefully acknowledge Neus Sabater for her valuable comments.
References


Figure captions

**Figure 1:** Map of the south pacific basin showing the location of the PACANTARCTIC1 and PACANTARCTIC2 cruises with respect to the Pacific Antarctic Ridge (PAR) and the East Pacific Rise (EPR). Hotspot locations are shown in orange on the map.

**Figure 2:** Examples of binary plots showing correlations within the MORB field. Samples devoid of plume influence are shown in blue whereas samples from plume-ridge interactions are in red. Blue circles represent samples collected along the PAR and blue crosses for EPR samples (all symbols are kept the same in all figures). Mantle reference lines defined in the literature are shown in dashed lines: in the Sr/Nd plot (A), the Pacific Reference Line (PRL) (Vlastélic et al., 1999), in the Nd/Hf plot (B), the mantle array (Vervoort et al., 1999), in the Pb/Pb plot (D), the North Hemisphere Reference Line (NHRL) (Hart, 1984). Since no reference line had yet been recognized in plots involving He isotopes, we show here a regression line ($^3\text{He}/^4\text{He} = 0.63 \varepsilon_{\text{Nd}} + 1.74, r = 0.89$) (C). The projection of the two first principal components (PC1 and PC2) calculated in our study are shown (solid black and grey lines, see text for additional information). Data references can be found in supplementary materials.

**Figure 3:** Geochemical variations along the PAR and the EPR from 66°S to 10°S. Dashed lines represent the location of major transform faults in the area and a grey shade is used for the Juan Fernandez and Easter Island microplates. The grey curve underlines the large-scale isotopic variation along the pacific ridges from 10 to 66°S and defines the Pacific Isotopic Bump (PIB).
Figure 4: Plot of the two first principal components along the Pacific Ridge. The similarity between PC2 and δ notation is illustrated by plotting δ(Sr-Pb) as defined by Vlastélic et al. (1999) and Δ²⁰⁷Pb as defined by Hart (1984) versus latitude.

Figure 5: Plot of PC2 versus PC1 for a principal component analysis computed using Sr, Nd, Hf and Pb isotopes. The hashed blue field defines the depleted trend and includes EPR and PAR samples. Data points are drawn as ellipses representing the 95% confidence domain of the components as calculated in Debaille et al. (2006). The insert shows the location of the classical mantle end-members in the PC2 vs PC1 space.

Figure 6: Binary plot of $^{3}\text{He}/^{4}\text{He}$ vs $^{206}\text{Pb}/^{204}\text{Pb}$. The dashed regression line ($^{3}\text{He}/^{4}\text{He} = -2.33^{206}\text{Pb}/^{204}\text{Pb} + 51.0$) is defined by samples devoid of plume influence. Plume-ridge interactions (17°S and Easter hotspots) are characterized by positive trends emerging from the PAR-EPR array and pointing towards higher $^{3}\text{He}/^{4}\text{He}$ (R/Ra) ratios and higher $^{206}\text{Pb}/^{204}\text{Pb}$.

Table 1: Complete Sr, Nd, Pb, Hf and He isotope data table for PACANTARCTIC 1 and 2 samples. In italic font, previously published data (Vlastélic et al., 1999; Vlastélic et al., 2000; Moreira et al., 2008; Hamelin et al., 2010).
He/He (R/Ra)

Depleted trend

17°S Easter
Table 1: Complete Sr, Nd, Pb, Hf and He isotope data for PACANTARCTIC 1 and 2 samples. In italic font, previously published data (Vlastélic et al., 1999; Vlastélic et al., 2000; Moreira et al., 2008; Hamelin et al., 2010).

<table>
<thead>
<tr>
<th>PACANTARCTIC 1</th>
<th>PACANTARCTIC 2</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Sr/Sr</strong></td>
<td><strong>Sr/Sr</strong></td>
</tr>
<tr>
<td><strong>Nd/Nd</strong></td>
<td><strong>Nd/Nd</strong></td>
</tr>
<tr>
<td><strong>He/He</strong></td>
<td><strong>He/He</strong></td>
</tr>
<tr>
<td><strong>Long. (°)</strong></td>
<td><strong>Long. (°)</strong></td>
</tr>
<tr>
<td><strong>Lat. (°)</strong></td>
<td><strong>Lat. (°)</strong></td>
</tr>
<tr>
<td><strong>Depth</strong></td>
<td><strong>Depth</strong></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Sample</th>
<th>Sr/Sr</th>
<th>Nd/Nd</th>
<th>He/He</th>
<th>Long.</th>
<th>Lat.</th>
<th>Depth</th>
</tr>
</thead>
<tbody>
<tr>
<td>PAC1DR01-2</td>
<td>0.702358 0.513138</td>
<td>0.75</td>
<td>18.404</td>
<td>15.489</td>
<td>16.02</td>
<td>7.6</td>
</tr>
<tr>
<td>PAC1DR03-1</td>
<td>0.702530 0.513102</td>
<td>9.05</td>
<td>18.776</td>
<td>15.524</td>
<td>38.379</td>
<td>0.283214</td>
</tr>
<tr>
<td>PAC1DR04-4</td>
<td>0.702557 0.513116</td>
<td>9.62</td>
<td>18.769</td>
<td>15.523</td>
<td>38.370</td>
<td>0.283218</td>
</tr>
<tr>
<td>PAC1DR13-2g</td>
<td>0.702556 0.513142</td>
<td>9.82</td>
<td>18.508</td>
<td>15.503</td>
<td>37.954</td>
<td>0.283204</td>
</tr>
<tr>
<td>PAC1DR15-3</td>
<td>0.702570 0.513145</td>
<td>9.89</td>
<td>18.508</td>
<td>15.503</td>
<td>37.963</td>
<td>0.283200</td>
</tr>
<tr>
<td>PAC1DR15-1g</td>
<td>0.702510 0.513150</td>
<td>9.99</td>
<td>18.604</td>
<td>15.467</td>
<td>37.468</td>
<td>0.283235</td>
</tr>
<tr>
<td>PAC1DR12-3g</td>
<td>0.702375 0.513146</td>
<td>9.91</td>
<td>18.194</td>
<td>15.473</td>
<td>37.627</td>
<td>0.283217</td>
</tr>
<tr>
<td>PAC1DR12-3r</td>
<td>0.702395 0.513170</td>
<td>10.38</td>
<td>18.192</td>
<td>15.470</td>
<td>37.620</td>
<td>0.283216</td>
</tr>
<tr>
<td>PAC1DR11-1g</td>
<td>0.702435 0.513125</td>
<td>9.50</td>
<td>18.420</td>
<td>15.491</td>
<td>37.877</td>
<td>0.283171</td>
</tr>
<tr>
<td>PAC1DR11-3</td>
<td>0.702469 0.513129</td>
<td>9.58</td>
<td>18.397</td>
<td>15.491</td>
<td>37.816</td>
<td>0.283182</td>
</tr>
<tr>
<td>PAC1DR10-1g</td>
<td>0.702473 0.513112</td>
<td>9.25</td>
<td>18.426</td>
<td>15.492</td>
<td>37.881</td>
<td>0.283149</td>
</tr>
<tr>
<td>PAC1DR09</td>
<td>-117.5</td>
<td>49.9</td>
<td>-117.2</td>
<td>-50.7</td>
<td>-117.2</td>
<td>-50.7</td>
</tr>
<tr>
<td>PAC1DR08</td>
<td>-112.7</td>
<td>-45.85</td>
<td>-112.3</td>
<td>-47.51</td>
<td>-113.3</td>
<td>-47.51</td>
</tr>
<tr>
<td>PAC1DR07</td>
<td>-111.8</td>
<td>-43.59</td>
<td>-111.3</td>
<td>-42.27</td>
<td>-111.3</td>
<td>-42.27</td>
</tr>
<tr>
<td>PAC1DR06</td>
<td>-110.8</td>
<td>-42.27</td>
<td>-110.3</td>
<td>-41.80</td>
<td>-110.3</td>
<td>-41.80</td>
</tr>
<tr>
<td>PAC1DR05</td>
<td>-110.0</td>
<td>-42.27</td>
<td>-110.0</td>
<td>-42.27</td>
<td>-110.0</td>
<td>-42.27</td>
</tr>
<tr>
<td>PAC1DR04</td>
<td>-109.2</td>
<td>-42.27</td>
<td>-109.2</td>
<td>-42.27</td>
<td>-109.2</td>
<td>-42.27</td>
</tr>
<tr>
<td>PAC1DR03</td>
<td>-108.4</td>
<td>-42.27</td>
<td>-108.4</td>
<td>-42.27</td>
<td>-108.4</td>
<td>-42.27</td>
</tr>
<tr>
<td>PAC1DR02</td>
<td>-107.6</td>
<td>-42.27</td>
<td>-107.6</td>
<td>-42.27</td>
<td>-107.6</td>
<td>-42.27</td>
</tr>
<tr>
<td>PAC1DR01</td>
<td>-106.8</td>
<td>-42.27</td>
<td>-106.8</td>
<td>-42.27</td>
<td>-106.8</td>
<td>-42.27</td>
</tr>
</tbody>
</table>
New algorithm to compute a PCA:

The PCA method is a linear transformation that converts the data into a new coordinate system in which the direction along which the greatest data variance is expressed, becomes the first axis (PC1). The direction along which the greatest data variance is expressed orthogonal to PC1 becomes the second axis (PC2), and so on. The first step, called the whitening, consists in normalizing the data. This is done by subtracting the mean value from each data point and dividing the result by the standard deviation of each variable. The correlation matrix (Pearson’s correlation matrix) of the reduced variables can then be calculated. The next step consists into diagonalizing the correlation matrix in order to find the eigenvalues, which correspond to the dimensions that have the strongest correlation in the data set. The principal components are the coordinates of the data points in the eigenvector referential.

The PCA method used with our geochemical dataset has been initially developed for low-rank matrix approximations (Srebro and Jaakkola, 2003) and was recently adapted for tectonic problems using incomplete geodetic times series (Kositsky and Avouac, 2010). The algorithm used to compute our principal components has been written specifically to address the major problem of a geochemical database: missing data in the matrix. Traditionally, for a given sample, it is necessary to acquire as many different isotope measurements as the number of dimensions used in the PCA. The paucity of combined Sr-Nd-Pb-Hf isotope data in individual samples is therefore an important issue for standard PCA algorithms. The more dimensions we use in the PCA, the fewer samples meet the required conditions: from 210 samples for 2-dimension (Sr-Nd) space to only 99 samples for a 6-dimension space. The main difference with a classical PCA is that we have replaced the standard Singular Value Decomposition (SVD) with a more sophisticated decomposition proposed by Srebro and Jaakkola [2003]. In this approach the data matrix X is decomposed into components U, S, V, with each data point weighted according to the square of its standard error. This decomposition is particularly adapted to take into account individual measurement errors and deal with missing data points. It allows us to compute the Principal Components on the entire data set, even if some isotopic ratios are missing. Each missing data point is weighted with infinite standard error. We want to emphasize that, consequently, no interpolation is made, which could erase some local effect. This technique is particularly suitable to geochemical data as it allows us to complete the principal components using the whole dataset, increasing therefore the accuracy of the calculation.


Limits of PCA:

Classical limits related to PCA calculation:

Because the PCA method is an orthogonal linear transformation, it assumes the linearity of the data co-variations. In most binary isotopic diagrams, mixing processes are expressed by hyperboles whose curvatures depend on the elemental concentration ratios of the involved end-members. But in the case of MORB, pseudo-linear correlations are observed (Fig. 2) indicating that denominator elements are in approximately constant proportions in the mixing components. It is very difficult to evaluate the extent of non-linear relationships concealed within the analytical noise. However, it is interesting to note that the geographical variations of the components calculated with our method are approximately the same as those computed in the 3-dimensional space of Pb isotopes, in which relationships are linear. In order to minimize the correlations induced by the $^{204}\text{Pb}$ analytical noise (it represents only about 1.4% of the total Pb), the computation has been made in the $^{204}\text{Pb}/^{206}\text{Pb}$, $^{207}\text{Pb}/^{206}\text{Pb}$ and $^{208}\text{Pb}/^{206}\text{Pb}$ space. Considering the problematic of our study, this observation suggests that the curvature can be neglected.

Limits related to our new algorithm:

Compared to traditional PCA algorithms, principal components are computed here altogether rather than separately, because the number of components affects the subspace in which these components reside. A limit to this method appears if one variable is represented by fewer samples than compared to other variables. In this particular case, a sample with a value corresponding to this “rare” variable is artificially given more weight. In return, each variable is also weighed in proportion to its number of samples. Because the number of He isotope analyses along the studied ridge section is too low compared to other isotopes, the information provided by this parameter has not been included in the PCA calculation.
Distribution of the variance among the principal components:

![Plot of the residual variance ($\chi^2$) versus the principal component number](image)

Figure 2 data references:


