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Edited by:

Nathalie Vigier,
UMR7093 Laboratoire
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Reviewed by:

Emmanuelle Albalat,
UMR5276 Laboratoire de géologie de
Lyon Terre, planètes et environnement
(LGL-TPE), France
João Canário,
University of Lisbon, Portugal

***Correspondence:**

F. Lacan
francois.lacan@legos.obs-mip.fr

[†]Present address:

Lise Artigue,
GET, CNRS, CNES, IRD, UPS,
University of Toulouse, Toulouse,
France

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Interferences and Matrix Effects on Iron Isotopic Composition Measurements by ⁵⁷Fe-⁵⁸Fe Double-Spike Multi-Collector Inductively Coupled Plasma Mass Spectrometry; the Importance of Calcium and Aluminum Interferences

F. Lacan^{1*}, L. Artigue^{1†}, J. K. Klar¹, C. Pradoux¹, J. Chmeleff² and R. Freydl³

¹LEGOS, CNRS, CNES, IRD, UPS, University of Toulouse, Toulouse, France, ²GET, CNRS, CNES, IRD, UPS, University of Toulouse, Toulouse, France, ³HydroSciences Montpellier, Université de Montpellier, CNRS, IRD, Montpellier, France

Multi-collector inductively coupled plasma mass spectrometers (MC-ICPMS) are widely used for Fe isotope measurements. The latter may be perturbed by interferences (notably from Cr and Ni) and matrix effects (notably from major elements), caused by elements remaining in the samples after purification. We quantified some of these perturbations and our ability to correct them whenever possible, using Thermo Neptune and Neptune Plus MC-ICPMS with a ⁵⁷Fe-⁵⁸Fe double-spike mass bias correction. ⁵⁴Cr and ⁵⁸Ni isobaric interference corrections were found to be extremely efficient up to Cr/Fe=0.12 and Ni/Fe=0.04 (g/g natural Fe). Matrix effects were found negligible up to at least Na/Fe=175, Mg/Fe=10, K/Fe=1.5, and Mo/Fe=75 (g/g natural Fe). ²⁸Si₂⁺ interference was found negligible up to Si/Fe=50. Finally, we found that calcium and aluminum could cause significant interferences (e.g., ⁴⁰Ca¹⁶O and ²⁷Al₂⁺), for Ca/Fe ≥ 2.5 and Al/Fe ≥ 2.5. The perturbation intensity relative to the Ca/Fe ratio was found dependent on the measurement conditions (plateau width). While working with samples with potentially high calcium or aluminum contents (such as calcite minerals or tests, bones and teeth, or marine samples and crustal rocks), we recommend to carefully take into account Ca and Al while tuning the instrument and checking the measurement accuracy with isotopic standards (i.e., doping the isotopic standard with Ca and Al levels comparable to those of the samples).

Keywords: iron isotopes, MC-ICPMS, double-spike, interferences, matrix effects, calcium, aluminum

INTRODUCTION

Iron isotopic compositions are nowadays widely used in Earth and planetary sciences, environmental research, and also in biological and medical research (Beard and Johnson, 2004; Abadie et al., 2017; Albarède et al., 2017; Dauphas et al., 2017). They are most commonly expressed as $\delta^{56}\text{Fe}$ defined as follows:

$$\delta^{56}\text{Fe} (\text{‰}) = \left[\frac{(^{56}\text{Fe}/^{54}\text{Fe})_{\text{sample}}}{(^{56}\text{Fe}/^{54}\text{Fe})_{\text{IRMM-14}}} - 1 \right] \times 10^3.$$

Almost all of these measurements are now carried out with multi-collector inductively coupled plasma mass spectrometers (MC-ICPMS). While these instruments allow very reliable and accurate measurements when operated in certain conditions, the measurements may suffer from a variety of perturbations, which include spectral interferences and matrix effects.

Samples are usually purified before MC-ICPMS analyses, in order to introduce pure Fe samples (dissolved in a dilute acid) in the instrument. However, even the highest performance purification procedure does not allow the entire elimination of other elements initially present in the sample. Some of these elements may cause spectral interferences and/or matrix effects. A spectral interference is the process by which an ion other than the analyte is detected as if it was the analyte because its mass-to-charge ratio is indistinguishable from that of the analyte. There are three main types of spectral interferences in ICPMS: isobaric interferences, produced by mono-atomic ions (e.g., $^{54}\text{Cr}^+$ and $^{58}\text{Ni}^+$ may interfere with $^{54}\text{Fe}^+$ and $^{58}\text{Fe}^+$, respectively), poly-atomic interference, produced by molecular ions (e.g., $^{14}\text{N}^{40}\text{Ar}^+$ or $^{27}\text{Al}_2^+$ may interfere with $^{54}\text{Fe}^+$, $^{16}\text{O}^{40}\text{Ca}^+$ may interfere with $^{56}\text{Fe}^+$), and doubly charged ion interferences (e.g., $^{112}\text{Cd}^{++}$ on $^{56}\text{Fe}^+$). A matrix effect is the process by which the instrument behavior varies in response to a variation in the sample composition. It is caused by dissolved ions in the solution (other than that of interest). It can be divided into two types: a signal drift with time due to salt deposition on the cones and/or a modification of the mass bias (instrumental mass fractionation) due to a modification of the ionization environment in the plasma (i.e., of the isotope distribution in the plasma; the so-called space charge effect; Gillson et al., 1988; Barling and Weis, 2012). For instance, the mass bias may suddenly vary between a very well-purified isotopic standard solution containing almost exclusively the analyte and an insufficiently purified sample solution containing significant levels of other compounds. Instrumental mass bias correction is a major aspect of the Fe isotopic ratio measurements because it is typically of the order of 2.5–4.5% per atomic mass unit (this study; Dauphas and Rouxel, 2006), which is about 200–400 times larger than the order of magnitude of the typical targeted accuracy of 0.1‰. Three main methods are commonly used for Fe mass bias corrections: standard bracketing; internal doping with another element of similar mass, usually Ni or Cu; and double-isotope spiking.

Some of the abovementioned perturbations, interferences, and matrix effects may be quantified and corrected for (e.g., Cr and Ni isobaric interferences), but corrections are never perfect. In addition, some perturbations cannot be quantified or corrected (e.g., matrix

effects). Therefore, these perturbations degrade the measurement trueness and precision.¹ The thresholds above which these degradations become significant need to be determined. They depend on many aspects of the analysis, including the type of instrument and its setting, the type of the introduction system, and the mass bias correction method. Several works have documented these perturbations and thresholds in different conditions, including different instruments [e.g., Micromass IsoProbe (Albarède and Beard, 2004), Thermochemical Axiom (Dideriksen et al., 2006), Nu Plasma (Millet et al., 2012; Sun et al., 2013; Finlayson et al., 2015; Zheng et al., 2018), and Thermo-Scientific Neptune (Schoenberg and von Blanckenburg, 2005; Dauphas et al., 2009; He et al., 2015)], different introduction systems [liquid, with or without desolvation (Schoenberg and von Blanckenburg, 2005; Dideriksen et al., 2006; Millet et al., 2012; Sun et al., 2013; Finlayson et al., 2015) (Schoenberg and von Blanckenburg, 2005; Dauphas et al., 2009; He et al., 2015), and laser ablation (Zheng et al., 2018)], and different mass bias correction methods [standard bracketing (Schoenberg and von Blanckenburg, 2005; Dauphas et al., 2009; Sun et al., 2013; He et al., 2015), Ni or Cu doping (Schoenberg and von Blanckenburg, 2005), and double-spiking (Dideriksen et al., 2006; Millet et al., 2012; Finlayson et al., 2015)]. For about a decade, Fe isotope analyses with ^{57}Fe – ^{58}Fe double-spike mass bias corrections are increasingly used in some fields, for instance, seawater studies (Conway et al., 2013; Dauphas et al., 2017; Klar et al., 2018; Lacan et al., 2008; Rudge et al., 2009). The latter relies on the combined use of the three Fe isotope ratios (i.e., the combined use of the four isotope measurements). This implies that isobaric interferences on both ^{54}Fe and ^{58}Fe (from ^{54}Cr and ^{58}Ni) can impact the final result, while in the case of standard bracketing or internal doping with another element, the final result may be obtained independently of one or both of these interferences (for instance when looking at the $^{57}\text{Fe}/^{56}\text{Fe}$ ratio only with the standard bracketing method, the final result is independent of the ^{54}Fe and ^{58}Fe measurements; or when looking at the $^{56}\text{Fe}/^{54}\text{Fe}$ and $^{57}\text{Fe}/^{54}\text{Fe}$ ratios, with the Ni doping method, the result is

¹The metrology vocabulary used in the present study refers to the “International vocabulary of metrology—Basic and general concepts and associated terms (VIM)” (2012) and the International Standard ISO 5725-1 (1994).

ISO 5725 uses two terms “trueness” and “precision” to describe the accuracy of a measurement method. “Trueness” refers to the closeness of agreement between the arithmetic mean of a large number of test results and the true or accepted reference value. “Precision” refers to the closeness of agreement between test results (ISO 5725-1, 1994).

The measurement “trueness” is inversely related to systematic measurement error, the latter being estimated by the “measurement bias.” “Measurement accuracy” is sometimes erroneously used instead of “measurement trueness” (VIM 2012).

“Measurement precision is used to define measurement repeatability, [...] and measurement reproducibility.” “Sometimes measurement precision is erroneously used to mean measurement accuracy” (VIM 2012).

The measurement “accuracy” is the “closeness of agreement between a measured quantity value and a true quantity value of a measurand” (VIM 2012). A measurement that is characterized at the same time by a good trueness (i.e. small bias, i.e. small systematic error) and a good precision (i.e. a good repeatability for instance) is characterized by a good accuracy.

For the sake of clarity, the term “perturbation” is used here instead of the term “influence quantity” as defined in the 2nd edition of the VIM as follows: “quantity that is not the measurand but that affects the result of the measurement.”

TABLE 1 | MC-ICPMS and desolvator operation parameters.

Neptune MC-ICPMS (Thermo-Scientific)	
RF power	1230 W
Acceleration voltage	10 kV
Mass analyzer pressure	$1.9 \cdot 10^{-8}$ mbar
Extraction lens	-2000 V
Overall transmission efficiency	0.03%
Neptune Plus MC-ICPMS (Thermo-Scientific)	
RF power	1230 W
Acceleration voltage	10 kV
Mass analyzer pressure	$1.9 \cdot 10^{-8}$ mbar
Extraction lens	-2000 V
Overall transmission efficiency	0.05%
Apex-IR desolvator (Elemental Scientific)	
Heater temperature	105°C
Chiller temperature	2°C
Nitrogen gas flow	0
Nebulizer	ESI MicroFlow PFA 75
Sample uptake rate	65–85 μLmin^{-1}
Overall mass discrimination	2.1–3.2% per atomic mass unit

independent of the ^{58}Fe measurement). Consequently, the impact of these perturbations may be different when using double-spike mass bias corrections compared to the other methods. In addition, the contribution of potential interferences cannot be verified in a three-isotope plot when using double-spiking (while this is possible with standard bracketing). However, the double-spike mass bias correction technique has several advantages: it allows to 1) measure the instrumental mass fractionation at the time of sample analysis (not before and after as with standard bracketing), 2) measure the Fe instrumental mass fractionation with Fe itself (not with another element as with Ni or Cu doping), and 3) correct for potential chemical mass fractionation during sample processing (by adding the

double-spike to the sample prior the chemical treatments, such as purification procedures, which other techniques do not allow). While some studies have documented thresholds for these isobaric Cr and Ni interferences below which reliable measurements could be obtained with standard bracketing and internal doping mass bias corrections (Schoenberg and von Blanckenburg, 2005; Dauphas et al., 2009; He et al., 2015), these thresholds have not been documented to date to our knowledge for double-spike mass bias corrections. In addition, while Thermo-Scientific Neptune and Neptune Plus MC-ICPMS are the most commonly used instruments nowadays for Fe isotope analyses, the impact of potentially major molecular ion interferences, such as $^{16}\text{O}^{40}\text{Ca}^+$ on $^{56}\text{Fe}^+$, or potential matrix effects associated to major elements such as Si or Na, have not been reported for Fe isotope measurements with these instruments to date to our knowledge.

The present work therefore presents for the first time a study of the impact of Cr, Ni, Ca, Si, and Al interferences and Na, Mg, Ca, K, and Mo matrix effects on iron isotope measurements using Thermo-Scientific Neptune and Neptune Plus MC-ICPMS with desolvation and a mass bias correction with a ^{57}Fe - ^{58}Fe double-spike. These results document thresholds below which reliable measurements could be obtained, and point toward elements, which require particular attention. They should provide a useful framework for future Fe isotope analyses.

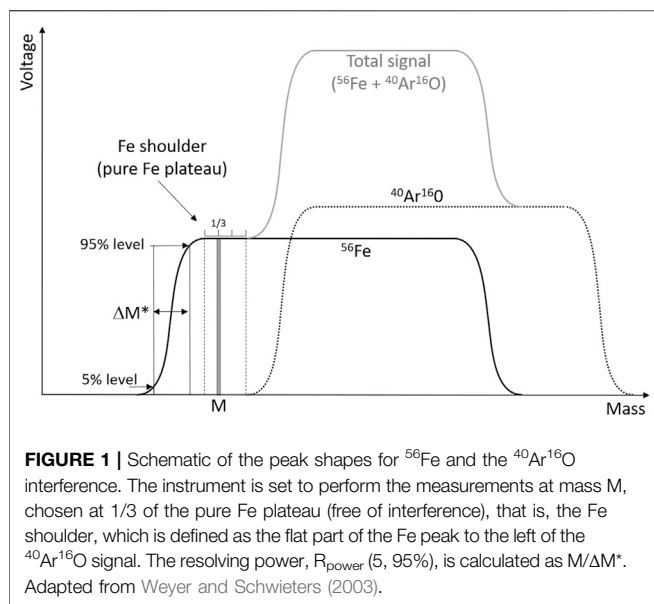
EXPERIMENTAL

The major part of the analyses was made using the Neptune MC-ICPMS (Thermo-Scientific) of the Observatoire Midi Pyrenees (Toulouse, France), while some measurements (detailed below) were made with the Neptune Plus MC-ICPMS (Thermo-Scientific) of the Observatoire de Recherche Méditerranéen de l'Environnement (Université de Montpellier, France). They were equipped with Ni X skimmer cones, with a standard sampler cone for the Neptune and a Jet sampler cone for the Neptune Plus and operated with high-resolution slits (25 μm). All samples, dissolved in $0.32 \text{ mol L}^{-1} \text{ HNO}_3$, were introduced *via* a $75\text{-}\mu\text{L}/\text{min}$ PFA MicroFlow nebulizer (Elemental Scientific) and an Apex-IR desolvator (Elemental Scientific). Operation parameters are detailed in **Table 1**. Collector configurations are given in **Table 2**.

With such instruments, Fe isotope measurements may suffer from poly-atomic ion interferences due to argon oxy-hydroxides

TABLE 2 | Faraday cup configuration and isotopic abundances for Fe, Cr, and Ni.

Nominal Mass		53	54	56	57	58	60	61
Isotope abundance (%)	Cr	9.5	2.37	—	—	—	—	—
	Fe	—	5.8	91.7	2.2	0.28	—	—
	Ni	—	—	—	—	68.3	26.1	1.13
Neptune collector configuration		L4	L2	L1	H1	H2	H3	H4
Neptune Plus collector configuration		L3	L2	C	H1	H2	H4	—



(e.g., $^{40}\text{Ar}^{16}\text{O}$ and $^{40}\text{Ar}^{16}\text{O}^1\text{H}$) and argon–nitrogen species (e.g., $^{40}\text{Ar}^{14}\text{N}$). These elements, such as Ar, O, H, and N, were present in all the samples of the present study, either supplied by the argon gas, ambient air, or by the HNO_3 . The mass resolving power (Weyer and Schwieters, 2003) of this type of instrument does not allow to fully separate the Fe peaks from the interference peaks. However, because the Fe isotopes are slightly lighter than these interferences, their peaks only partially overlap (Figure 1). The light part of the Fe peaks is virtually free of interference, that is, the light peak tails of the interferences do not significantly contribute to the Fe signals (given that Fe to interference intensity ratios are not too small). This has been extensively described previously (Weyer and Schwieters, 2003) and is commonly referred to as measuring on the Fe “shoulder” of the peaks.

We used a ^{57}Fe – ^{58}Fe double-spike composed of 0.11, 3.21, 59.2, and 37.5% (mol/mol) of ^{54}Fe , ^{56}Fe , ^{57}Fe , and ^{58}Fe , respectively. The sample-to-spike Fe mass ratio was 2/3 (i.e., the sample double-spike mixture is composed of 40% w/w of sample and 60% w/w), to maximize beam intensities on the four Fe collectors and minimize error propagation (Rudge et al., 2009; Lacan et al., 2010; John, 2012).

Samples were run following the typical sequence: IRMM14—HEM—IRMM14—SAMPLE-1—IRMM14—SAMPLE-2—IRMM14—HEM—IRMM14—SAMPLE-3 . . . and so on, where “HEM” is an in-house hematite standard (produced at ETH Zurich) of known Fe isotopic composition ($\delta^{56}\text{Fe} = 0.53 \pm 0.06\%$, 2SD) and negligible contents of other elements (Poirasson and Freydiser, 2005; Lacan et al., 2008; Lacan et al., 2010), which allowed monitoring measurement accuracy.

In between each of the above, the background was measured *via* the measurement of a 0.32 mol L^{-1} HNO_3 solution in which all samples were dissolved.

Data extraction was performed as follows. The mean background signals (over a sequence, typically 12 h) were subtracted from the raw data of each sample or standard. Cr

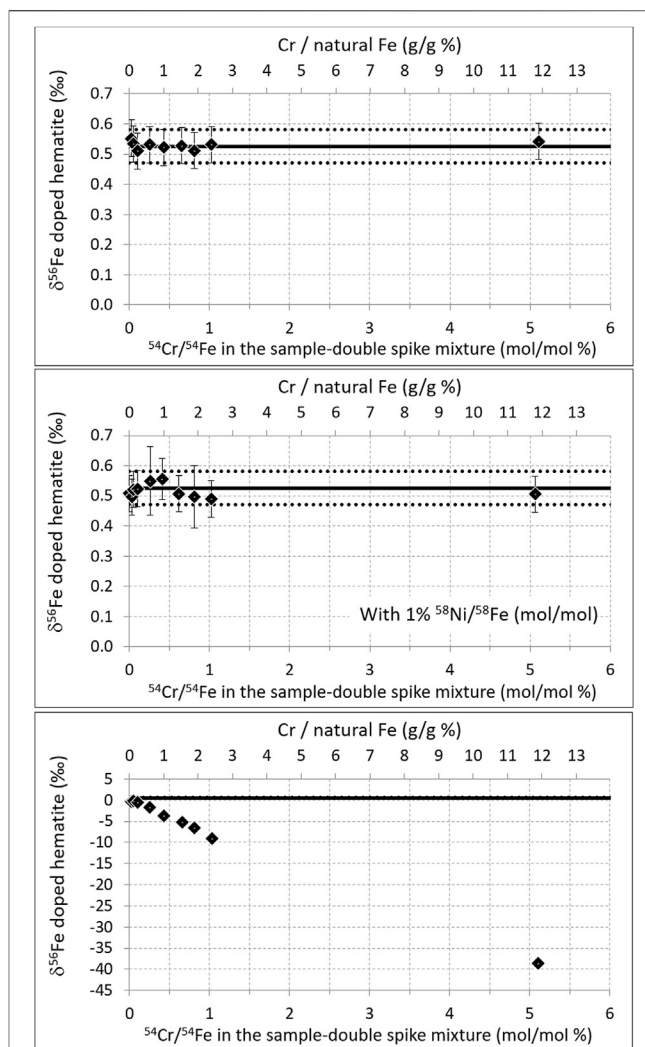
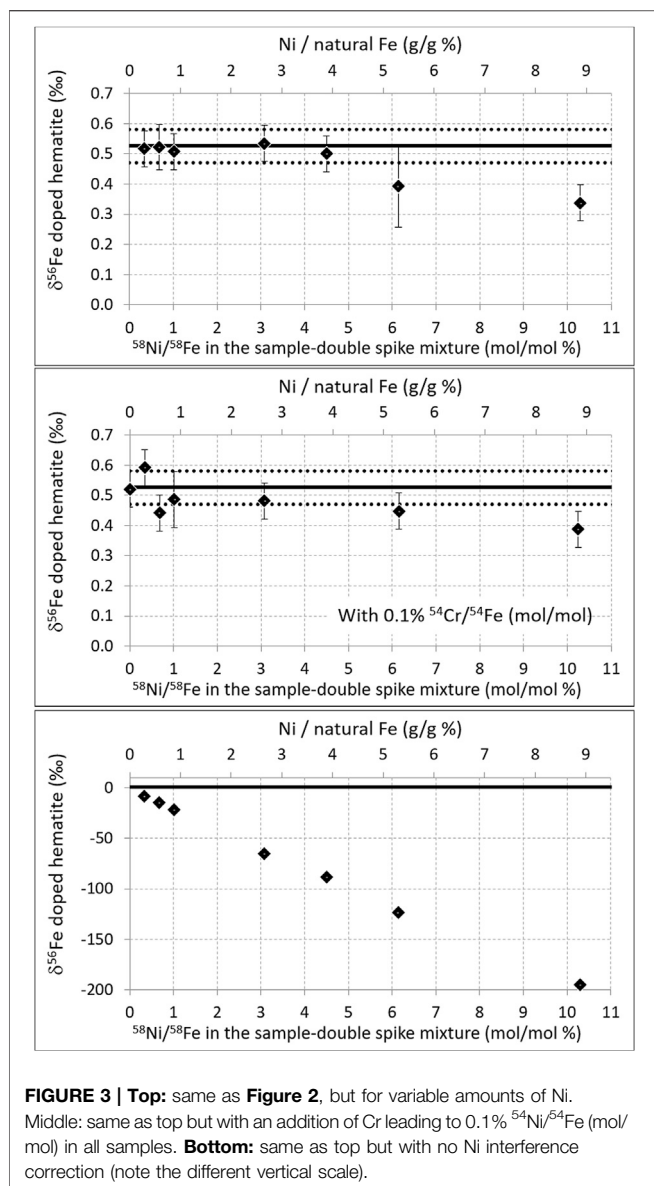


FIGURE 2 | Top: $\delta^{56}\text{Fe}$ of hematite (200 ppb natural Fe) doped with variable amounts of Cr. Middle: same as top but with an addition of Ni leading to 1% $^{58}\text{Ni}/^{58}\text{Fe}$ (mol/mol) in all samples. Bottom: same as top but with no Cr interference correction (note the different vertical scale). The level of Cr doping is expressed in two ways: i) the level of Cr (all isotopes) compared to that of natural Fe (i.e., not taking into account the double-spike, top x axis), and ii) the level of ^{54}Cr compared to ^{54}Fe in the analyzed solution, that is, in the sample double-spike mixture (bottom x axis). Diamonds are the averages of duplicate measurements (two Neptune analyses were performed for each data point reported on the plot). Error bars are 2 standard deviations of the duplicate measurements or the long-term internal precision (2SD) of our instrument (0.06‰), whichever the greatest (they are smaller than the symbol size on the bottom panel). The horizontal black lines show the $\delta^{56}\text{Fe}$ of pure (un-doped) hematite measured during the same analytical session; mean value (solid line) \pm 2 standard deviations (dotted lines).

and Ni isobaric interferences (on ^{54}Fe and ^{58}Fe) were corrected for by estimating ^{54}Cr and ^{58}Ni , from the ^{53}Cr and ^{60}Ni signals, the natural Cr and Ni abundances, and the instrumental isotopic fractionation. The latter was computed for Fe from the background and isobaric interference-corrected signals, using the Siebert et al. iterative calculation (Siebert et al., 2001). Instrumental mass fractionations for Cr and Ni were therefore



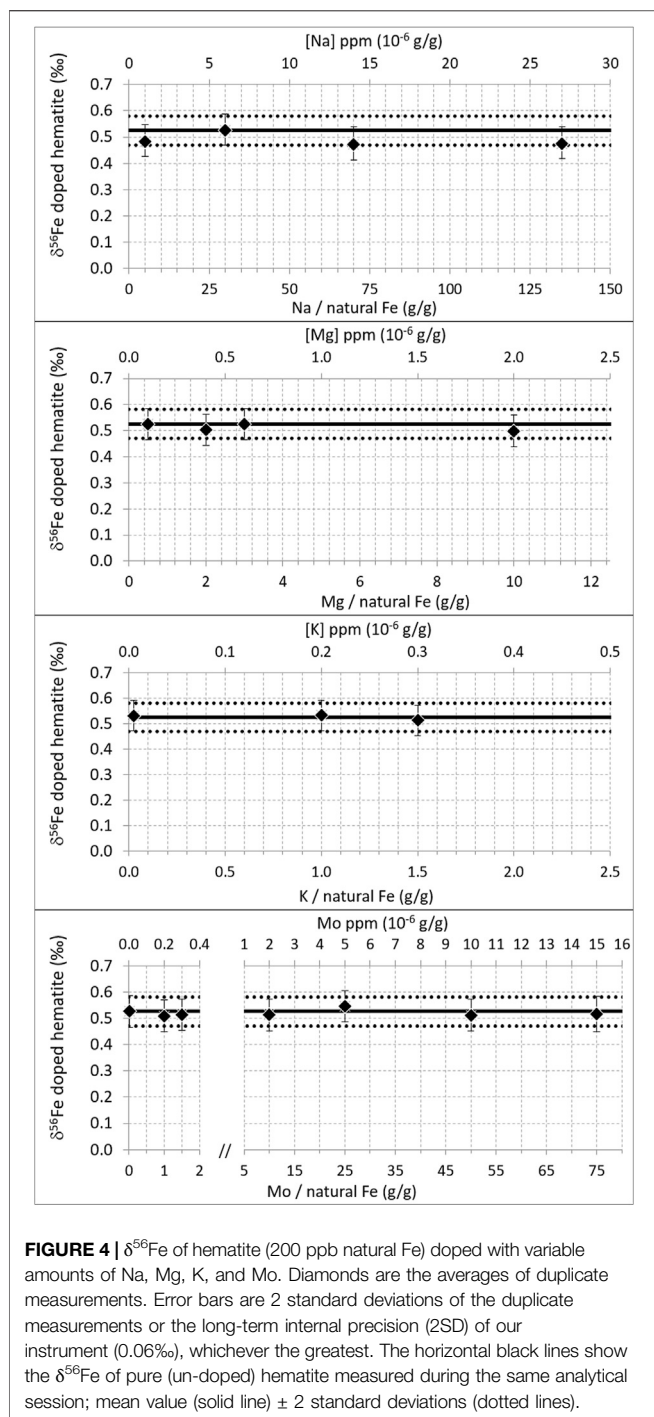
assumed to be the same as for Fe. The above procedure is circular because it requires the instrumental mass fractionation to estimate the Cr and Ni interferences, but it also requires the Cr and Ni interferences to estimate the instrumental mass fractionation. This problem was circumvented through another iterative calculation. Finally, for each sample, the $\delta^{56}\text{Fe}$ value was calculated relative to the mean of its two bracketing IRMM14. Precision and accuracy of this method were evaluated through repeated measurements of the pure ETH hematite standard. Considering all its measurements during sessions dedicated to the present study, between June 2015 and October 2018, its mean value was $\delta^{56}\text{Fe} = 0.525 \pm 0.054\text{‰}$ (2SD, $N = 89$). This is in perfect agreement with data previously reported by different groups (Lacan et al., 2010 and references therein). Average background signals were 0.02% of the sample signals, on average for the four isotopes (in any case between 0.01 and 0.04%).

As mentioned above, the double-spike mass bias correction involves the combined use of the three Fe isotope ratios, that is, the combined use of the four Fe isotopes. A perturbation on any of the four isotope measurements could therefore impact the final result. To determine the impact of interferences and matrix effects, samples were prepared as follows. A 200 ppb [10^{-9} g (Fe)/g (HNO_3)] hematite solution, mixed with the double-spike (300 ppb), was doped with variable amounts of mono-elemental standard solutions of Cr, Ni, Cr + Ni, Al, Si, Na, Mg, K, Ca, Mo, and Na + Mg + K + Ca + Mo. These elements were chosen either because they produce spectral interferences (Cr, Ni, Al, Si, and Ca) or because they might be found at significant levels in the final purified solutions (either because of their high initial abundances in natural samples or because of imperfect chemical separation). Most Fe isotope measurements at LEGOS are performed on seawater samples. Although Na, Mg, Ca, and K are very efficiently separated from Fe through preconcentration and purification procedures [chelating and ion exchange resins, e.g., (Schoenberg and von Blanckenburg, 2005; Borrok et al., 2007; Lacan et al., 2010)], even very small fractions of the initial contents of these elements may result in significant quantities in purified samples. For example, 0.0001% of Na in an open ocean surface seawater sample represents a mass 2000 times greater than its typical Fe content. Other types of samples and other elements may present similar problems. For example, 0.0001% of Ca initially present that tooth samples (enamel or dentin) can represent ~ 50 times their iron content (Teruel et al., 2015). Similarly, 0.1% of the calcium initially present in some iron-depleted calcites can represent ten times their iron content (von Blanckenburg et al., 2008). Additionally, Mo was chosen because the efficiency of its separation on the most common Fe purification procedures (using notably AG 1-x4, AG 1-x8, or AG MP-1 ion exchange resins) depends on its oxidation state, which makes this separation more uncertain than for other elements (Lacan et al., 2010). Therefore, although this study has been motivated by the seawater work at LEGOS, it aims at providing a general framework for iron isotope measurements on any kind of samples for which imperfect purification may be expected.

The abovementioned mixtures were measured to assess whether or not the correct isotopic composition of the hematite could be determined. All measurements were performed with resolving powers $R_{\text{power}} (5.95\%) = M/\Delta M^* \geq 8,000$ and plateau width ≥ 200 ppm [that is sufficient for Fe isotope measurements (Weyer and Schwieters, 2003)]. The doubly charged effect is often negligible as doubly charged ions are rare in the conditions of the plasma (Houk, 1986); therefore, Cd was not investigated.

RESULTS AND DISCUSSION

All results presented below were obtained with the Neptune MC-ICPMS, unless otherwise indicated (for Neptune Plus data). The effect of ^{54}Cr and ^{58}Ni isobaric interferences on the hematite measurements is displayed in Figure 2 and Figure 3. Figure 2 shows that even when the hematite was doped with



11.9% g (Cr)/g (hematite Fe) (i.e., the solution analyzed was 200 ppb hematite Fe + 24 ppb Cr + 300 ppb double-spike; 200 ppb \times 11.9 % = 24 ppb), which lead to a $^{54}\text{Cr}/^{54}\text{Fe}$ slightly larger than 5% (mol/mol), the correct isotopic composition of the hematite was found. The presence of a small quantity of Ni (leading to 1% $^{58}\text{Ni}/^{58}\text{Fe}$ (mol/mol) in the analyzed solution, level to which interference corrections have been validated for standard bracketing measurements (Schoenberg and von Blanckenburg, 2005) did not significantly change the results.

This demonstrates that up to this level, the ^{54}Cr interference can be successfully corrected for. This value is similar to the threshold documented earlier with the same type of instrument (Neptune) but with a standard bracketing mass bias correction (Dauphas et al., 2009). Note that these results are obtained, thanks to the interference correction; without correction, significant errors (of the order of 1‰) occur as soon as the Cr content reaches 0.1% g (Cr)/g (hematite Fe), and this error increases dramatically with increasing Cr content (it reaches 39‰ for 11.9% g (Cr)/g (hematite Fe), cf. **Figure 2** bottom panel).

Figure 3 shows that when the hematite was doped with up to 3.9% g (Ni)/g (hematite Fe) (i.e., 200 ppb hematite + 8 ppb Ni + 300 ppb double-spike; i.e., $^{58}\text{Ni}/^{58}\text{Fe} \approx 4.5\%$ mol/mol), the correct hematite isotopic composition was found. Above that threshold, the values obtained are significantly deviated from the true value. Again, this threshold is similar to that documented earlier with a Neptune and a standard bracketing mass bias correction (Dauphas et al., 2009). The presence of a small quantity of Cr (leading to 0.1% $^{54}\text{Cr}/^{54}\text{Fe}$ (mol/mol) in the analyzed solution, again the level to which interference corrections have been validated for standard bracketing measurements (Schoenberg and von Blanckenburg, 2005), did not significantly change the results. As for Cr, these results are obtained, thanks to the interference corrections; without correction, very significant errors (of the order of 8‰) occur as soon as the Ni content reaches 0.3% g (Ni)/g (hematite Fe), and this error increases dramatically with increasing Ni content [it reaches 88‰ for 3.9% g (Ni)/g (hematite Fe), see **Figure 3** bottom panel].

Figure 4 shows the effect of Na, Mg, K, and Mo doping (on the 200 ppb natural Fe solution, mixed with 300 ppb double-spike). It shows that below certain doping concentrations (up to 27 ppm Na, 2 ppm Mg, 0.3 ppm K, and 15 ppm Mo), the correct isotopic composition of the hematite was found. This demonstrates that

TABLE 3 | First ionization energies and degrees of ionization for each element of interest (Houk, 1986). Degree of ionization calculated from Saha equation with T plasma: 7,500 K and $n_e = 10^{15} \text{ cm}^{-3}$ (where n_e is the electron density) (Boumans, 1966; Houk, 1986).

Element	Ionization energy (eV)	Degree of ionization (%)
Na	5.14	100
Mg	7.64	98
Al	5.99	98
Si	8.15	85
K	4.34	100
Ca	6.11	99
Ar	15.76	0.04
Fe	7.90	96
Ni	7.64	91
Mo	7.10	98

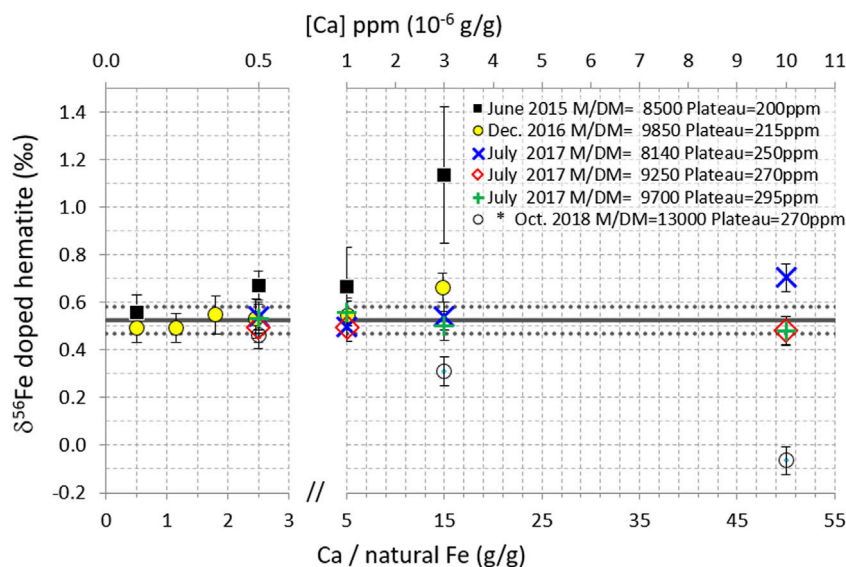


FIGURE 5 | $\delta^{56}\text{Fe}$ of hematite (200 ppb natural Fe) doped with variable amounts of Ca, for five measurement sessions on the Neptune and (*) one session on the Neptune Plus (2018). Data points are the averages of replicate measurements ($n = 2-5$), except for 2017 data (no replicate). Error bars are 2 standard deviations of the replicate measurements for the 2015–16 data and the long-term internal precision (2SD) of the instrument (0.06‰) for the 2017 data (no replicate) and for the 2018 Neptune Plus data (2SD of replicates being smaller than long term internal precision). The horizontal black lines show the $\delta^{56}\text{Fe}$ of pure (un-doped) hematite measured during the same analytical sessions; mean value (solid line) \pm 2 standard deviations (dotted lines).

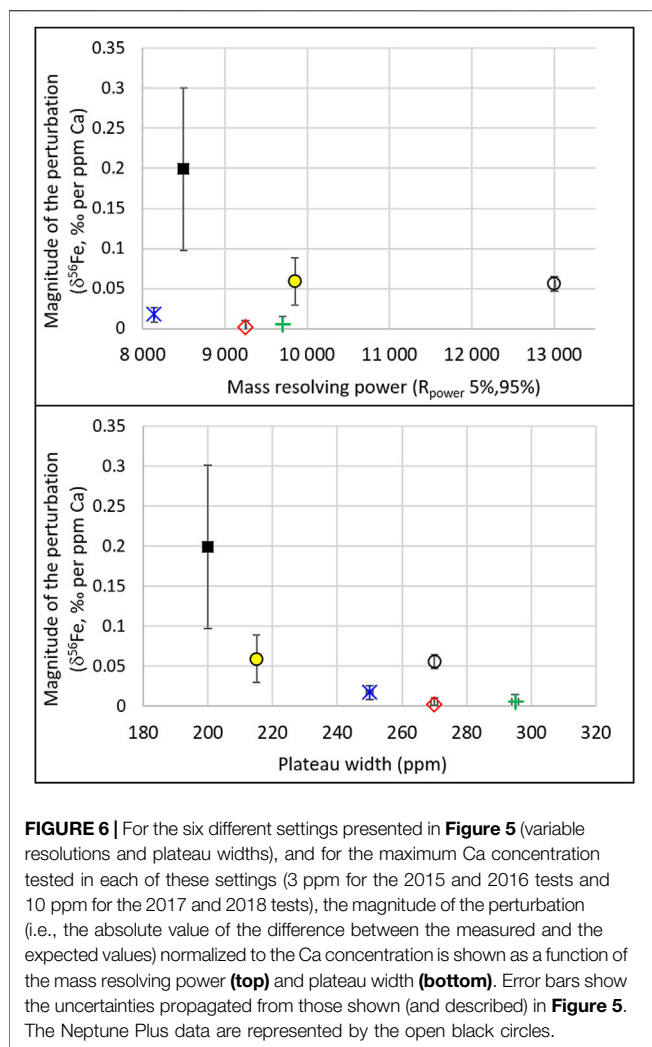
up to these levels and despite the lower ionization energies (and thus higher degrees of ionization) of these elements relative to Fe (Table 3), these matrix effects are negligible, or at least accurately corrected for with the double-spike. Larger concentrations have not been tested. These levels were chosen because they represent the maximum values we encounter with purified seawater samples we usually work on. They correspond to Na/Fe = 135, Mg/Fe = 10, K/Fe = 1.5, and Mo/Fe = 75 (g/g, where Fe is natural Fe). For Na and Mg, these ratios are significantly larger than those documented previously (Na/Fe = 5 and Mg/Fe = 2 g/g, with a ^{54}Fe - ^{58}Fe double-spike and a different instrument, Thermochemical AXIOM MC-ICPMS) (Dideriksen et al., 2006), which allows extending the range within which Na and Mg matrix effects can be neglected. For K and Mo, these thresholds have not been documented before to our knowledge.

The effects of calcium doping, as shown in Figure 5, are more significant and more complicated. Calcium may perturb the $\delta^{56}\text{Fe}$ measurements in two ways. Residual amounts of Ca after sample purification may lead to matrix effects due to its high concentrations in certain natural samples (e.g., rocks, waters, and shells). In addition, similarly to Ar, Ca may also cause polyatomic interferences on Fe isotopes, as shown previously (Dideriksen et al., 2006). The molecular ions such as $^{40}\text{Ca}^{14}\text{N}$, $^{40}\text{Ca}^{16}\text{O}$, $^{40}\text{Ca}^{16}\text{O}^{1}\text{H}$, and $^{42}\text{Ca}^{16}\text{O}$ may interfere on ^{54}Fe , ^{56}Fe , ^{57}Fe , and ^{58}Fe , respectively. The weak Ca ionization energy compared to Fe (and thus high degree of ionization, cf. Table 3) potentially reinforces both perturbations.

Measurements performed in June 2015, together with the other tests presented above, show that with 0.5 ppm Ca (Ca/Fe = 2.5 g/g where Fe is natural Fe), the measurement of the hematite

$\delta^{56}\text{Fe}$ was significantly biased; it was found to be 0.67‰ instead of $0.52\text{‰} \pm 0.06$ (2SD) (black squares in Figure 5). This evidences a much higher $\delta^{56}\text{Fe}$ measurement sensitivity to the Ca contents compared to other elements tested for matrix effects (e.g., no perturbation up to Na/Fe = 135 or Mo/Fe = 75 g/g). This observation suggests that the observed perturbations are more likely due to interferences rather than to matrix effects. This confirms previous works (Dideriksen et al., 2006) and is further confirmed by additional information given below. These levels of Ca impurities (Ca/Fe = 2.5 g/g) after purification are not frequent. However, they may be encountered in some cases. For instance, out of 18 marine sediment interstitial water samples purified on AG-MP1 column, we found seven with Ca/Fe (g/g) ratios >1 (with a maximum value of 4.6).

As explained at the beginning of the experimental section, measurements are performed on the peak shoulders, to stay away from the ArO, ArOH, and ArN interferences (cf. Figure 1). This is usually done by looking at the peak shapes produced by pure Fe solutions and by tuning the instrument to maximize the width of the Fe shoulder, that is, the Fe plateau, and the slope of the peak flanks, that is, the mass resolving power. Then the absence of significant interference contributions to the Fe signal is checked by comparing the peak shapes produced by pure Fe solutions (where both Fe and the interferences contribute to the peak shapes) and HNO₃ solutions (where only interferences are present) (Weyer and Schwieters, 2003). ^{40}Ca is very slightly heavier than ^{40}Ar ($M^{40}\text{Ca} = 39.96259$ u, $M^{40}\text{Ar} = 39.96238$ u). Therefore, if the Fe plateau is free of the argon molecular ion interference and if calcium molecular ions are not significantly more abundant than argon ones, then calcium should not cause



any significant interference on the Fe measurements. However, if the calcium molecular ions are significantly more abundant than the argon ones (because of a large Ca concentration in the sample or because of a large calcium molecular ion formation rate compared to that of argon), then the calcium molecular ion peak tail may be larger at a given mass than to that of the argon molecular ions, despite the fact that ^{40}Ca is slightly heavier than ^{40}Ar . In summary, resolving the argon molecular ion interferences is not a warranty that calcium molecular ion interferences will be resolved. The effects of potential calcium molecular ion interferences need, therefore, to be quantified.

Figure 5 displays the results obtained for the hematite doped with variable Ca amounts, during five different measurement sessions on the Neptune instrument (Neptune Plus data will be discussed next), for which the mass resolving power [R_{power} 5–95% (Weyer and Schwieters, 2003)] and plateau width were set to different values (8,500–13,000 and 200–295 ppm, respectively) notably by varying of the “Shape” parameter in the instrument tuning. It shows that during the sessions with the largest mass resolving power and plateau width, the perturbations

were the smallest. For instance, with a mass resolving power of 9,700 and a plateau of 295 ppm, no perturbation was detected even with 10 ppm Ca (for 200 ppb natural Fe). This corresponds to $\text{Ca}/\text{Fe} = 50$ (g/g where Fe is natural Fe). This is 500 times larger than the threshold documented previously ($\text{Ca}/\text{Fe} = 0.1$, with a ^{54}Fe - ^{58}Fe double-spike and a different instrument, Thermochemical AXIOM MC-ICPMS) (Dideriksen et al., 2006).

The relations among mass resolving power, plateau width, and perturbations are further displayed in **Figure 6**. The latter shows, for the maximum Ca concentration of the 5 different Neptune sessions (3 ppm for the 2015 and 2016 tests and 10 ppm for the 2017 tests), the magnitude of the perturbation (i.e. the absolute value of the difference between the measured and the expected values) normalized to the Ca concentration, as a function of the mass resolving powers and the plateau widths. For instance, in June 2015, the maximum Ca concentration tested was 3 ppm, and the magnitude of the perturbation for this maximum Ca concentration was 0.6‰; this leads to 0.2‰ perturbation per ppm Ca, for a resolution and plateau width of 8,500 and 200 ppm, respectively. While no relation is seen as a function of mass resolving power, the figure shows that the perturbation decreases with increasing plateau width. The relationship between plateau width and magnitude of the perturbation confirms that the perturbations are caused by interferences and not by matrix effects.

Calcium doping tests were also performed on the Neptune Plus instrument (**Figure 5**, **Figure 6**, open black circles). Results significantly differ from those obtained with the Neptune. First, perturbation magnitudes seem to be larger with the Neptune Plus than with the Neptune, for instance, for the same Ca concentration [$\text{Ca} = 10$ ppm], and the same plateau width (270 ppm), whereas an insignificant perturbation was measured on the Neptune (hematite $\delta^{56}\text{Fe} = 0.48$ instead of 0.53), and a large perturbation was measured on the Neptune Plus ($\delta^{56}\text{Fe} = -0.07$ instead of 0.53). Moreover, while the effect of Ca doping on the Neptune systematically led to positive biases, negative ones were obtained with the Neptune Plus (**Figure 5**). The principal difference between the two systems is the interface (cones and vacuum), named “Jet interface” on the Neptune Plus. Enhanced levels of polyatomic interferences, notably $^{40}\text{Ar}^{14}\text{N}^+$, have been reported with the Jet interface compared to the standard interface (Chernonozhkin et al., 2017). High levels of Ca may similarly lead to enhanced levels of $^{40}\text{Ca}^{14}\text{N}^+$ with the Jet interface compared to the standard one. This could explain the larger and negative biases observed on the Neptune Plus compared to those observed on the Neptune.

Interferences may also result from homonuclear diatomic ions such as $^{27}\text{Al}_2^+$ and $^{28}\text{Si}_2^+$, interfering on $^{54}\text{Fe}^+$ and $^{56}\text{Fe}^+$, respectively. This was tested on the Neptune Plus. Results are shown in **Figure 7**. For silicon, the correct isotopic composition of the hematite was found up to the maximum doping concentration tested here, 10 ppm ($\text{Si}/\text{Fe} = 50$ g/g where Fe is natural Fe). This demonstrates that up to this level, this interference is negligible. Such tests have not been previously published to our knowledge.

For Al, the correct isotopic composition is found up to $[\text{Al}] = 500$ ppb ($\text{Al}/\text{Fe} = 2.5$ g/g where Fe is natural Fe). However, significant perturbations are detected for concentrations equal and above $[\text{Al}] = 1$ ppm ($\text{Al}/\text{Fe} = 5$ g/g), with the perturbation

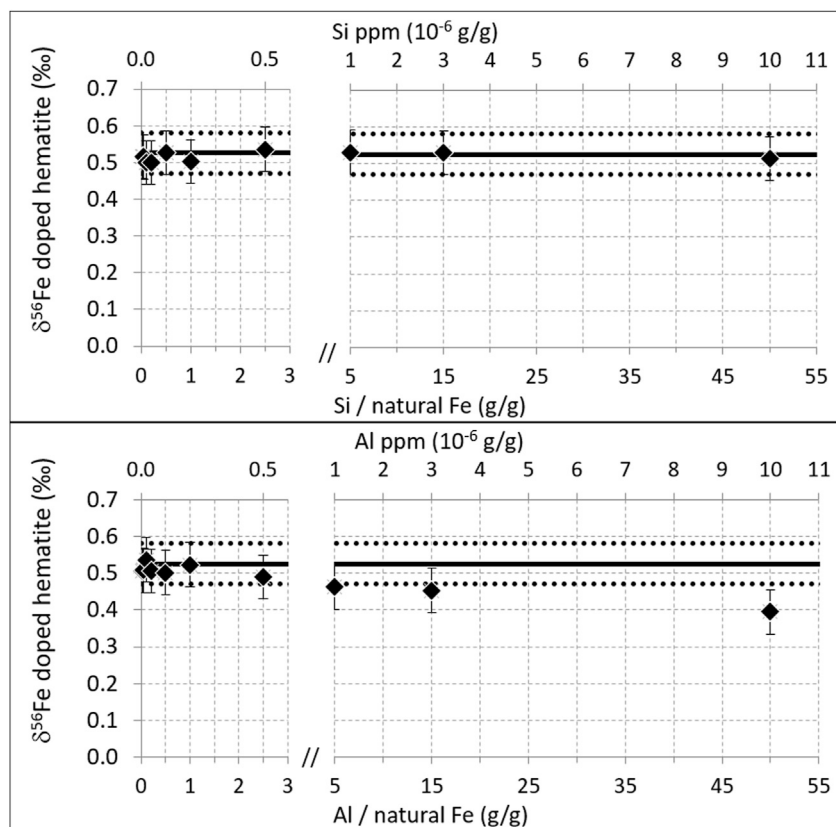


FIGURE 7 | Same as **Figure 4**, for Al and Si. Measurements performed on the Neptune Plus, in October 2018. Resolution 13,000, plateau 270 ppm, natural Fe concentration 200 ppb.

magnitude increasing with increasing Al concentration. This effect had been previously shown on a Nu Plasma instrument with a $^{57-58}\text{Fe}$ double-spike (Millet et al., 2012). However, this latter study found significant perturbations ($\delta^{56}\text{Fe}$ difference of 0.1‰) for much smaller Al contents, Al/Fe = 0.05 (g/g), while here no perturbation is detected up to Al/Fe = 2.5 (50 times higher).

Unlike Cr and Ni isobaric interferences, Ca and Al polyatomic ion interferences cannot be corrected for. While ^{54}Cr and ^{58}Ni interferences can be deduced from the measurement of ^{53}Cr and ^{60}Ni (cf. Experimental section), the intensity of Ca and Al polyatomic ion interferences depend on 1) their concentration, 2) the rate of production of the polyatomic ions, and 3) the precise mass (M in **Figure 1**) at which the measurements are performed. While the first point could be addressed by its measurement (in dynamic mode), the two last points depend on many uncontrolled and potentially variable parameters, such as the efficiency of the desolvating system, the plasma temperature, environment of the cones (vacuum, size of hole, and distances), or the magnetic field intensity (that varies very slightly). Because Ca and Al polyatomic ion interferences cannot be corrected for, their concentrations in the samples have to be below the indicated thresholds.

We preconize the following procedure. The calcium, aluminum, and iron concentrations should be determined in all purified samples prior to the MC-ICPMS measurements.

Then, while tuning the MC-ICPMS (cups alignments and peak shape optimization), in addition to pure Fe and HNO_3 solutions, pure Ca and Al solutions, with concentrations at least equal to their maximum values in samples, should be introduced and their peak shape measured, in order to verify that their polyatomic ion interference level at mass M are comparable to (and not larger than) that of the HNO_3 . Finally, and most importantly, a Fe isotopic standard doped with these Ca and Al levels should be regularly measured, to make sure that these elements do not significantly perturb the measurements. This is especially important for samples with high initial calcium contents, such as calcite minerals or tests, bones and teeth, but also marine samples, including seawater (Heimann et al., 2010; Jaouen et al., 2012; Emmanuel et al., 2014; Labatut et al., 2014; Revels et al., 2015; Klar et al., 2017), or high initial aluminum content such as silicate rocks or atmospheric dust samples.

CONCLUSION

While samples are usually purified before Fe isotopic composition MC-ICPMS analysis, some elements may remain in the purified samples at significant concentrations, in some circumstances. They may degrade the measurement accuracy. We have tested the effect of

the following perturbations on the accuracy of the Fe isotopic composition measurement with a Neptune and a Neptune Plus MC-ICPMS, using a ^{58}Fe - ^{57}Fe double-spike mass bias correction:

- Cr and Ni isobaric interferences;
- Na, Mg, K, Ca, and Mo matrix effects;
- Ca polyatomic ion spectral interferences;
- Al and Si homonuclear diatomic ion spectral interferences.

We found that for 200 ppb natural Fe solutions (mixed with 300 ppb of double-spike), the measurements (after Cr and Ni interference and mass bias corrections) were unperturbed up to the following levels (in the following ratios, Fe stands for natural Fe; i.e., the double-spike contribution is not included; results were obtained with the Neptune unless otherwise indicated):

- [Cr] = 24 ppb, that is, Cr/Fe = 0.12 (g/g)
- [Ni] = 8 ppb, that is, Ni/Fe = 0.04 (g/g)
- Na/Fe = 175, Mg/Fe = 10, K/Fe = 1.5, and Mo/Fe = 75 (g/g)
- For Ca, the perturbation was shown to be related to the Fe plateau width, and therefore to be due to Ca polyatomic ion spectral interferences ($^{40}\text{Ca}^{14}\text{N}$, $^{40}\text{Ca}^{16}\text{O}$, $^{40}\text{Ca}^{16}\text{O}^1\text{H}$, and $^{42}\text{Ca}^{16}\text{O}$), rather than to matrix effects. With the Neptune MC-ICPMS, it was found that significant perturbations could occur with only 0.5 ppm Ca (Ca/Fe = 2.5) for a plateau width of 200 ppm, while no perturbation could be detected with 10 ppm Ca (Ca/Fe = 50) for a plateau width of 295 ppm. The Neptune Plus MC-ICPMS was found more sensitive than the Neptune to the Ca polyatomic ion interferences.
- $^{28}\text{Si}_2^+$ interference was found insignificant up to Si/Fe = 50 (on the Neptune Plus).
- $^{27}\text{Al}_2^+$ interference was found insignificant up to Al/Fe = 2.5, but significant over this threshold (on the Neptune Plus).

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These results imply that while working with samples with potentially high calcium or aluminum contents (including purified seawater or silicate rocks), Ca and Al perturbations should carefully be taken into account (in addition to Cr and Ni) while 1) tuning the instrument and 2) checking the measurement accuracy with isotopic standards (i.e., doping the isotopic standard with Ca and Al levels comparable to those of the samples).

AUTHOR CONTRIBUTIONS

FL conceived the study. CP, LA, and JK prepared the samples. FL, CP, LA, and JK performed the MC-ICPMS measurements. JC and RF brought their expertise on MC-ICPMS, and FL wrote the manuscript with contributions from JK, JC, LA, and RF.

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