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# Revealing the incorporation of cerium in fluorapatite

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## ABSTRACT

Fluorapatite (FAP, nominally  $\text{Ca}_5(\text{PO}_4)_3\text{F}$ ) is the most common phosphate mineral at the Earth's surface and a main host for rare earth elements (REE) in magmatic and hydrothermal ore deposits and in marine sediments. Our understanding of the enrichment process of REE in FAP rests upon two foundations: (1) being able to elucidate the thermodynamic driving force for their partitioning between the Ca1 and Ca2 structural sites, and (2) being able to determine how the substitution of REE(III) for Ca(II) is charge compensated. A main unsolved question is the marked preference of the larger light REE (lanthanum  $\rightarrow$  samarium) for the smaller Ca2 site. We used density functional theory (DFT) and high-energy resolution fluorescence-detected extended X-ray absorption fine structure (HERFD-EXAFS) spectroscopy to gain detailed insight into the bonding energy, electronic structure, and short-range order of cerium (Ce) in natural FAP. Results show that Ce(III) has a marked preference for a Ca2 site where the nearest five-valent phosphorous cation is replaced with a tetravalent silicon cation, thus balancing the charge excess of the Ce impurity locally. Atomic charge calculations show that the Ca2 site is more ionic than the Ca1 site and that the energetics of the site preference is linearly correlated to the ionization energy of the substituent. Cations with a low energy

of ionization, like Ce, preferably occupy the Ca2 site. Novel combination of HERFD-EXAFS spectroscopy and DFT appears to be the most straightforward and reliable way to assess the crystal chemistry of trace elements in compositionally complex natural materials, and opens a previously unavailable avenue for mechanistic investigation of metal enrichment in ore deposits.

## INTRODUCTION

FAP widely occurs in calcium-rich igneous rocks<sup>1,2</sup> and as biogenic remain in ancient (e.g., phosphorite)<sup>3-7</sup> and modern (e.g., deep-sea mud) marine deposits.<sup>8-12</sup> The crystal structure of FAP permits a wide range of cation and anion substitutions.<sup>13-18</sup> In particular, it has two calcium sites, a larger Ca1 site surrounded by 6 + 3 oxygen and a smaller Ca2 site surrounded by 6 oxygen and one fluorine,<sup>19,20</sup> which can accommodate monovalent, divalent, and trivalent cations as substituents, such as Na, Sr, Cd, and rare-earth elements of the lanthanide series (REE) and yttrium (REY).<sup>8-12,21-25</sup> Igneous and sedimentary fluorapatites commonly contain ore grades of several percent (w/w) REY.<sup>9,26-29,12,30,31</sup> REY enrichment occurs in contact with magmatic and hydrothermal fluids for igneous FAP,<sup>30,32-34</sup> and in contact with bottom seawaters and pore waters for detrital biogenic FAP.<sup>11,35-37</sup>

In addition to having a different size, the Ca1 and Ca2 sites also differ in their exposure to fluids. The crystal framework of apatite consists of face-shared Ca1 octahedra linked through oxygen atoms shared with PO<sub>4</sub> tetrahedra (Figure 1a).<sup>19,20</sup> The Ca1-PO<sub>4</sub> framework circumscribes one-directional tunnels along the *c* axis demarcated by CaO<sub>6</sub>F polyhedra (Ca2 site) on the walls that are in contact with circulating solutions in the lumen of the tunnels and along the external surfaces of the mineral grains. The distribution of REE between the two Ca sites of FAP has been addressed in many studies performed on natural and synthetic minerals.<sup>19,38-46</sup> REE prefer the Ca2 site over the Ca1 site with a Ca2/Ca1 site occupancy ratio that decreases from light (LREE) to heavy (HREE) REE. Site preference in FAP has been rationalized variously in terms of ionic radii of REE(III) and Ca(II), equalization of Ca1, Ca2, and F bond valences, bond covalencies, and minimization of strain effects.<sup>40-43,46-48</sup> Still, no satisfactory explanation has been offered to support the preference of LREE with a radius larger than Ca(II) for the smaller Ca2 site.

An estimated Ca2/Ca1 occupancy ratio of 3.67 has been estimated for Ce in singly-REE substituted synthetic FAP by Fleet and Pan in 1995 using single-crystal X-ray diffraction.<sup>42</sup> As comprehensive as this study is, Fleet and Pan further noted in 1997 that “quantitative transference of laboratory REE site preferences to natural apatites is frustrated by the compositional complexity in nature”.<sup>43</sup> Crystallographic data on natural apatites indeed provide site occupancies averaged over all

substituents, and therefore lack structural sensitivity to specific REE. The site occupancy of a given REE can be appraised in principle by extended X-ray absorption fine structure (EXAFS) spectroscopy as this technique is element selective. However, a major roadblock in analyzing multi-elemental Earth materials by EXAFS is the overlap of the X-ray fluorescence lines from two or more elements,<sup>8,49,50</sup> and therefore its application to REE-containing natural apatites is unprecedented.

Another key question in understanding the uptake of REY into FAp is how the increase of positive electronic charge resulting from the REE(III) for Ca(II) substitution is compensated chemically. Two coupled substitutions have been proposed:  $\text{REE(III)} + \text{Na}^+ \leftrightarrow 2\text{Ca(II)}$  and  $\text{REE(III)} + \text{Si}^{4+} \leftrightarrow \text{P}^{5+} + \text{Ca(II)}$ .<sup>39-43,46</sup> X-ray refinement of singly La, Nd, Gd, and Dy substituted synthetic FAp found that Na occupies the Ca1 site<sup>42,46</sup> (Na1), and that charge balance is maintained with both Na and Si ( $\text{Na} > \text{Si}$ ).<sup>46</sup> Atomistic calculation of the formation enthalpy of  $\text{Ca}_9\text{Nd(PO}_4)_5\text{SiO}_4\text{F}_2$  and  $\text{Ca}_8\text{NdNa(PO}_4)_6\text{F}_2$  showed that the first compound is thermodynamically more stable than the second, and therefore that the heterovalent REE(III) - Ca(II) substitution is more likely charge balanced by parallel  $\text{Si}^{4+} \leftrightarrow \text{P}^{5+}$  substitution than by  $\text{Na}^+ \leftrightarrow \text{Ca}^{2+}$  substitution,<sup>47</sup> at odds with diffraction results.<sup>46</sup> Still, the theoretical study confirmed the preference of Na for the Ca1 site and of Nd for the Ca2 site, and predicted that the Nd-Si coupled substitution takes place at short distance in the FAp structure.

Despite the large amount of literature available on REE-containing FAp, several questions and ambiguities on the atomic-scale structure of this common phosphate mineral remain. Although LREE most likely occupy the Ca2 site in natural FAp, direct confirmation by EXAFS spectroscopy is clearly desirable. Here, the complication arising from the multi-elemental nature of natural FAp was circumvented by filtering the unwanted X-ray fluorescence lines with analyzer crystals in high energy resolution fluorescence detection mode (HERFD-EXAFS).<sup>51-53,50</sup> The nature of the charge compensating cation and the energetic driving force for the site preference of REE remain unresolved. The two questions were addressed theoretically by optimizing the geometries and calculating the total energies of (1) Na/Si charge-compensated Ce(III)-FAp models, and (2) Mg(II), Sr(II), Ba(II), and Cd(II)-substituted FAp models. The choice of Ce as REE was dictated because of its broad interest in earth science aiming at inferring magmatic and marine sediment processes, and paleo-redox and paleo-environmental reconstructions.<sup>8,36,54-60</sup> For the sake of convenience, we will start by presenting the quantum-mechanical results. The geometrically-optimized structures will be used next to interpret the Ce-EXAFS data.

## MATERIALS AND METHODS

**DTF.** Density functional theory (DFT) computations were performed in the Vienna Ab-initio Simulation Package (VASP) software.<sup>61,62</sup> The projector augmented wave (PAW) pseudopotentials<sup>63,64</sup> were used for the core-electron interaction, in combination with a converged energy cutoff of 500 eV for the planewave basis set. The energies and forces were evaluated at the generalized gradient approximation (GGA) with a Perdew-Burke-Ernzerhof (PBE)<sup>65</sup> exchange-correlation functional in conjunction with the density-dependent dispersion correction dDsC.<sup>66,67</sup> The Brillouin zone of 2 x 2 x 2 supercells with stoichiometry Ca<sub>80</sub>P<sub>48</sub>O<sub>192</sub>F<sub>16</sub> (FAp has two formula units in the primitive cell) and cell vector lengths of about 18.8 Å and 13.7 Å for in-plane and out-of-plane directions was probed at the Gamma point. A Fermi-smearing of 0.026 eV (~300 K) was applied, but had no impact on the electronic energy for this insulating material. During the self-consistent field procedure, the wave functions were optimized to an energy-change below 10<sup>-6</sup> eV. The numerical settings for the fast Fourier transform grids was set to “accurate” and the real-space projector operators were determined via the recommended automatic optimization scheme. For geometry optimizations all atoms and the unit cell were fully relaxed until the maximum force was below 0.05 eV/Å. The atomic Hirshfeld charges, which are known to depend little on the electronic structure and the basis set and are well suited for assessing charge-transfers,<sup>68</sup> have been taken from the standard output of the dDsC dispersion correction. The maximum error in the interatomic distances of FAp calculated with VASP relative to the diffraction values<sup>19</sup> is 1.15% (Table S1).

**Samples and EXAFS Spectroscopy.** Two magmatic-hydrothermal FAp were studied by EXAFS spectroscopy, one from Cerro de Mercado near Durango, Mexico,<sup>69</sup> and another from Imilchil, Morocco.<sup>70-72</sup> The Durango FAp was characterized previously by laser ablation inductively coupled plasma mass spectrometry (LA-ICP-MS) and electron probe microanalysis (EPMA).<sup>8</sup> It contains 4,890 mg/kg (ppm) Ce, 3,340 mg/kg La, and 971 mg/kg Pr. The chemical composition of the Imilchil FAp is reported in Table S2. It contains 4,864 mg/kg (ppm) Ce, 2,678 mg/kg La, and 492 mg/kg Pr. The Ce L<sub>3</sub>-edge (5723 eV) HERFD-EXAFS spectra were measured at the European Synchrotron Radiation Facility (ESRF), at room temperature on ID24-DCM and at 8-12 K on FAME-UHD (BM16) for the Durango FAp, and at room temperature on BM20 for the Imilchil FAp. No changes in spectral features were noted during the courses of data collection that would indicate radiation damage. Experimental details are provided in the Supporting Information (SI).

The benefits of HERFD-EXAFS are shown in Figure 2. First, a spectrometer acts both as a low-pass and high-pass filter, eliminating (1) the undesired fluorescence lines at lower and higher energies than that of the Ce L <sub>$\alpha$ 1</sub> line at 4839 eV, here the L <sub>$\alpha$ 1+2</sub> (~4647 eV) and L <sub>$\beta$ 1</sub> (~5038 eV) lines of La and the L <sub>$\alpha$ 1+2</sub> (~5035 eV) of Pr, and (2) the elastic and inelastic rays at higher energy (Figure

2a). As a result, the background signal is lower, and therefore the signal/noise ratio higher and the detection limit lower (Figure 2b). Second, filtering the Ce  $L_{\alpha 1}$  line allows extending the  $k$ -range of EXAFS beyond the La  $L_2$ -edge at 5891 eV and the Pr  $L_3$ -edge at 5964 eV (Figure 2b). This phenomenon was first observed in 2005 for La in LaOCL.<sup>73</sup>

## RESULTS AND DISCUSSION

**DFT.** The first series of calculation focuses on the site preference of Na when Ce occupies the Ca1 site (Ce1) or the Ca2 site (Ce2) site (see polyhedral drawings in Figure 3). This was accomplished by calculating the energy difference of models Ce1-Na1-close and Ce1-Na2-close, on one hand, and Ce2-Na1-close and Ce2-Na2-close, on the other hand, in which Na is positioned in closest proximity to Ce to balance locally the excess of positive charge resulting from the Ce(III) for Ca(II) substitution. Na in the Ce2 site (Na2) is less favorable than in the Ce1 site (Na1) by -0.30 eV/cell when Ce is in the Ca1 site and by -0.18 eV/cell when Ce is in the Ca2 site (Table 1). Na clearly prefers the Ca1 site, in agreement with the structural refinement of synthetic FAp using X-ray diffraction.<sup>42,46</sup> Therefore, Na will be placed in this site in all further calculations. Next, we interrogated the site preference of Ce when the charge excess is compensated by Na (Ce1-Na1-close and Ce2-Na1-close models) or by Si (Ce1-Si-close and Ce2-Si-close models). The Ce1-Na1-close and Ce2-Na1-close models have the same total energy, whereas the Ce2-Si-close model is -0.08 eV/cell lower in energy than the Ce1-Si-close model (Table 1). The finding predicts that Ce prefers the Ca2 site when the charge is balanced by Si, otherwise Ce has no preference when the charge is balanced by Na.

The question that remains to handle is which of the Na or Si substituents provides the lowest energy structure. The answer was obtained from the energy difference between models with the compensating Na and Si cations placed either in closest proximity or far away from Ce ( $d > 9 \text{ \AA}$ ). Configurations that promote local charge balance are always the most energetically competitive structures (Table 1). The gain in energy obtained with the Na short-range surrounding of Ce is -0.09 eV/cell for Ce2-Na1-close and -0.18 eV/cell for Ce1-Na1-close. With Si, the gains are -0.36 eV/cell for Ce2-Si-close and -0.34 eV/cell for Ce1-Si-close. This result shows that the Ce(III) substitution is better charge compensated by Si than by Na, regardless of the site occupancy of Ce. However, given that the Ce2-Si-close model is 0.08 eV/cell more stable than the Ce1-Si-close model, we conclude that Ce substitutes into the Ca2 site with a Si atom next to it at thermodynamic equilibrium.

To be examined now is whether the reason for the Ca2-Si preference is structural or electronic. In FAp, each apex of a  $\text{Ca}_1\text{O}_6$  octahedron shares one oxygen with a  $\text{PO}_4$  tetrahedron (corner-sharing

linkage), whereas four apices of the  $\text{Ca}_2\text{O}_6\text{F}$  polyhedron share one oxygen with a  $\text{PO}_4$  tetrahedron (corner-sharing linkage) and two apices share their oxygen atoms with a single  $\text{PO}_4$  tetrahedron (edge-sharing linkage, Figure 1b). The Ca2-P distance is 3.07 Å across the shared edge and 3.26-3.68 Å across the shared apices. The most favorable configuration from an energetic standpoint is that in which the nearest P atom is replaced with a Si atom, yielding a Ce2-Si distance of 3.09 Å (Ce2-Si-close model, Figure 3a).

Examination of the Ce1-Si-close atomic arrangement shows that one of the three second-nearest O atoms of the Ca1 site at 2.81 Å and bonded to P (1.53 Å) and Ca2 (2.35 Å) in FAp,<sup>19</sup> bonds also the Ce1 atom at 2.54 Å in the optimized structure. The Ce1 coordination is 7 + 2, instead of 6 + 3 for Ca1 in FAp (Figure 3b). Interestingly, the displaced O atom is the one bonded to Si in Ce1-Si-close, instead of P in FAp. As a result of the O displacement, the  $\text{SiO}_4$  tetrahedron shares an edge, and not an apice, with the  $\text{Ce}_1\text{O}_7$  polyhedron at a Ce-Si distance of 3.16 Å (Figures 1b and 3). The relaxation of the atomic positions around the substitutional Ce1 atom allows local balancing of the charge excess of the Ce1 site similarly to the Ce2 site. The charge compensating Si atom is 3.16 - 3.09 = 0.07 Å closer to Ce in the Ca2 site than in the Ca1 site. It is difficult to assess if the reduction in distance is sufficient to account for the 0.08 eV/cell gain in thermodynamic stability for the Ce2 partitioning relative to the Ce1 partitioning (Table 1). Compelling evidence for electronic effects on site preference is presented next.

Electronic effects were addressed with considerations of four divalent Me(II) substituents: Mg(II), Cd(II), Sr(II), and Ba(II). The first parameter examined is the cation size, reasoning that cations smaller than Ca (1.00 Å) should substitute into the smaller Ca2 site ( $\langle d(\text{Ca-O}) \rangle = 2.40$  Å for the six-fold coordination) and larger cations should substitute into the larger Ca1 site ( $\langle d(\text{Ca-O}) \rangle = 2.43$  Å for the six-fold coordination).<sup>19</sup> A regression analysis of the effective ionic radius<sup>74</sup> of the Me(II) substituents against the difference of total energy between the two Ca sites ( $E(\text{Me}_2) - E(\text{Me}_1)$ , Table S3) shows moderate correlation ( $R^2 = 0.79$ , Figure 4a). Furthermore, the Ce2-Si-close model, and to a lesser extent the Ce1-Na-close/Ce2-Na-close models, are distant from the regression line.

Besides their size difference, another distinction of the Ca1 and Ca2 sites is their ionicity. Calculation of atomic charges shows that Ca has a charge of 0.429e in the Ca1 site and 0.475e in the Ca2 site, indicating that the Ca2 site has a higher ionic character. Therefore, we sought whether a correlation existed between the atomic charge and the site preference of the four Me(II) cations (Table S3). Figure 4b shows that the two variables are strongly correlated ( $R^2 = 0.92$ ): the higher the difference of atomic charge of a Me(II) cation between the Ca2 and Ca1 sites, the higher is its preference for the Ca2 site, in line with the higher ionicity of the Ca2 site of FAp. The relationship

between Me(II) partitioning and ionicity of the Ca site is further supported by the fact that Ca is close to the regression line, as this atom was not considered in the linear fit. The Ce2-Si-close and the Ce1-Na-close/Ce2-Na-close models are far apart from the regression line due to the formal trivalent charge of Ce(III) (Figure 4b, Table S4).

Which distinctive electronic property causes a Me(II) cation to acquire a specific atomic charge in a Ca site that eventually determines its affinity for this site? What immediately comes to mind is its ionization energies. Figures 4c and 4d show that the first and second ionization energies are linearly correlated to  $E(\text{Me}2) - E(\text{Me}1)$  ( $R^2 = 0.90$  and  $0.93$ ), thereby backing up our hypothesis. Remarkably, the Ce(III) cation from the Ce2-Si-close model is close to the two regression lines, which indicates that this cation is energetically and electronically well stabilized in the Ca2 site. This finding suggests that ionization energy is the main thermodynamic driving force for the partitioning of Ce(III) among the Ca1 and Ca2 sites in FAp, rather than the  $0.07 \text{ \AA}$  difference in the Ce-Si interatomic distances between the two sites.

**EXAFS Spectroscopy.** The EXAFS spectra and Fourier transforms (FT) of the Durango FAp measured at room and low temperature (Durango-RT, Durango-HeT) and of the Imilchil FAp measured at room temperature (Imilchil-RT) are presented in Figure 5. The Durango spectrum has a higher amplitude at HeT, which is reflected in the FT by a considerable enhancement of the second and third peaks attributed to the dominant contribution of the Ce-P and Ce-Ca atomic pairs (Figure 3). The thermal motion of the Ce-P and Ce-Ca pairs is much more pronounced than that of the Ce-O pair at shorter bond distance (1<sup>st</sup> FT peak). Comparing the imaginary parts of the FT (i.e., oscillatory curves on Figure 5b), we find that they are superimposed at RT and HeT, which excludes a temperature-dependence of the bond distances. Thus, the Ce-P and Ce-Ca motions are harmonic with respect to the equilibrium position of the Ce, P, and Ca atoms. Large temperature-dependence of the cation shells has been observed previously for the Mn-Mn and Pb-Mn pairs in phyllosulfates,<sup>75,76</sup> but the physical motion was anharmonic, perhaps as a result of the layered structure of phyllosulfates and the implication of the two-dimensionality on atomic vibrational modes. The Durango-RT and Imilchil-RT spectra and FT are much alike in their amplitude and phase, indicating that Ce has similar bonding environment in the two FAp (Figure 5c,d). The site occupancy of Ce and the charge compensation mechanism are examined next with the better quality Durango-RT data.

Apatite is chemically simple, comprising only two cations, Ca and P, yet its structure is complex with two Ca sites, each surrounded by several and partly overlapping P and Ca shells between  $3.0 \text{ \AA}$  and  $4.1 \text{ \AA}$  (Figure 3). Least-squares fitting the average distances and coordination numbers ( $CN$ ) of the 4-5 nearest O, P, and Ca shells around Ce, and the mean-square displacement of the constitutive



atoms in each of them (Debye-Waller term,  $\sigma$ ), gives multiple optima in the parameter space. Additional complications arise from multiple scattering events of the photoelectron that are not negligible at  $R_{\text{eff}} \geq 3.6 \text{ \AA}$ , where  $R_{\text{eff}}$  is the half-path length of the scattering path (Figure S1). It is critical not to overparametrize any of the models tested, and good quality fits that are both statistically robust and structurally meaningful are difficult to obtain. In our hands, only the average distance of the first Ce-O shell can be obtained reliably, and can be used to distinguish unequivocally between the Ce1 and the Ce2 sites. In the DFT model, the 9 O atoms of the Ce1 site are at 2.549  $\text{\AA}$  on average ( $\sigma = 0.078 \text{ \AA}$ ) and the 6 nearest O atoms of the Ce2 site are at 2.428  $\text{\AA}$  on average ( $\sigma = 0.035 \text{ \AA}$ ) (Figure 3). The best-fit EXAFS distance is 2.43  $\text{\AA}$ , identical to the DFT Ce2-O distance. Best-fit coordination numbers are highly correlated to  $\sigma$ , and therefore do not yield reliable results. In addition, the distribution of Ce-O distances is asymmetric in the lowest energy Ce2-Si-close model (Figure 3b), and its EXAFS analysis using a standard Gaussian-type distribution of interatomic distances will exclude its full distribution and lose O atoms ( $CN_{\text{eff}} < CN_{\text{str}}$ ).

A holistic alternative to determination of the bonding environment of Ce in apatite consists of comparing data with DFT-derived EXAFS models. The EXAFS spectra of five lowest energy DFT structures were generated using the FEFF 8.2 code.<sup>77</sup> The degree to which a theoretical spectrum resembles the Durango-RT FAp spectrum was evaluated with the normalized sum-of-squares ( $NSS = \sum(k^2\chi_{\text{exp}} - k^2\chi_{\text{fit}})^2 / \sum(k^2\chi_{\text{exp}})^2$ ) parameter.

The theoretical Ce2-Si-close and Ce1-Si-close spectra and FT are compared to the Durango-RT data in Figure 6. The Ce2-Si-close model well replicates the phase and amplitude of the experimental wave and FT (Figure 6a,b,  $NSS = 0.05$ ). This good match was obtained by optimizing only six EXAFS parameters: the threshold Fermi level of the photoelectron ( $\Delta E = (k/0.512)^2$ ) for the wave phase, four Debye-Waller (DB) values for single-scattering events between the central atom (Ce) and the O, P1, P2, and Ca shells up to 4.15  $\text{\AA}$  (Figure 3b), and one DB value for multiple scattering (MS) events ( $\sigma(\text{O}) = 0.12 \text{ \AA}$ ,  $\sigma(\text{Si,P1}) = 0.07 \text{ \AA}$ ,  $\sigma(\text{P2}) = 0.12 \text{ \AA}$ ,  $\sigma(\text{Ca}) = 0.11 \text{ \AA}$ ,  $\sigma(\text{MS}) = 0.17 \text{ \AA}$ ). Lower DB values for the cation shells were obtained with Durango-HeT ( $\sigma(\text{P1}) = 0.04 \text{ \AA}$ ,  $\sigma(\text{P2}) = 0.09 \text{ \AA}$ ,  $\sigma(\text{Ca}) = 0.08 \text{ \AA}$ ), consistent with the reduction of atomic motion at low temperature. In contrast to Ce2-Si-close, Ce1-Si-close failed to replicate the experimental wave phase and amplitude, yielding considerable disparity between measurement and prediction ( $NSS = 0.30$ ) (Figure 6c,d). The theoretical wave is shifted to lower wavevector values ( $k$ ) in the 3.0-6.0  $\text{\AA}^{-1}$   $k$ -range, and this displacement can be used to estimate the maximum amount, or detection limit, of Ce in the Ca1 site.

The Ce1 component makes up <10% of Ce total, on the basis of linear-combinations of the Ce1-Si-close and Ce2-Si-close spectra.

EXAFS ordinarily fails to distinguish between P and Si atomic neighbors for their photoelectron scattering amplitudes are similar ( $Z = 15$  and  $14$ ), and it has little sensitivity to the Na shell at  $d = 3.46 \text{ \AA}$  (Ce1-Na1-close) and  $3.98 \text{ \AA}$  (Ce2-Na1-close, distance histograms of Figures S3a, S4a). Therefore, the Ce-Na and Ce-Si models are virtually impossible to differentiate with the usual shell-by-shell least-squares fitting analysis. Using a whole-structure approach, like here, allows differentiation of the two models, because the Si for P, and the Na for Ca, substitutions modify more than just two discrete atomic shells, but a larger portion of the short and medium range orders around the Ce impurity. Si for P distinction is illustrated in Figure S2 with the poorer match to data of Ce2-Si-far ( $NSS = 0.06$ ) than Ce2-Si-close ( $NSS = 0.05$ ), indicating that the Ce(III) for Ca substitution is paired with a Si for P substitution at short distance in the apatite structure. Ce2-Si-far has longer Ca-Ca pairs than Ce2-Si-close (histograms of Figures 3b and S2a), which are seen in the FT of Ce2-Si-far by a shift to longer distance of the Ce-Ca peak (Figure S2c). Similarly, Ce2-Na1-close ( $NSS = 0.08$ ) and Ce1-Na1-close ( $NSS = 0.28$ ) have EXAFS and FT features that differentiate them from other models (Figures S3b,c and S4b,c).

## CONCLUDING REMARKS

The results of this study lead us to revisit a previous suggestion that the distribution of a substituent between the Ca1 and Ca2 sites is a marker for the enrichment mechanism, which distinguishes secondary diffusion process through ion-exchange from authigenic precipitation or recrystallization.<sup>78</sup> The idea was proposed after a study on the incorporation of Cd in sedimentary FAp using EXAFS spectroscopy.<sup>22</sup> It was found that Cd occupied the Ca2 site in the sedimentary FAp and was substituted in the Ca1 site in a synthetic Cd-doped hydrothermal reference. It was hypothesized that the Cd enrichment in conditions of the Earth's surface occurred by a diffusion mechanism into the apatite nanopores or along the external surfaces of the mineral grains, because the Ca2 sites are exposed to circulating fluids (Figure 1a). An authigenic formation process was discounted. This hypothesis is supported here and in the DFT study of Tamm and Peld<sup>79</sup> by the energetic preference of Cd for the Ca1 site (Figure 4). Thus, substitution of Cd in the Ca2 site of the natural sedimentary FAp was metastable and, consequently, the site occupancy of Cd had registered the geochemical process that the FAp had undergone. Figure 4 shows that Mg may also be used as a structural probe for past geochemical processes, whereas Ce clearly cannot.

This study presents the first demonstration that Ce(III) is incorporated in the Ca2 site of natural magmatic/hydrothermal FAp and shows that the excess of positive charge resulting from the Ce(III) for Ca(II) substitution is balanced by a Si for P substitution at short distance. It remains to be seen whether this finding can be extended to biogenic FAp, such as those occurring in marine sediments, which are prospective resources for REY.<sup>80</sup> Bioapatites from ocean-floor sediments have similar Ce L<sub>3</sub>-edge X-ray absorption near-edge structure (XANES) spectra as magmatic/hydrothermal FAp,<sup>8</sup> indicating that Ce is also trivalent. But XANES spectroscopy does not inform on the chemical form of REY, and therefore on ore enrichment processes which can be different for high-temperature magmatic/hydrothermal and low-temperature sedimentary deposits. From a broader earth sciences perspective, the new analytical possibilities offered by the combination of HERFD-EXAFS and DFT is perhaps the most exciting finding of this work, as it opens up the possibility to differentiate metal enrichment by ion-exchange and precipitation process.

## **ASSOCIATED CONTENT**

### **Supporting Information**

EXAFS spectroscopy and LA-ICP-MS experimental methods, optimized (VASP) versus experimental (XRD) interatomic distances for FAp, total energies and atomic charges of the Me(II)-FAp models, atomic charges of the Ce-Si and Ce-Na models, chemical analysis of the Imilchil apatite, Cartesian coordinates of the Ce1-Na1-close, Ce2-Na1-close, Ce1-Si-close, Ce2-Si-close, and Ce2-Si-far models, EXAFS spectra (PDF).

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**Table 1. Total Energies of the Ce-Si and Ce-Na Models**

<b>Model</b>	<b>Total energy <i>E</i>(eV/cell)</b>	<b>Energy difference</b>
<i>Na prefers the Ca1 site, regardless of the Ce site</i>		
Ce2-Na1-close	-2447.40	-0.18
Ce2-Na2-close	-2447.22	-
Ce1-Na1-close	-2447.40	-0.30

Ce1-Na2-close	-2447.10	-
<i>Ce has no site preference in the Ce-Na model</i>		
Ce2-Na1-close	-2447.40	0.00
Ce1-Na1-close	-2447.40	
<i>Ce prefers the Ca2 site in the Ce-Si model</i>		
Ce2-Si-close	-2451.30	-0.08
Ce1-Si-close	-2451.22	
<i>Ce2-Na1 charge compensation at short distance</i>		
Ce2-Na1-close	-2447.40	-0.09
Ce2-Na1-far	-2447.31	-
<i>Ce1-Na1 charge compensation at short distance</i>		
Ce1-Na1-close	-2447.40	-0.18
Ce1-Na1-far	-2447.22	-
<i>Ce2-Si charge compensation at short distance</i>		
Ce2-Si-close	-2451.30	-0.36
Ce2-Si-far	-2450.94	
<i>Ce1-Si charge compensation at short distance</i>		
Ce1-Si-close	-2451.22	-0.34
Ce1-Si-far	-2450.88	-

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## FIGURE CAPTION

**Figure 1.** (a) Structure of FAp projected in the  $ab$  plane.<sup>19</sup> (b)  $\text{CaO}_6\text{-PO}_4$  and  $\text{CaO}_6\text{F-PO}_4$  linkages at the Ca1 and Ca2 sites of FAp, and  $\text{CeO}_7\text{-(PO}_4\text{,SiO}_4\text{)}$  linkage of the Ce1-Si-close model. The F atom is located in the middle of the tunnel (a) and is unshared (b).

**Figure 2.** (a) Monte Carlo simulation of the XRF spectrum for the Durango FAp calculated with the resolution of a SDD and an excitation energy of 6 keV using the Xraylib database.<sup>81,82</sup> (b) Ce  $L_3$ -edge X-ray absorption spectrum of the Durango FAp measured with a SDD and a five-crystal spectrometer (HERFD-XAS).<sup>53</sup>

**Figure 3.** Polyedral sketch of the DFT local structure of Ce in the Ca1 and Ca2 sites (a), and population histogram of the Ce-(O,F,Si,P,Ca) distances (b,c). (a) Brown, Ca in the Ca1 site; green,

Ca in the Ca2 site; cyan, Ce in the Ca1/Ca2 site; purple, P corner-linked to Ca2/Ce2; blue, Si edge-linked to Ce2/Ce1; red, oxygen; light green, F. (b) Ce-2-Si-close DFT model. Average distance of the 6 nearest O,F = 2.428 Å,  $\sigma = 0.035$  Å. (c) Ce1-Si-close DFT model. Average distance of the 9 O = 2.549 Å,  $\sigma = 0.078$  Å. Number of atoms on the histograms are counted in intervals of 0.05 Å.

**Figure 4.** Dependence of the relative stability of substituents in the Ca1 and Ca2 site of FAp on their electronic properties. (a) Effect of the effective ionic radius expressed as the difference between the Me(II) and Ca(II) radii ( $\Delta IR$ ). (b) Effect of the atomic charge expressed as the difference of its value for Me(II) in the Ca2 and Ca1 sites ( $\Delta AC$ ). (c,d) Effect of the first (c) and second (d) ionization energies expressed as the difference between the Me(II) and Ca(II) energies ( $\Delta IE$ ).

**Figure 5.** Experimental Ce L<sub>3</sub>-edge EXAFS spectra and Fourier transforms (magnitude and imaginary parts) of the Durango FAp measured at room temperature (Durango-RT) and at 8-12 K (Durango-HeT) (a,b), and of Durango-RT and the Imilchil FAp measured at room temperature (Imilchil-RT) (c,d).

**Figure 6.** Experimental and DFT-derived theoretical Ce L<sub>3</sub>-edge EXAFS spectra and Fourier transforms (magnitude and imaginary parts) of FAp. (a,b) Durango FAp measured at room temperature (Durango-RT) and Ce2-Si-close spectra. (c,d) Durango-RT and Ce1-Si-close spectra.

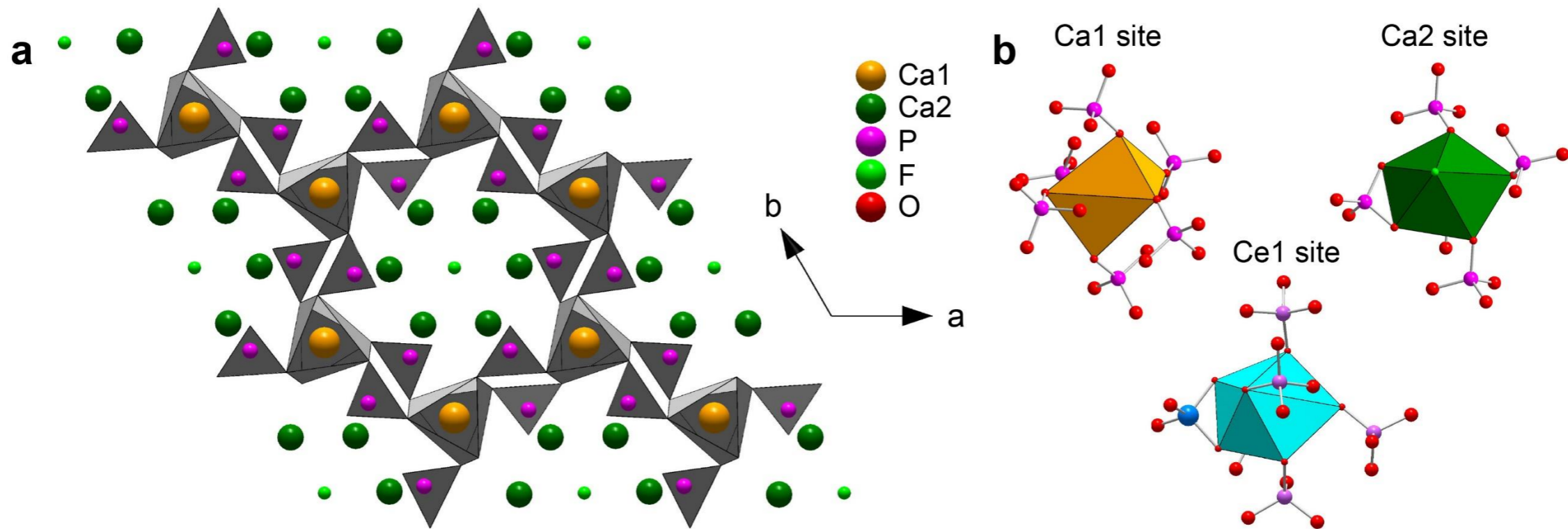
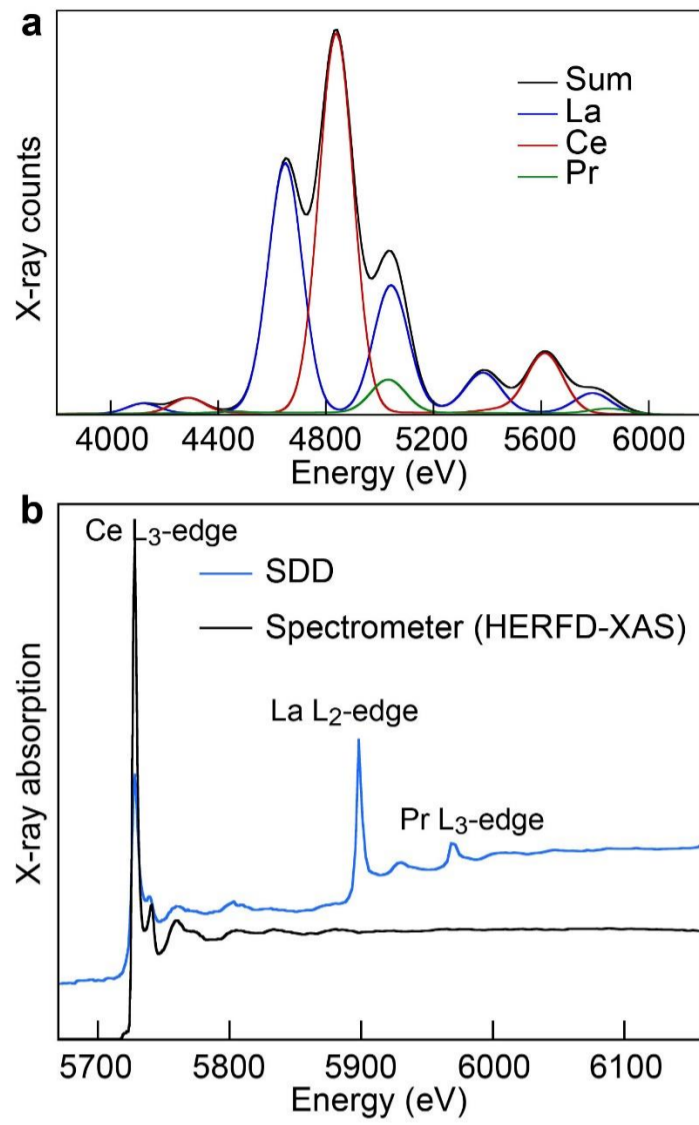
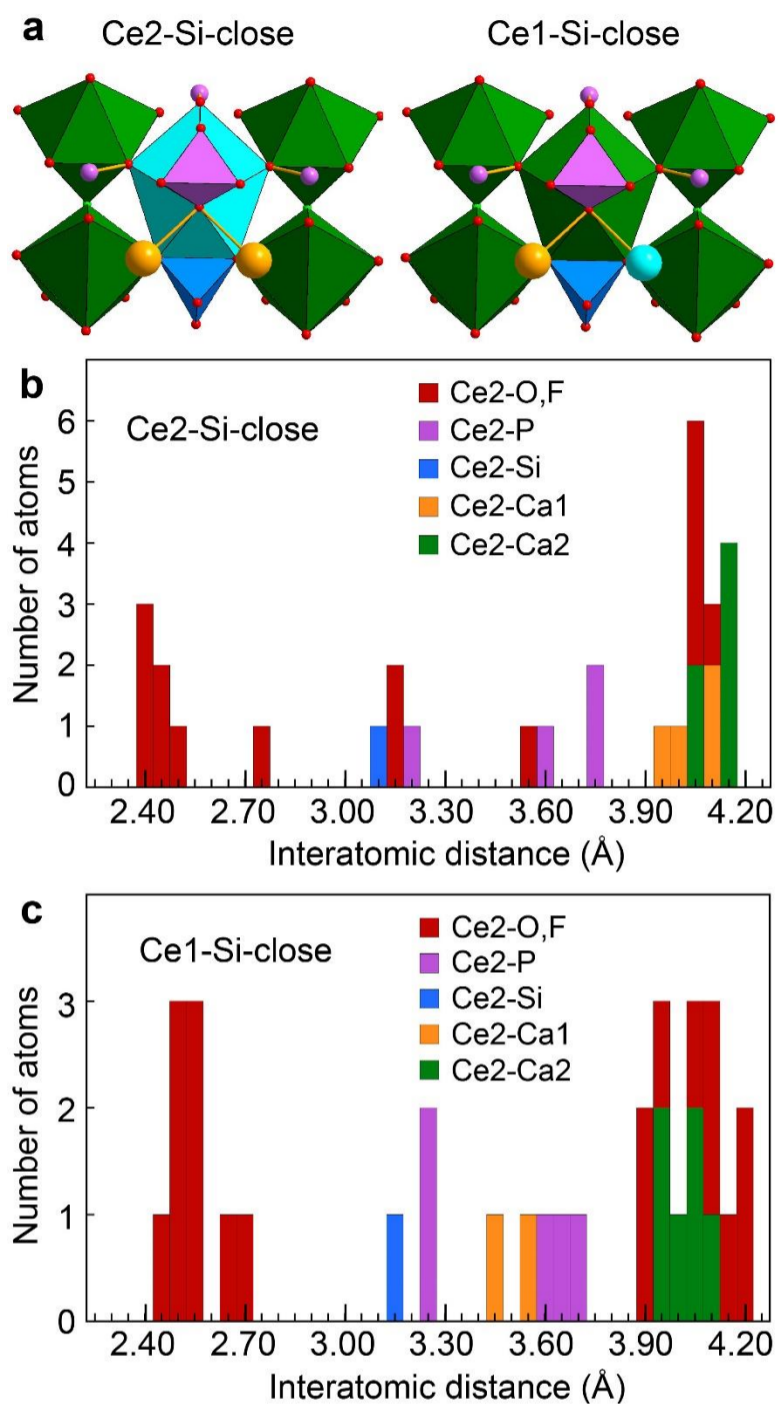


Figure 1



**Figure 2**



**Figure 3**

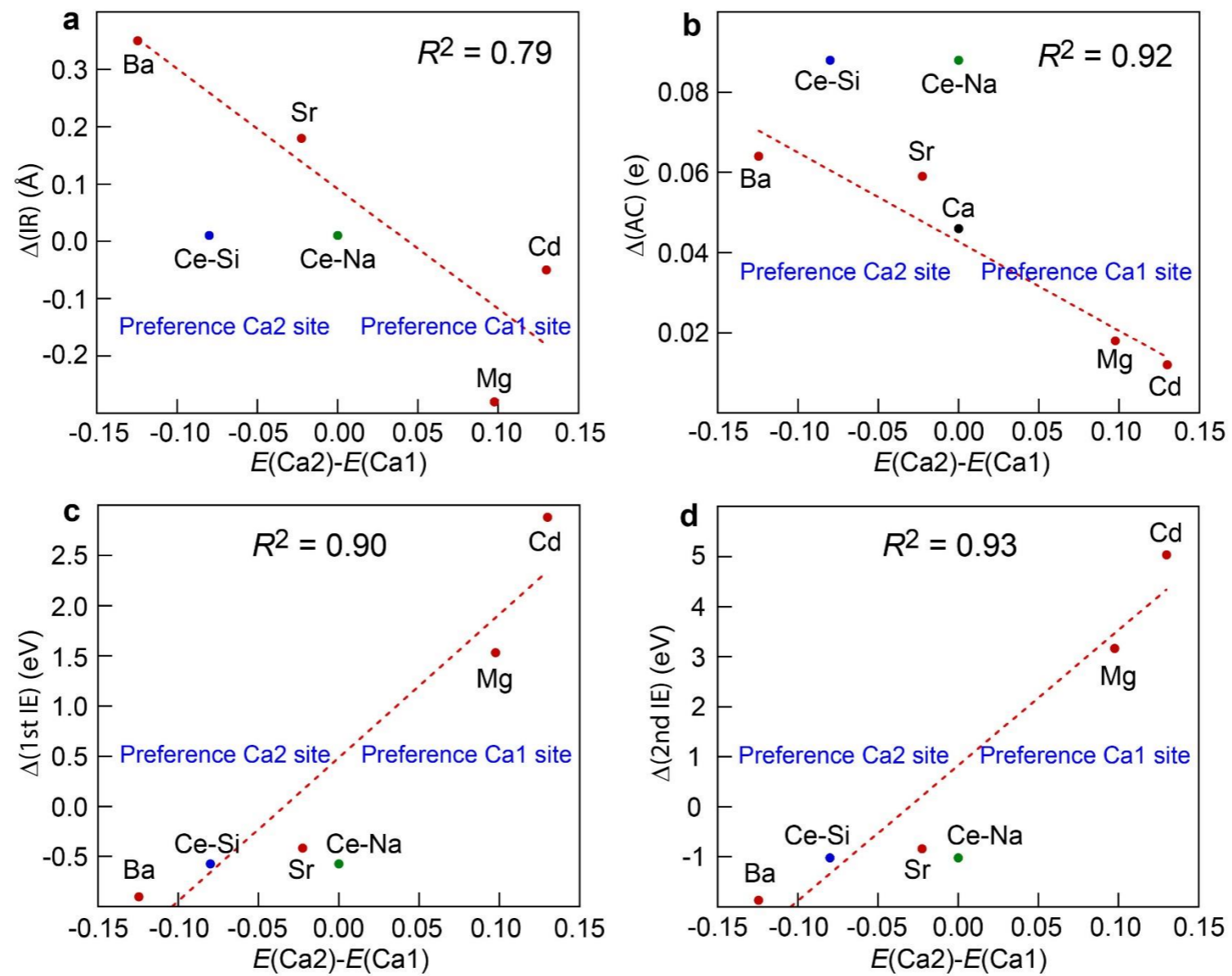


Figure 4



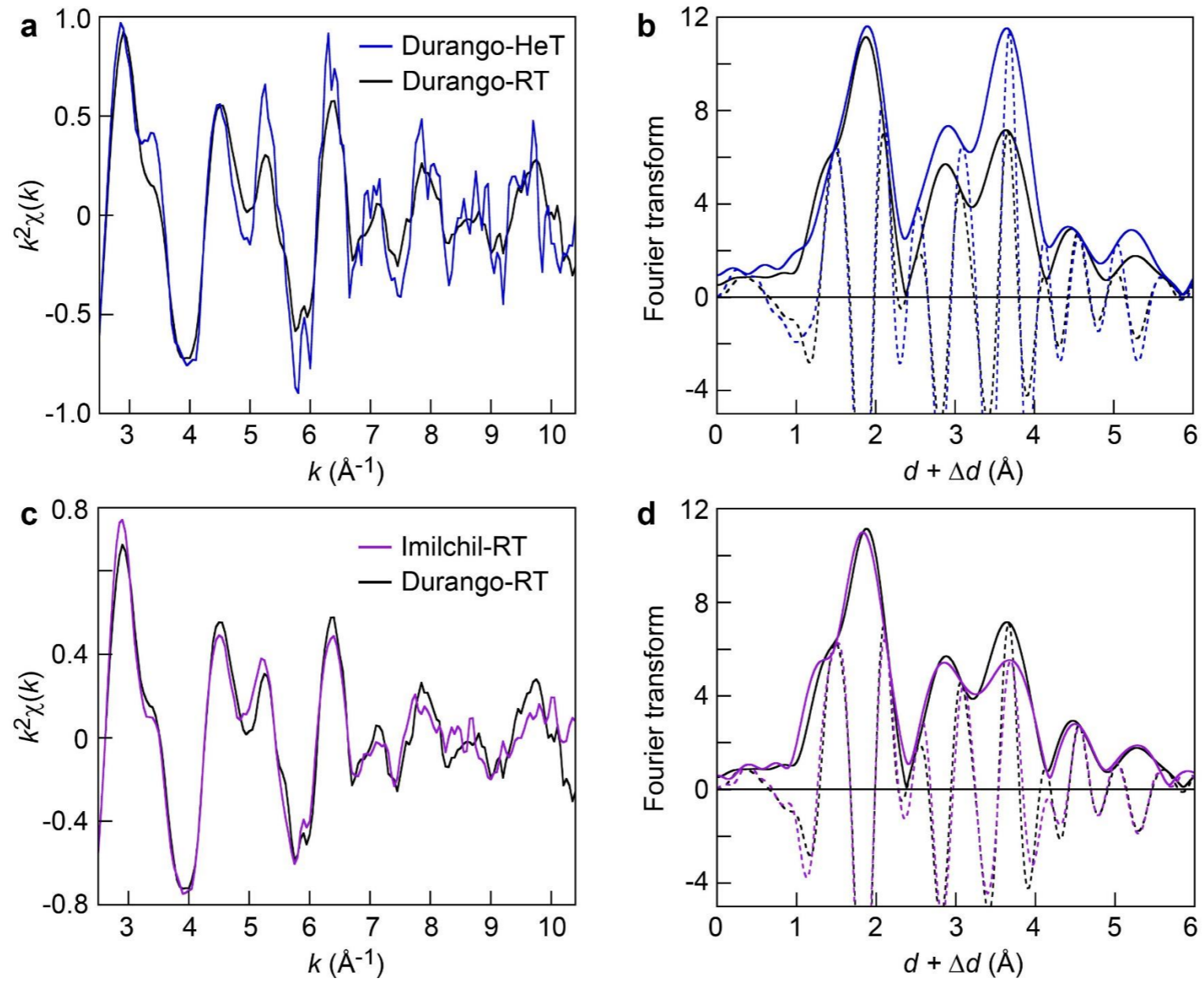


Figure 5

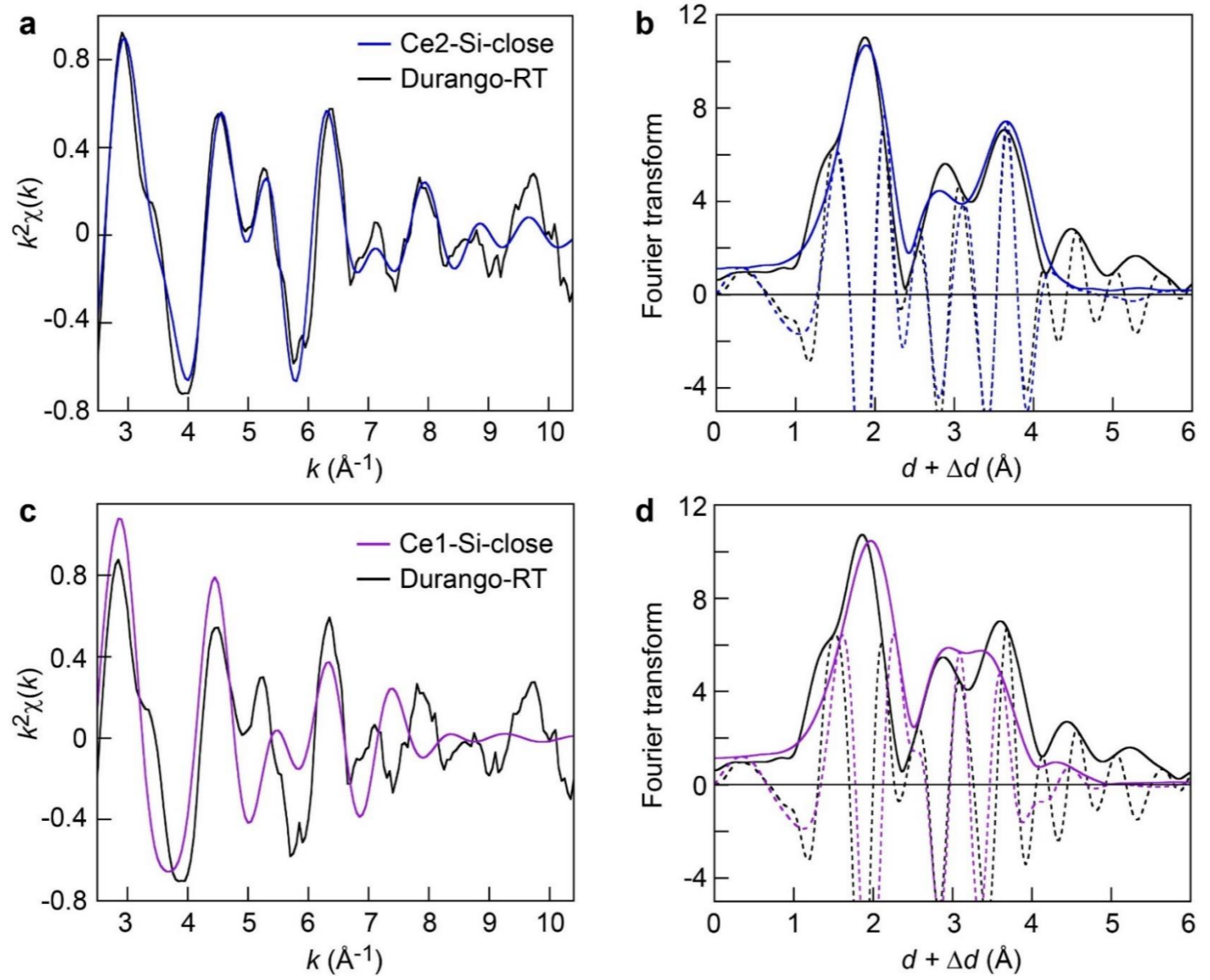


Figure 6

## Supporting Information

### Revealing the incorporation of cerium in fluorapatite

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### EXAFS Spectroscopy

**Measurement of the Durango FAp at RT on ID24-DCM.** The incident photon energy was selected using the Si(111) reflection from a liquid-nitrogen-cooled double-crystal monochromator, and the rejection of higher harmonics was achieved with two Si mirrors at an incidence angle of 4 mrad (7 keV cutoff). The beam size on the sample was 80 (H) × 80 (V) μm<sup>2</sup>, and the incident X-ray flux was ~5 × 10<sup>12</sup> photons per s at the sample position. The Ce L<sub>α1</sub> (3d<sub>5/2</sub> → 2p<sub>3/2</sub>) fluorescence line (4839 eV) was selected using the 400 reflection of five Si crystals of 100 mm diameter, spherically bent to a radius of 0.5 m and aligned at a Bragg angle of 70.6° in a vertical Rowland geometry.<sup>1</sup> The diffracted intensity was measured with a Si drift diode detector (SDD) in single photon counting mode. Forty-four spectra of 7 min were acquired in continuous scan mode, and averaged to improve statistics.

**Measurement of the Durango FAp at 8-12 K on FAME-UHD (BM16).** The incident photon energy was selected using the Si(220) reflection from a liquid-nitrogen-cooled double-crystal monochromator, and the rejection of higher harmonics was achieved with two Rh-coated cylindrical mirrors at an incidence angle of 6.2 mrad (9.5 keV cutoff). The beam size on the sample was 300 (H) × 100 (V) μm<sup>2</sup>, and the incident X-ray flux was ~10<sup>11</sup> photons per s at the sample position. The Ce L<sub>α1</sub> fluorescence line was selected using the 331 reflection of five Ge crystals of 100 mm diameter, spherically bent to a radius of 1.0 m and aligned at a Bragg angle of 80.7° in a vertical Rowland geometry.<sup>2</sup> The diffracted intensity was measured with a Si drift diode detector (SDD) in single photon counting mode. Forty-seven spectra of 31 min were acquired and averaged to improve statistics. Measurements were performed using a helium flow cryostat.

**Measurement of the Imilchil FAp on BM20.** The incident photon energy was selected using the Si(111) reflection from a liquid-nitrogen-cooled double-crystal monochromator, and the rejection of higher harmonics was achieved by two Si mirrors at an incidence angle of 2.5 mrad. The beam size on the sample was 400 (H) × 150 (V) μm<sup>2</sup>, and the incident X-ray flux was ~10<sup>10</sup> photons per s at the sample position. The Ce L<sub>α1</sub> fluorescence line was selected using the 331 reflection of five Ge crystals of 100 mm diameter, spherically bent to a radius of 0.5 m and aligned at a Bragg angle of 80.7° in a vertical Rowland geometry.<sup>1,3</sup> Eighty-three spectra of 20 min were acquired and averaged to improve statistics.

EXAFS spectra collected on different beamlines were intercalibrated on the same energy scale. Data were reduced with the Athena software.<sup>4</sup>

### LA-ICP-MS analysis

Laser sampling was performed with a 193 nm GeoLas Pro ArF Excimer laser (Microlas®, Göttingen, Germany) equipped with beam homogenization optics. An ablation crater of 60 μm was performed by focusing the beam on the sample surface with a constant fluence of 10 mJ/cm<sup>2</sup> and constant repetition rate of 5 Hz. Helium was used as a carrier gas to transport the laser-generated aerosols from the ablation cell to the ICP-MS analyzer. Argon as make-up gas was mixed with the carrier gas before entering the ICP torch. A typical flow rate of 0.5 l.min<sup>-1</sup> for He and 0.96 l.min<sup>-1</sup> for Ar was used. These values were optimized with the analysis of NIST610 SRM silicate glass. Ablated material was analyzed by an Agilent 7500c Quadrupole ICP-MS (Agilent®, Santa Clara, California) equipped with an Octopole Reaction

System with enhanced sensitivity optional lenses (Cs type, Agilent). Peak hopping ablation mode began 20 s after the beginning of the signal acquisition, in order to sample the background signal before ablation (used for data reduction), and was stopped after 200 pulses (40 s). External standard calibration was performed with the NIST610 SRM reference. Data reduction was carried out using Iolite software (version 4),<sup>5</sup> following the standard methods of Longerich et al., (1996).<sup>6</sup>

### Supplementary References

- (1) Rovezzi, M.; Lapras, C.; Manceau, A.; Glatzel, P.; Verbeni, R. High Energy-Resolution x-Ray Spectroscopy at Ultra-High Dilution with Spherically Bent Crystal Analyzers of 0.5 m Radius. *Rev. Sci. Instr.* **2017**, *88*, 013108.
- (2) Proux, O.; Lahera, E.; Del Net, W.; Kieffer, I.; Rovezzi, M.; Testemale, D.; Irar, M.; Thomas, S.; Aguilar-Tapia, A.; Bazarkina, E.; Prat, A.; Tella, M.; Auffan, M.; Rose, J.; Hazemann, J. L. High-Energy Resolution Fluorescence Detected X-Ray Absorption Spectroscopy: A Powerful New Structural Tool in Environmental Biogeochemistry Sciences. *J. Environ. Qual.* **2017**, *46*, 1146–1157.
- (3) Kvashnina, K.; Scheinost, A. A Johann-Type X-Ray Emission Spectrometer at the Rossendorf Beamline. *J. Synchrotron Rad.* **2016**, *23*, 836–841.
- (4) Ravel, B.; Newville, M. ATHENA, ARTEMIS, HEPHAESTUS: Data Analysis for X-Ray Absorption Spectroscopy Using IFEFFIT. *J. Synchrotron Radiat.* **2005**, *12*, 537–541.
- (5) Paton, C.; Hellstrom, J.; Paul, B.; Woodhead, J.; Hergt, J. Iolite: Freeware for the Visualisation and Processing of Mass Spectrometric Data. *J. Anal. At. Spectrom.* **2011**, *26*, 2508–2518.
- (6) Longerich; Jackson, S. E.; Gunther, D. Laser Ablation Inductively Coupled Plasma Mass Spectrometric Transient Signal Data Acquisition and Analyte Concentration Calculation. *J. Anal. At. Spectrom.* **1996**, *11*, 899–904.

**Table S1. Optimized (VASP) versus experimental (XRD)<sup>a</sup> interatomic distances (Å) for FAp**

Ca1 site	VASP	XRD	%Diff
3O1	2.397	2.399	0.08
3O2	2.446	2.457	0.45
3O3	2.806	2.807	0.04
3P	3.206	3.207	0.03
Ca1	3.427	3.425	-0.06
Ca1	3.447	3.453	0.17
3P	3.586	3.585	-0.03
3O3	3.931	3.916	-0.38
<b>Ca2 site</b>			
F	2.305	2.311	0.26
2O3	2.341	2.348	0.30
O2	2.372	2.374	0.08
2O3	2.493	2.501	0.32
O1	2.731	2.700	-1.15
P	3.053	3.075	0.72
P	3.285	3.265	-0.61
2O3	3.33	3.347	0.51
O1	3.493	3.493	0.00
P	3.5	3.494	-0.17
2P	3.684	3.68	-0.11
2Ca1	3.974	3.961	-0.33

<sup>a</sup>Hughes et al. *Am. Miner.*, 1989, 74, 870-876.

**Table S2. Chemical analysis of the Imilchil FAp in mg/kg (ppm).**

Technique	Element	Point 1	Point 2	Point 3	Point 4	Average	STD
ICP-AES	Na	1332	1353	1076	1531	1323	187
ICP-AES	Mg	92	93	70	88	85.8	10.9
ICP-AES	K	10	2	8	4	6.0	3.6

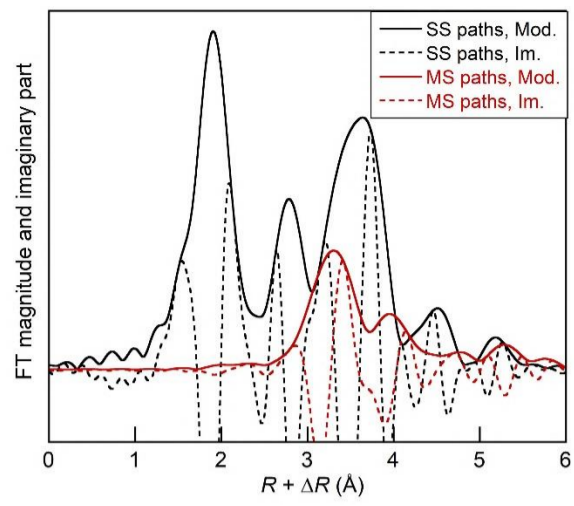
ICP-AES	Sc	0.2	0.2	0.3	0.3	0.3	0.1
ICP-AES	Ti	18	24	22	23	22	3
ICP-AES	V	3	3	3	4	3.3	0.4
ICP-AES	Fe	613	628	606	694	635	40
ICP-AES	Co	0.2	0.1	0.2	0.1	0.2	0.1
ICP-AES	Ni	0.5	3.2	3.4	0.2	1.8	1.7
ICP-AES	Cu	0.4	0.1	0.1	0.0	0.2	0.2
ICP-AES	Zn	0.1	0.0	0.1	0.2	0.1	0.1
ICP-AES	As	12	12	11	14	12.4	1.3
ICP-AES	Sn	4	4	5	6	4.8	0.6
ICP-AES	Ba	0.8	0.9	0.9	1.4	1.0	0.3
LA-ICP-MS	Mn	220	211	234	233	225	11
LA-ICP-MS	Sr	672	668	664	668	668	3
LA-ICP-MS	Y	1226	1280	1219	1289	1254	36
LA-ICP-MS	La	2307	2366	3006	3033	2678	395
LA-ICP-MS	Ce	4764	4617	5068	5006	4864	210
LA-ICP-MS	Pr	487	491	500	490	492	5
LA-ICP-MS	Nd	1917	1964	1898	1917	1924	28
LA-ICP-MS	Sm	397	409	382	389	395	12
LA-ICP-MS	Eu	41	41	38	39	40.0	1.5
LA-ICP-MS	Gd	389	401	370	380	385	13
LA-ICP-MS	Tb	49	50	47	48	48.3	1.2
LA-ICP-MS	Dy	264	274	259	269	266	7
LA-ICP-MS	Ho	47	49	47	49	47.8	1.0
LA-ICP-MS	Er	108	113	110	114	111	3
LA-ICP-MS	Tm	12	12	12	13	12.2	0.3
LA-ICP-MS	Yb	57	59	61	62	59.7	2.2
LA-ICP-MS	Lu	5	6	6	6	5.7	0.4
LA-ICP-MS	Th	59	61	82	88	72.7	14.5
LA-ICP-MS	U	7	7	9	9	7.8	1.0

**Table S3. Total energies and atomic charges of the Me(II)-FAP models**

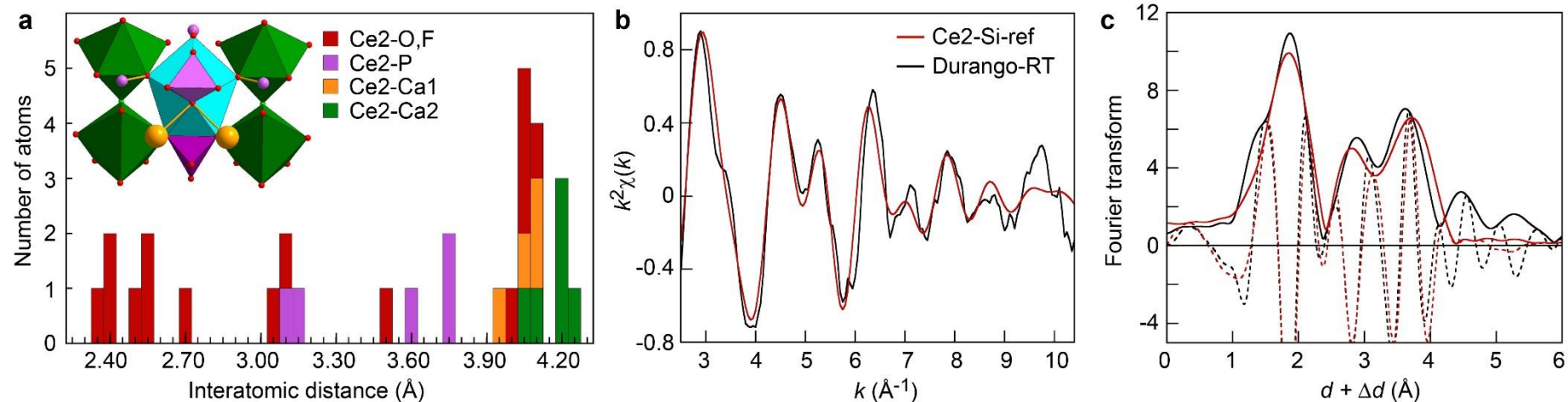
<b>Model</b>	<b>Total energy E (eV/cell)</b>	<b>Energy difference</b>	<b>Atomic charge AC (e-)</b>	<b>AC difference</b>
Mg2	-2444.91	0.10	0.64	0.02
Mg1	-2445.01	-	0.62	-
Cd2	-2441.03	0.13	0.82	0.01
Cd1	-2441.16	-	0.81	-
Sr2	-2446.67	-0.02	0.49	0.06
Sr1	-2446.65	-	0.43	-
Ba2	-2446.75	-0.12	0.44	0.06
Ba1	-2446.63	-	0.38	-

**Table S4. Atomic charges of the Ce-Si and Ce-Na models**

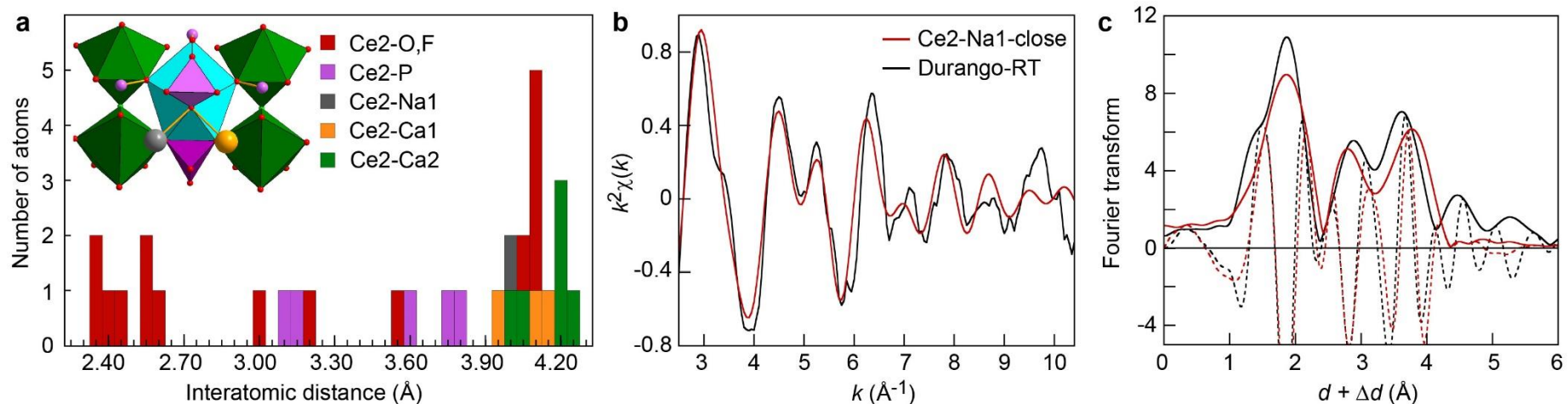
<b>Model</b>	<b>Atomic charge AC (e-)</b>	<b>AC difference</b>
Ce2-Si-close	0.77	0.08
Ce1-Si-close	0.69	-
Ce2-Na1-close	0.78	0.09
Ce1-Na1-close	0.69	-



**Fig. S1.** Fourier transforms (modulus and imaginary parts) of the EXAFS single (SS) and multiple (MS) scattering paths calculated with FEFF 8.2.



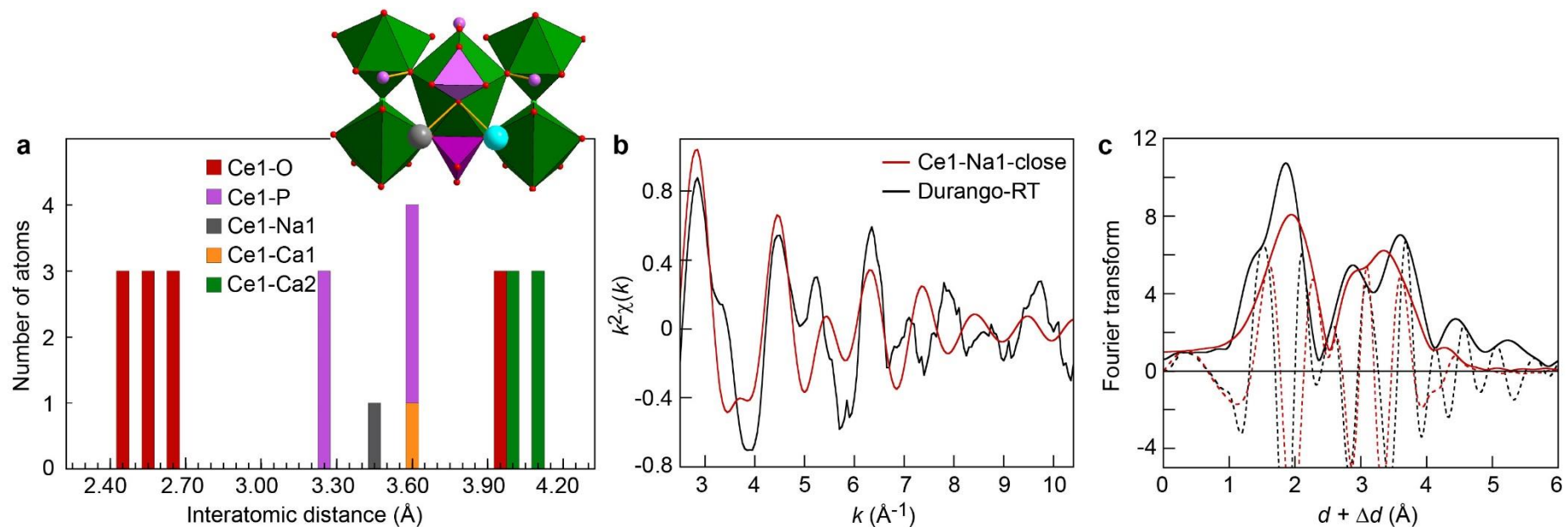
**Fig. S2.** (a) Population histogram of the Ce-(O,F,P,Ca) distances for the Ce2-Si-ref model and polyedral sketch of the local structure of Ce in the Ca2 site. Brown, Ca in the Ca1 site; green, Ca2 polyhedron; cyan, Ce2 polyhedron; dark purple, P tetrahedron edge-linked to Ce2 polyhedron and corner-linked to Ca1, light purple, P tetrahedron corner-linked to Ce2 polyhedron; red, oxygen; light green, F. (b,c) EXAFS spectra and Fourier transforms (magnitude and imaginary parts) of Durango-RT and the Ce2-Si-ref model.



**Fig. S3.** (a) Population histogram of the Ce-(O,F,P, Na, Ca) distances for the Ce2-Na1-close model and polyedral sketch of the local structure of Ce in the Ca2 site with Na at short distance. Brown, Ca in the Ca1 site; green, Ca2 polyhedron; cyan, Ce2 polyhedron; grey, Na in the Ca1 site



(Na1); dark purple, P tetrahedron edge-linked to Ce2 polyhedron and corner-linked to Ca1 and Na1, light purple, P tetrahedron corner-linked to Ce2 polyhedron; red, oxygen; light green, F (b,c) EXAFS spectra and Fourier transforms (magnitude and imaginary parts) of Durango-RT and the Ce2-Na1-close model.



**Fig. S4.** (a) Population histogram of the Ce-(O,F,P, Na, Ca) distances for the Ce1-Na1-close model and polyedral sketch of the local structure of Ce in the Ca1 site with Na at short distance. Green, Ca2 polyhedron; cyan, Ce in the Ca1 site (Ce1); grey: Na in the Ca1 site (Na1); dark purple, P tetrahedron edge-linked to Ca2 polyhedron and corner-linked to Ce1 and Na1, light purple, P tetrahedron corner-linked to Ca2 polyhedron; red, oxygen; light green, F (b,c) EXAFS spectra and Fourier transforms (magnitude and imaginary parts) of Durango-RT and the Ce1-Na1-close model.



**Cartesian coordinates ( $a = 100 \text{ \AA}$ ,  $b = 100 \text{ \AA}$ ,  $c = 100 \text{ \AA}$ ) - Ce1-Na1-close model**

58	9.392709952	5.422883296	6.839256000
11	9.392709952	5.422883296	3.379071143
20	5.877256564	6.156736541	5.096284453
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20	4.697584079	2.696475410	6.862156680
20	14.093982154	2.723370343	-0.004956409
20	14.101411952	2.719989430	6.862156680
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20	0.000000149	10.845767321	6.871653625
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20	0.003369815	5.419532776	13.733762578
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20	-4.700941526	13.555965644	13.733762578
20	4.698092822	13.550323170	6.853543179
20	4.697571705	13.561802339	13.733762578
20	8.222441124	1.994306495	1.709854358
20	8.182734047	1.935578419	8.605319945
20	17.599224232	1.972908709	1.701696449
20	17.603587025	1.969186298	8.584268320
20	3.511620251	10.120966507	1.700871240
20	3.507541904	10.104939874	8.582015432
20	12.895781687	10.078816032	1.707031633
20	12.915528214	10.129204287	8.589157044
20	3.609018414	6.128666644	1.707031633
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20	13.017792876	6.118666465	8.605319945
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20	-1.112195556	14.253801439	8.582015432
20	8.277218804	14.254921035	1.701696449
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20	10.514901521	2.011485213	5.096284453
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20	2.301686764	0.040821234	1.701696449
20	2.296281665	0.038904151	8.584268320
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20	6.977602877	8.214406094	8.605319945
20	2.397109069	8.096621674	5.145426921
20	2.392338199	8.096817423	12.031569390
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20	-3.493619021	6.164122728	5.136950696
20	-3.506714076	6.159688336	12.037500494
20	5.866746606	6.163168048	12.036604549
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20	-8.207942778	14.297361805	12.029491596
20	1.182275371	14.296297373	5.145426921
20	1.184491282	14.292067804	12.031569390
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20	4.696870915	13.553487742	10.304540600
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20	4.675865847	2.712256563	10.308786748
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15	1.588607550	3.237954186	8.581877887
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15	6.288611118	11.370975756	8.587745128
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15	3.098123193	4.891763411	12.021727636
15	12.525314488	4.901528689	5.129507908
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15	-1.590695510	13.026796189	12.022509424
15	7.800403102	13.033752591	5.145957554
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15	1.104050185	7.889404863	1.703909799
15	1.087113198	7.893916147	8.584439133
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8	5.442453779	4.188485519	1.676557339
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8	14.894931247	4.184244076	1.710733781
8	14.870012787	4.192237521	8.596601768
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### Cartesian coordinates – Ce2-Na1-close model

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8	-1.685116164	10.785427257	8.575983213
8	7.717842680	10.782618534	1.718711241
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8	1.674042855	5.487442242	5.134489260
8	1.667909166	5.475692532	12.031859289
8	11.121834339	5.532534872	5.112409200
8	11.077799329	5.458068557	12.013174908
8	-3.033362383	13.621894734	5.139742599
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8	6.369845750	13.618066896	12.023417091
8	5.573189803	1.274399114	1.684846324
8	5.576225187	1.272325018	8.589908314
8	14.988035186	1.287878058	1.718171567
8	15.012227158	1.290513912	8.575653261
8	0.876412044	9.431119039	1.703390184
8	0.875761738	9.425929529	8.585513686
8	10.256335340	9.412789628	1.702389046

8	10.287561857	9.447625221	8.617042286
8	-0.858286083	6.856129838	5.138019990
8	-0.879216938	6.842935462	12.019852711
8	8.500533935	6.828975981	5.091977795
8	8.534470394	6.846025037	12.013663321
8	-5.603230559	14.971875119	5.137646605
8	-5.594831950	14.968172725	12.024728034
8	3.833438575	14.976079443	5.128694039
8	3.812893171	14.971506544	12.026286420
8	5.494585557	4.167768647	1.701855593
8	5.453253780	4.174255810	8.623430123
8	14.892308775	4.181554442	1.691846675
8	14.894940806	4.182763904	8.578392652
8	0.781592016	12.329812435	1.697425357
8	0.786595703	12.330644057	8.585153625
8	10.179526174	12.321690823	1.696595319
8	10.162855456	12.347524283	8.597908343
8	-0.773595122	3.945650678	5.140559422
8	-0.783418276	3.936785862	12.026233924
8	8.603229635	3.921703483	5.113348751
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8	11.050416701	4.753955862	8.546068827
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8	3.094254468	3.358704065	5.144332655
8	3.073995546	3.336540218	12.021631831
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8	12.490599494	3.326091796	12.036856922
8	-1.640697689	11.472599505	5.151234332
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8	-0.268742161	7.141442084	1.681241485
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8	9.136513649	7.167732048	8.528778809
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8	4.422316630	15.284984903	8.582650284
8	4.963024698	0.984386775	5.166055694
8	4.965776283	0.982183044	12.038837722
8	14.371451191	1.003567942	5.143508272
8	14.364761596	0.988819971	12.017879750
8	0.220847112	9.168974709	5.147679165
8	0.270764975	9.130104875	12.029030410
8	9.591144738	9.142620419	5.196215767
8	9.671214136	9.141046080	12.030899730
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8	8.010306127	4.300311481	8.583122956
8	-6.058768602	12.463076005	1.697577734
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8	3.339754720	12.460142460	1.687352542
8	3.343644630	12.472126160	8.574848512
8	6.043557334	3.817244589	5.174465869
8	6.039491521	3.784430206	12.034803366
8	15.466526781	3.790703815	5.162811288
8	15.454278766	3.792535193	12.035373383
8	1.336315732	11.964421523	5.147846471



8	1.358063773	11.932734871	12.029277427
8	10.746769711	11.959074207	5.139004869
8	10.750325104	11.926217761	12.017438648
8	0.808520789	2.766496851	0.460311017
8	0.800910389	2.768898127	7.336839649
8	10.198433584	2.749339454	0.461805412
8	10.222793668	2.631896213	7.383227123
8	-3.904532145	10.895637884	0.456747784
8	-3.918837611	10.889496924	7.336851799
8	5.496199098	10.861023039	0.470299228
8	5.474462121	10.926551005	7.325781720
8	3.902292522	5.402725225	6.379136911
8	3.893411668	5.348394828	13.280979367
8	13.335222301	5.371992801	6.373887112
8	13.301281364	5.364305626	13.265602449
8	-0.806226100	13.499585457	6.383021148
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8	11.281698440	7.432629749	0.453332718
8	11.398256320	7.536969693	7.389403573
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8	6.581595967	15.550283303	7.327943033
8	2.786199292	0.695351482	6.381776956
8	2.796993250	0.691668140	13.270816581
8	12.210490717	0.680280920	6.383291730
8	12.213762515	0.693720313	13.280081029
8	-1.937888575	8.806937771	6.383319533
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8	-2.659134186	6.046483887	7.332856361
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8	-7.393506357	14.191544497	7.339052006
8	2.003590564	14.194347620	0.454249621
8	1.998977999	14.198294387	7.335950321
8	7.375456137	2.060157242	6.372616497
8	7.398833133	2.068675052	13.267748902
8	16.826418532	2.057204326	6.381151270
8	16.795774235	2.062625525	13.270551167
8	2.654076653	10.226966383	6.387416339
8	2.699559492	10.205616112	13.268890474
8	12.051598599	10.224615159	6.385457412
8	12.080341709	10.210086556	13.280009397
8	3.915646092	5.399326682	3.906829464
8	3.892532306	5.374099907	10.791760541
8	13.343907881	5.301910209	3.876781815
8	13.308801403	5.339032138	10.772456916
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8	-0.800017640	13.472559679	10.774709024
8	8.592044343	13.453950971	3.885717098
8	8.594233202	13.481873350	10.775779620
8	0.804217961	2.778086093	2.953562892
8	0.800729947	2.786076892	9.832398446
8	10.195989611	2.788566686	2.955173064
8	10.201571698	2.794094813	9.855094198
8	-3.911998118	10.911303024	2.949058049
8	-3.907426370	10.907246499	9.830866892
8	5.485980608	10.948784327	2.959168409
8	5.502164232	10.891502107	9.818321782
8	2.823882395	0.711152394	3.888261412
8	2.812973294	0.697354806	10.778346783
8	12.211552540	0.696607513	3.892735612

8	12.189605493	0.682718478	10.788213913
8	-1.925956143	8.814678045	3.892082830
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8	7.449965523	8.799163957	3.877635997
8	7.518406989	8.841033659	10.776465588
8	1.875029751	7.439768821	2.952534405
8	1.899872515	7.442104508	9.829533930
8	11.290883584	7.439531312	2.945626264
8	11.243674399	7.412248271	9.858144961
8	-2.817792393	15.565109645	2.945466854
8	-2.832871039	15.542389073	9.837313005
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8	7.308624824	2.092226404	3.874742231
8	7.377715512	2.070482814	10.775874245
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8	16.787598371	2.071093148	10.778336468
8	2.645992129	10.234980016	3.890232196
8	2.692535205	10.210913773	10.775290899
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8	12.104168391	10.196241018	10.786356527
8	-2.662076321	6.038316254	2.950337091
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8	6.722787514	6.025219735	9.876936711
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8	2.014760664	14.189517669	9.833042198
9	0.003997232	-0.003455785	1.720068551
9	0.004637052	-0.003827621	8.598089094
9	9.397444017	-0.008035196	1.725667339
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9	4.711456022	8.119953060	1.765715316
9	4.702729781	8.139484855	8.583836582
9	0.003771870	0.003547548	5.121795076
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9	9.397337978	-0.005943723	12.037734117
9	-4.737456042	8.132905115	5.124029893
9	-4.698895424	8.130266690	12.036505995
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### Cartesian coordinates – Ce1-Si-close model

58	9.427270083	5.364236686	6.929563874
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20	5.864955817	6.153124530	5.121935832
20	4.699450849	2.701037191	13.760096296
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20	14.095556873	2.702745577	13.753497404
20	14.090474238	2.677767682	6.901948511
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20	9.391555874	10.861092997	6.868036529
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20	4.706155152	13.535072432	13.743608183
20	8.213590191	1.967989915	1.701372484
20	8.267200255	1.969864156	8.623468398
20	17.600087742	1.966797205	1.704671153
20	17.609876943	1.962749201	8.589844535

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20	3.506950509	10.105333315	8.582174658
20	12.913690076	10.100784466	1.700133872
20	12.923708012	10.102346852	8.599041675
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20	3.563976983	6.126998370	8.599550477
20	12.982665465	6.112664776	1.699479395
20	12.999043685	6.100194126	8.610767922
20	-1.121525300	14.256186151	1.700256473
20	-1.114771261	14.256697895	8.582436958
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20	8.284286460	14.233034461	8.586584149
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20	1.103608234	2.007557404	12.029370161
20	10.521464576	2.008100433	5.162153671
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20	-3.575170947	10.148892833	12.031471835
20	5.801896051	10.137844458	5.142466699
20	5.815799534	10.142630179	12.031546763
20	2.297287790	0.038163842	1.703166673
20	2.303106167	0.036536437	8.588380758
20	11.704506691	0.031136211	1.704026305
20	11.724797993	0.069241470	8.599435199
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20	-2.395545671	8.174341601	8.582779247
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20	6.985874671	8.205917320	8.606503738
20	2.389189876	8.091642194	5.149049381
20	2.384920866	8.091484205	12.032491560
20	11.795716886	8.101035112	5.128435565
20	11.815862032	8.098176806	12.024842722
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20	-2.321872062	16.222862961	12.032787845
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20	7.096462790	16.215551029	12.030343092
20	-3.504895927	6.167645011	5.145774346
20	-3.496214254	6.150780203	12.033084921
20	5.856162669	6.164475333	12.029915433
20	-8.214097846	14.287278068	5.142893160
20	-8.212535027	14.292117032	12.029754477
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20	4.699631264	13.560993312	10.311163073
20	4.661878850	2.695166633	3.416860967
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20	9.397723254	10.865967778	3.414697210
20	9.397787570	10.851101555	10.307898368
15	1.582006502	3.234063766	1.699948250
15	1.586827262	3.237969050	8.586842781
15	10.989445136	3.236686338	1.700367943
15	-3.107605056	11.371118285	1.701022558
15	-3.102752475	11.378500261	8.586957872
15	6.290449647	11.362482454	1.703122593
15	6.286288739	11.357995814	8.582714786
15	3.099955821	4.905237232	5.151491236
15	3.090209132	4.883338829	12.034174884
15	12.526504020	4.878967093	5.122139670
15	12.526212422	4.895861873	12.049792657
15	-1.590210384	13.028581746	5.144478458

15	-1.589390802	13.028340696	12.025584404
15	7.800362458	13.009086561	5.143179970
15	7.808580186	13.010652781	12.030227132
15	1.093024517	7.894596392	1.703578822
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15	10.499323334	7.896057897	1.697163354
15	10.481862592	7.907667650	8.613748166
15	-3.620341584	16.017221518	1.706138553
15	-3.604359350	16.030569893	8.582260816
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15	5.825524866	16.009062210	8.586938823
15	3.600278943	0.236262061	5.145928624
15	3.590763167	0.237225531	12.031019159
15	12.997368016	0.226279689	5.135334536
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15	-1.099134679	8.376505216	12.027184996
15	8.310383204	8.385670046	5.125105216
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15	2.027010016	13.269047364	8.586178780
15	7.342311991	2.994423579	5.119609697
15	7.361654819	2.986157971	12.050258059
15	16.779301370	2.995465227	5.147427485
15	16.783755713	2.984200787	12.031212286
15	2.687879460	11.128338147	5.145676228
15	2.689100470	11.123497822	12.026658789
15	12.094568629	11.117677830	5.152427973
15	12.083277666	11.134294435	12.032207197
8	3.012629053	2.646170298	1.696963377
8	3.016665768	2.648652408	8.591638185
8	12.424100730	2.664766235	1.712558287
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8	-1.677688123	10.783565907	1.704242021
8	-1.670934515	10.795780665	8.588795002
8	7.719568317	10.771412316	1.705452274
8	7.720939314	10.776062798	8.586662768
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8	1.659559088	5.469638947	12.032304292
8	11.076805259	5.439826357	5.080701210
8	11.101355012	5.507486208	12.069013150
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8	-3.020371046	13.616734202	12.021495386
8	6.375705281	13.606513946	5.141394127
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8	5.562599931	1.280429364	1.714761697
8	5.577004433	1.294044973	8.590261462
8	14.978124402	1.273567166	1.700142542
8	15.073534780	1.279512232	8.595532457
8	0.887674670	9.427132346	1.705806161
8	0.877420662	9.423419184	8.589987213
8	10.291143990	9.427668823	1.711120764
8	10.253656526	9.430545948	8.561253803
8	-0.888316876	6.846130207	5.143717572
8	-0.891912298	6.843760866	12.026216912
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8	8.480937978	6.815481765	12.009497545
8	-5.599522834	14.969530582	5.138791953
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8	3.807240285	14.960349119	5.123622867
8	3.830114415	14.956209145	12.055157580
8	5.469950227	4.189816295	1.709031836
8	5.449165260	4.200102630	8.549484913
8	14.898595255	4.169958659	1.713055944

8	14.869701381	4.202810387	8.590403743
8	0.781043945	12.320443067	1.703900000
8	0.800738279	12.325200000	8.586432275
8	10.179735539	12.317434088	1.694037708
8	10.187069473	12.316641573	8.589360287
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8	1.615463360	4.790728372	1.691708543
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8	11.011307411	4.796532175	1.665963949
8	11.055052324	4.871534678	8.698451281
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8	3.065161941	3.348635555	5.164893347
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8	12.506037584	3.327297219	5.147785140
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8	-4.980013818	15.258567739	1.694691853
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8	4.444592983	15.290573723	8.575823721
8	4.964278411	0.985521107	5.157678942
8	4.950496661	0.994147275	12.035302405
8	14.361904753	0.975161315	5.180063520
8	14.377342745	0.980401815	12.022793730
8	0.264861107	9.132050948	5.151878791
8	0.263824117	9.129119350	12.031048526
8	9.661340788	9.153867384	5.124826264
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8	6.005426368	3.788635838	5.118685189
8	6.041202824	3.812759472	12.022542332
8	15.449035567	3.801454603	5.177943806
8	15.448210337	3.781700851	12.010387695
8	1.357495325	11.936457735	5.146463373
8	1.358845857	11.932760692	12.035399273
8	10.763156748	11.924659140	5.157642719
8	10.745393511	11.929601946	12.048540158
8	0.784563490	2.760169707	0.459870640
8	0.793132865	2.759403516	7.345850715
8	10.194403819	2.740221908	0.467946346
8	10.119515580	2.937600827	7.257793306
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8	-3.897070549	10.909284904	7.342988793
8	5.494512693	10.887908890	0.461477177
8	5.499304001	10.878572498	7.338168388
8	3.903424665	5.387407132	6.386780946

8	3.887467158	5.364012683	13.273072585
8	13.267176643	5.399574412	6.375326141
8	13.327330443	5.321338005	13.305072024
8	-0.793911170	13.498761303	6.388176064
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8	11.119397565	7.379771523	7.291034846
8	-2.811338977	15.569332529	0.462403638
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8	2.784891665	0.684354000	13.275165579
8	12.156687602	0.659200945	6.360518711
8	12.239188131	0.693915170	13.304041363
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8	-1.898270314	8.829838881	13.275029774
8	7.516762707	8.786834139	6.391612983
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8	2.027662530	14.185186531	7.338694163
8	7.381367505	2.116764059	6.391524993
8	7.329824483	2.076304237	13.304497058
8	16.814859953	2.057296354	6.379175092
8	16.780350370	2.073707011	13.285691590
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8	12.098754397	10.201700453	13.269759867
8	3.888075150	5.357934266	3.895666420
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8	13.319920399	5.344118042	3.880846340
8	13.317725291	5.393594270	10.817179089
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8	8.604518710	13.455204914	10.776665648
8	0.794266300	2.779008729	2.954402342
8	0.793528046	2.791294912	9.840192370
8	10.193940614	2.808713221	2.961729834
8	10.131273051	2.707378773	9.887234505
8	-3.903567014	10.922307182	2.952927128
8	-3.898754901	10.916843223	9.835474587
8	5.492347595	10.914727949	2.953699105
8	5.496320624	10.886622459	9.829442089
8	2.814710623	0.706007697	3.894890552
8	2.794416043	0.692780424	10.782253909
8	12.236342508	0.703026580	3.873540071
8	12.197344457	0.654087270	10.815436270
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8	-1.896231066	8.835062746	10.780357613
8	7.488673374	8.829660735	3.894094331
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8	1.880982312	7.430986644	2.954695310
8	1.885149403	7.433905601	9.836745832
8	11.272652809	7.421272188	2.954083644
8	11.423442896	7.542477086	9.776501820
8	-2.828837782	15.554243991	2.955425300
8	-2.803963717	15.573237186	9.828045047
8	6.585405823	15.555999840	2.950507428

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8	7.361794913	2.060327829	3.888446570
8	7.362515083	2.061288526	10.810066805
8	16.752691229	2.088668130	3.888895333
8	16.821930786	2.052126430	10.795897129
8	2.690771287	10.208882530	3.898309905
8	2.681444479	10.206922039	10.777116736
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8	12.078829192	10.228338864	10.776303639
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8	2.031971410	14.181175542	9.838408546
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9	0.002952414	-0.009617332	8.606672056
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9	4.691577256	8.135378749	8.519611992
9	0.000120560	-0.009362929	5.123061362
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9	-4.684035485	8.135641273	12.036075597
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### Cartesian coordinates – Ce2-Si-close model

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20	5.958904535	6.095079503	5.145518040
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20	4.693408561	2.713379275	6.876911998
20	14.086877586	2.719733768	13.755360374
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20	8.208381147	1.969455574	1.698607485
20	8.207820740	1.970717912	8.587511429
20	17.598283938	1.985559755	1.700613090
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20	12.895954836	10.115763145	1.705875617
20	12.895404060	10.115265608	8.579450927
20	3.592378081	6.131338424	1.715461938
20	3.593729285	6.132664897	8.571564789
20	12.964795464	6.122156302	1.705056385
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20	1.111147533	2.012214701	5.142229079
20	1.117317294	2.019883362	12.023269433
20	10.498734633	1.998958040	5.143006728
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20	-3.584417612	10.155539142	12.023784740
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20	2.333618655	8.058570229	5.143412125
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20	7.092663027	16.260054547	12.023778017
20	-3.522024580	6.161214078	5.142379071
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20	5.877344733	6.160489197	12.024338781
20	-8.206858611	14.312818189	5.142718458
20	-8.207718599	14.312494268	12.023460623
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20	4.693874425	2.709986729	3.417361005
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15	-3.113639775	11.385621730	8.587491742
15	6.298935807	11.432346872	1.657619637
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15	7.799087546	13.052104237	12.023549962
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15	5.785554161	16.061709157	8.585577848
15	3.596372196	0.243111739	5.143507247



15	3.593371941	0.237504864	12.024227243
15	12.993994593	0.252159736	5.142777067
15	12.982761335	0.249139775	12.023813036
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8	12.415632152	2.653081819	8.592780681
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8	11.072204720	5.486681826	5.139117718
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8	6.381798712	13.646276180	12.021739097
8	5.594056486	1.277516824	1.731512514
8	5.598164198	1.277043827	8.558763066
8	14.978096657	1.292843533	1.723081538
8	14.980713030	1.293786556	8.567455922
8	0.897951112	9.453034278	1.740677699
8	0.897922040	9.450729627	8.556939181
8	10.260380970	9.426810618	1.709132451
8	10.261120556	9.426491176	8.585168053
8	-0.859758321	6.871010241	5.142374142
8	-0.885833432	6.861179224	12.024227698
8	8.484603107	6.838798450	5.141018139
8	8.517578595	6.857653291	12.024863024
8	-5.595360923	15.001989364	5.142024514
8	-5.642725633	14.890632191	12.025833685
8	3.831240687	15.019100787	5.139078789
8	3.793004109	15.002530185	12.020366451
8	5.480698477	4.175408901	1.692665652
8	5.480941566	4.174688163	8.602332102
8	14.865932212	4.207190291	1.707228687
8	14.865700668	4.206972572	8.585673310
8	0.799128344	12.325279827	1.718205340
8	0.801230210	12.324265381	8.568903265
8	10.177329707	12.326197463	1.719020544
8	10.175871074	12.327472702	8.569048603
8	-0.798702783	3.968251141	5.140797237
8	-0.818665637	3.982003860	12.021509248
8	8.576727187	3.961728261	5.143920897
8	8.603030668	3.978303344	12.024365948
8	-5.485867796	12.091898110	5.143136638
8	-5.437563328	12.019841470	12.023243828
8	3.893853802	12.143481269	5.146240897
8	3.911842375	12.092547453	12.025362867
8	1.614485528	4.840053501	1.719273809
8	1.612303243	4.838577104	8.561871991
8	11.020040826	4.801013815	1.691428548
8	11.020426358	4.800204818	8.579553092
8	-3.086669699	12.939508953	1.682471647
8	-3.087541553	12.939196767	8.589185813
8	6.323853937	12.959829392	1.646316106
8	6.322940446	12.957743208	8.614978539
8	3.053422598	3.327471255	5.144233674
8	3.076177864	3.401850083	12.027200973
8	12.448213729	3.339197066	5.147921773

8	12.458213931	3.343023964	12.031782985
8	-1.640827565	11.475600228	5.148892451
8	-1.623831240	11.483271975	12.029040309
8	7.772197424	11.447374525	5.152502520
8	7.777175129	11.493620610	12.034301927
8	-0.263123974	7.167468039	1.683611776
8	-0.264374100	7.165783759	8.576435509
8	9.128985418	7.132388680	1.713756972
8	9.129004377	7.132566422	8.558141403
8	-4.965364650	15.289605143	1.702801312
8	-4.965141607	15.291204217	8.574489387
8	4.417654307	15.307878134	1.660450912
8	4.417519818	15.309908235	8.622148217
8	4.963689996	0.987942962	5.148587493
8	5.042394928	1.016003704	12.029726087
8	14.339789145	1.023742532	5.146460061
8	14.341586458	0.991585187	12.028332404
8	0.283566517	9.161679464	5.149149844
8	0.279161788	9.143414116	12.032660104
8	9.621229734	9.129362295	5.146523406
8	9.658997710	9.148270375	12.027067379
8	-1.362926815	4.361703997	1.712057534
8	-1.363076988	4.359985570	8.569346260
8	8.041808234	4.322341352	1.688074589
8	8.042333552	4.320775223	8.594892285
8	-6.053538641	12.490290115	1.695287696
8	-6.054843922	12.490489865	8.573815263
8	3.355903246	12.476218188	1.682159340
8	3.358164706	12.475862244	8.596937421
8	6.036971105	3.733555503	5.146640787
8	6.050190446	3.794606435	12.026946965
8	15.432684341	3.811346976	5.145650208
8	15.415154449	3.812628133	12.029423618
8	1.357552767	11.862619160	5.140890574
8	1.354600910	11.956976628	12.025492324
8	10.733651576	11.959781590	5.148656093
8	10.774541682	11.974696945	12.029560347
8	0.781035086	2.842423248	0.459868094
8	0.774922749	2.814405808	7.328132482
8	10.197143779	2.764597368	0.466459429
8	10.192842586	2.786653971	7.330739819
8	-3.913105672	10.888413277	0.467209313
8	-3.901998487	10.919924239	7.330484270
8	5.498924924	10.898716477	0.451906467
8	5.499976216	10.953306737	7.366213998
8	3.910769532	5.327014557	6.393395095
8	3.888672823	5.439696079	13.269662683
8	13.281722846	5.357931372	6.389606762
8	13.290596769	5.368853212	13.271117534
8	-0.785405201	13.478166670	6.388996793
8	-0.787917493	13.495135122	13.271746351
8	8.567794027	13.487182495	6.388785436
8	8.606639680	13.516657120	13.269840517
8	1.934866563	7.493188774	0.486266065
8	1.899784947	7.449505329	7.330044575
8	11.286915146	7.448308379	0.462510130
8	11.301444293	7.459062879	7.337147874
8	-2.783710298	15.595173562	0.479172066
8	-2.810509562	15.560749545	7.319983732
8	6.612907635	15.615844448	0.472778175
8	6.535065010	15.563776489	7.313413473
8	2.795433255	0.700642607	6.390167459
8	2.774687630	0.764067426	13.336820182
8	12.179034899	0.687500061	6.389476079
8	12.174974112	0.691287566	13.270052565
8	-1.880861854	8.854478580	6.388145727
8	-1.888690117	8.850903450	13.268559663
8	7.440599884	8.817843279	6.378801492
8	7.498952275	8.838629595	13.270774276
8	-2.703139147	6.081833308	0.461171845

8	-2.711120422	6.085798135	7.335642224
8	6.697648946	6.057837945	0.462987948
8	6.704435595	6.032763643	7.340613435
8	-7.431048376	14.223331895	0.491265226
8	-7.401148621	14.200600073	7.312286470
8	1.990976960	14.205974154	0.468924737
8	2.041760355	14.181258943	7.319867094
8	7.434555407	2.047940304	6.387239412
8	7.441402782	2.089922901	13.271370322
8	16.775361722	2.085980150	6.387897339
8	16.758181482	2.096845054	13.272420198
8	2.846626791	10.191731602	6.373355052
8	2.685169784	10.230172508	13.269635244
8	12.055431863	10.229234333	6.389833825
8	12.048440175	10.208325549	13.273897825
8	3.909376000	5.330839088	3.899548381
8	3.886423593	5.440147318	10.784768767
8	13.279398964	5.353392963	3.897546259
8	13.289145302	5.359279974	10.777441428
8	-0.787270326	13.468245861	3.892808441
8	-0.787546843	13.486829016	10.773481530
8	8.565862982	13.469880328	3.889761410
8	8.604562791	13.497082750	10.769353722
8	0.776537860	2.818220428	2.957494703
8	0.781281327	2.842047407	9.826132121
8	10.194370752	2.801100901	2.961127540
8	10.198771909	2.778290521	9.826039723
8	-3.901734067	10.932304605	2.957231349
8	-3.911358054	10.897709521	9.819978504
8	5.502194937	10.982011593	2.929788253
8	5.501025833	10.919818288	9.844643470
8	2.802441351	0.704147497	3.895411905
8	2.780840764	0.765972256	10.714166614
8	12.184300354	0.693448919	3.894184095
8	12.179974651	0.697845134	10.777413524
8	-1.869712884	8.860050959	3.892389309
8	-1.874558213	8.856261602	10.773257130
8	7.444702551	8.823067719	3.905707901
8	7.501649945	8.840950240	10.778022528
8	1.873783768	7.438497269	2.972283785
8	1.907317255	7.476410038	9.817288440
8	11.288377289	7.450395899	2.956208306
8	11.273555323	7.437888622	9.830607019
8	-2.818055394	15.559104742	2.968125047
8	-2.789193090	15.589609794	9.809303584
8	6.534464615	15.560627704	2.970850625
8	6.611613664	15.611494964	9.811767356
8	7.431161425	2.050406536	3.898822183
8	7.438738597	2.089905692	10.779664814
8	16.770922704	2.087280414	3.897627523
8	16.749813345	2.099053323	10.776270391
8	2.855704705	10.187948873	3.919864513
8	2.685802528	10.228334087	10.782863152
8	12.050831823	10.233522348	3.896070852
8	12.043432285	10.212250759	10.773574416
8	-2.705028503	6.085706068	2.953072822
8	-2.694929141	6.079353550	9.827831575
8	6.709469718	6.032136254	2.951848556
8	6.705928199	6.054534655	9.829557623
8	-7.387146848	14.190148600	2.979551584
8	-7.419108390	14.215599454	9.800699231
8	2.042670799	14.180532979	2.965750710
8	1.996455031	14.203241905	9.817347016
9	18.761402812	0.017772909	1.742836816
9	18.759239437	0.017982470	8.552774037
9	0.000016929	16.280299798	1.730189332
9	0.000517931	16.278129318	8.569025847
9	-4.699737055	8.152092037	1.711997749
9	-4.699643320	8.151087790	8.590815058
9	4.714224225	8.166403057	1.784552077



9	4.713037272	8.162593140	8.538394290
9	9.362171451	16.281789382	5.134951596
9	-0.021644768	-0.001938490	12.015791884
9	0.009403638	16.246292022	5.137233836
9	0.002548070	16.278929019	12.022559957
9	14.076677297	8.144663117	5.143429081
9	-4.699635268	8.151922356	12.020778806
9	4.673684338	8.002370081	5.149652702
9	4.707692287	8.163216738	12.027868944

### EXAFS spectrum of Durango apatite measured on FAME-UHD

2.5000	-0.096252
2.5500	-0.045720
2.6000	-0.0034365
2.6500	0.036065
2.7000	0.071844
2.7500	0.098624
2.8000	0.11226
2.8500	0.11935
2.9000	0.11174
2.9500	0.095042
3.0000	0.083590
3.0500	0.065356
3.1000	0.045377
3.1500	0.039627
3.2000	0.035188
3.2500	0.035926
3.3000	0.034318
3.3500	0.037123
3.4000	0.035754
3.4500	0.030352
3.5000	0.020697
3.5500	0.0049137
3.6000	-0.0078926
3.6500	-0.023631
3.7000	-0.039874
3.7500	-0.045355
3.8000	-0.045444
3.8500	-0.046777
3.9000	-0.047993
3.9500	-0.048546
4.0000	-0.046454
4.0500	-0.044406
4.1000	-0.043535
4.1500	-0.034342
4.2000	-0.018100
4.2500	-0.0012503
4.3000	0.010947
4.3500	0.020293
4.4000	0.025090
4.4500	0.028016
4.5000	0.027668
4.5500	0.024052
4.6000	0.021243
4.6500	0.013914
4.7000	0.0072647
4.7500	0.0038517
4.8000	-8.9974e-05
4.8500	-0.0036865
4.9000	-0.0050456
4.9500	-0.0045652
5.0000	-0.0058546
5.0500	-0.0023051
5.1000	0.0088106
5.1500	0.016375
5.2000	0.022008
5.2500	0.023962
5.3000	0.017427
5.3500	0.015011
5.4000	0.011797

5.4500 0.0018258  
5.5000 5.1639e-05  
5.5500 -0.0030299  
5.6000 -0.011271  
5.6500 -0.015343  
5.7000 -0.020243  
5.7500 -0.026203  
5.8000 -0.026683  
5.8500 -0.017742  
5.9000 -0.014807  
5.9500 -0.017340  
6.0000 -0.021476  
6.0500 -0.011898  
6.1000 -0.0012348  
6.1500 0.0075337  
6.2000 0.0096617  
6.2500 0.019258  
6.3000 0.023097  
6.3500 0.015667  
6.4000 0.018029  
6.4500 0.016242  
6.5000 0.0086194  
6.5500 0.0081524  
6.6000 0.0011532  
6.6500 -0.0093705  
6.7000 -0.0061468  
6.7500 -0.0050710  
6.8000 -0.0030558  
6.8500 -0.0053557  
6.9000 0.0021153  
6.9500 0.0026021  
7.0000 -0.00028372  
7.0500 0.0029845  
7.1000 0.0020452  
7.1500 0.0035960  
7.2000 -0.0010280  
7.2500 -0.0035058  
7.3000 -0.0051100  
7.3500 -0.0060543  
7.4000 -0.0057650  
7.4500 -0.0072361  
7.5000 -0.0072989  
7.5500 -0.0054153  
7.6000 -0.0040626  
7.6500 -0.0026145  
7.7000 0.0024358  
7.7500 0.0060414  
7.8000 0.0067560  
7.8500 0.0078724  
7.9000 0.0044022  
7.9500 0.0012902  
8.0000 0.0038620  
8.0500 0.0040236  
8.1000 0.0032208  
8.1500 0.0029432  
8.2000 -0.0010131  
8.2500 -0.0032790  
8.3000 -0.0051100  
8.3500 -0.0040637  
8.4000 -0.0012968  
8.4500 -0.00019127  
8.5000 -0.00017960  
8.5500 0.0011338  
8.6000 -0.00042640  
8.6500 -0.0012367  
8.7000 0.00095895  
8.7500 0.0028951  
8.8000 0.00047861  
8.8500 -0.00094246  
8.9000 0.0017463

8.9500	0.0019876
9.0000	-0.0026151
9.0500	-0.0035535
9.1000	-0.0022670
9.1500	-0.0026673
9.2000	-0.0052786
9.2500	-0.0021929
9.3000	0.0017483
9.3500	0.0022302
9.4000	0.0022291
9.4500	0.0017168
9.5000	0.0015086
9.5500	0.0024608
9.6000	0.0011194
9.6500	0.0022258
9.7000	0.0050673
9.7500	0.0036302
9.8000	-0.0010479
9.8500	-0.0029347
9.9000	-0.00011018
9.9500	0.00049115
10.000	-0.0014126
10.050	-0.0014208
10.100	-0.0023414
10.150	-0.0028011
10.200	-0.0018214
10.250	-0.0019143
10.300	-0.0014082
10.350	-0.0014382
10.400	-5.2058e-05

#### **EXAFS spectrum of Durango apatite measured on ID24**

2.5000	-0.086069
2.5500	-0.051740
2.6000	-0.012006
2.6500	0.024093
2.7000	0.056118
2.7500	0.078940
2.8000	0.096976
2.8500	0.10708
2.9000	0.10992
2.9500	0.10263
3.0000	0.089178
3.0500	0.075551
3.1000	0.055826
3.1500	0.040439
3.2000	0.026840
3.2500	0.020297
3.3000	0.016386
3.3500	0.013639
3.4000	0.012592
3.4500	0.0088105
3.5000	0.0028457
3.5500	-0.0044123
3.6000	-0.014477
3.6500	-0.025397
3.7000	-0.034593
3.7500	-0.041820
3.8000	-0.046904
3.8500	-0.048100
3.9000	-0.047524
3.9500	-0.046103
4.0000	-0.045066
4.0500	-0.041537
4.1000	-0.036602
4.1500	-0.028246
4.2000	-0.016233
4.2500	-0.0036438
4.3000	0.0067434
4.3500	0.016268

4.4000 0.023809  
4.4500 0.026827  
4.5000 0.027379  
4.5500 0.026702  
4.6000 0.023845  
4.6500 0.019496  
4.7000 0.015085  
4.7500 0.011527  
4.8000 0.0080304  
4.8500 0.0043769  
4.9000 0.0016301  
4.9500 0.00064078  
5.0000 0.0012807  
5.0500 0.0015357  
5.1000 0.0033889  
5.1500 0.0071302  
5.2000 0.0097869  
5.2500 0.011063  
5.3000 0.010419  
5.3500 0.0079620  
5.4000 0.0030466  
5.4500 -0.0018494  
5.5000 -0.0044753  
5.5500 -0.0054697  
5.6000 -0.0069915  
5.6500 -0.010588  
5.7000 -0.014915  
5.7500 -0.017689  
5.8000 -0.016737  
5.8500 -0.014128  
5.9000 -0.013137  
5.9500 -0.014478  
6.0000 -0.012896  
6.0500 -0.0097491  
6.1000 -0.0046882  
6.1500 0.0020940  
6.2000 0.0076636  
6.2500 0.010890  
6.3000 0.013035  
6.3500 0.014282  
6.4000 0.014088  
6.4500 0.011869  
6.5000 0.0094003  
6.5500 0.0073621  
6.6000 0.0035624  
6.6500 -0.0018255  
6.7000 -0.0050688  
6.7500 -0.0042296  
6.8000 -0.0026977  
6.8500 -0.0021917  
6.9000 -0.0021680  
6.9500 -0.0016233  
7.0000 -0.00099464  
7.0500 -0.00016423  
7.1000 0.0011613  
7.1500 0.0010782  
7.2000 0.00058766  
7.2500 -0.00070671  
7.3000 -0.0031976  
7.3500 -0.0034309  
7.4000 -0.0036845  
7.4500 -0.0046083  
7.5000 -0.0031918  
7.5500 -0.0016818  
7.6000 -0.00030985  
7.6500 -0.00055444  
7.7000 0.00017722  
7.7500 0.0022999  
7.8000 0.0033400  
7.8500 0.0042637

7.9000	0.0034286
7.9500	0.0034087
8.0000	0.0028021
8.0500	0.0020395
8.1000	0.0013490
8.1500	0.0011640
8.2000	-0.0011175
8.2500	-0.0020062
8.3000	-0.0020420
8.3500	-0.0013772
8.4000	-0.0013329
8.4500	-0.00081975
8.5000	-0.00068989
8.5500	-0.00039273
8.6000	-0.00024817
8.6500	-0.00027684
8.7000	-0.00046088
8.7500	-0.00020332
8.8000	0.00028250
8.8500	-0.00051243
8.9000	-0.00058734
8.9500	-0.0015527
9.0000	-0.0023425
9.0500	-0.0020482
9.1000	-0.0013285
9.1500	-0.0011026
9.2000	-0.0022643
9.2500	-0.0012793
9.3000	-0.00078079
9.3500	-0.00012330
9.4000	0.00090597
9.4500	0.0015535
9.5000	0.0012893
9.5500	0.0018401
9.6000	0.0022821
9.6500	0.0023457
9.7000	0.0028299
9.7500	0.0029452
9.8000	0.0026821
9.8500	0.0019583
9.9000	0.00099696
9.9500	4.2563e-06
10.000	0.00015271
10.050	-0.00017758
10.100	0.00048826
10.150	-0.00036098
10.200	-0.0021954
10.250	-0.0018545
10.300	-0.0021366
10.350	-0.0028273
10.400	-0.0024002

### **EXAFS spectrum of Imilchil apatite measured on BM20**

2.5000	-0.087850
2.5500	-0.052709
2.6000	-0.012356
2.6500	0.030397
2.7000	0.067925
2.7500	0.096936
2.8000	0.11775
2.8500	0.12653
2.9000	0.12397
2.9500	0.11306
3.0000	0.094036
3.0500	0.070214
3.1000	0.048560
3.1500	0.034130
3.2000	0.022611
3.2500	0.013894
3.3000	0.0093002
3.3500	0.0088011

3.4000	0.0083194
3.4500	0.0059332
3.5000	0.0020222
3.5500	-0.0078052
3.6000	-0.019385
3.6500	-0.029284
3.7000	-0.038348
3.7500	-0.043864
3.8000	-0.045512
3.8500	-0.047336
3.9000	-0.049145
3.9500	-0.047538
4.0000	-0.045699
4.0500	-0.044039
4.1000	-0.037423
4.1500	-0.028534
4.2000	-0.017519
4.2500	-0.0029764
4.3000	0.0081335
4.3500	0.015620
4.4000	0.020620
4.4500	0.023665
4.5000	0.024272
4.5500	0.023245
4.6000	0.020623
4.6500	0.015378
4.7000	0.011366
4.7500	0.0082097
4.8000	0.0042483
4.8500	0.0035787
4.9000	0.0048005
4.9500	0.0045643
5.0000	0.0055831
5.0500	0.0078807
5.1000	0.010061
5.1500	0.012135
5.2000	0.014083
5.2500	0.013446
5.3000	0.011191
5.3500	0.0086495
5.4000	0.0033359
5.4500	-0.0024576
5.5000	-0.0061751
5.5500	-0.0094093
5.6000	-0.012237
5.6500	-0.014470
5.7000	-0.016747
5.7500	-0.018297
5.8000	-0.016915
5.8500	-0.013092
5.9000	-0.011263
5.9500	-0.012170
6.0000	-0.011168
6.0500	-0.0069276
6.1000	-0.0015312
6.1500	0.0030855
6.2000	0.0067398
6.2500	0.0096260
6.3000	0.011141
6.3500	0.011768
6.4000	0.011911
6.4500	0.010665
6.5000	0.0085288
6.5500	0.0057420
6.6000	0.0026985
6.6500	-0.0010908
6.7000	-0.0037000
6.7500	-0.0039228
6.8000	-0.0039742
6.8500	-0.0031791
6.9000	-0.0016465
6.9500	-0.0015549
7.0000	-0.0017399
7.0500	-0.0011457
7.1000	-0.00039838

7.1500	-0.00054303
7.2000	-0.00080400
7.2500	-0.00055174
7.3000	-0.0012504
7.3500	-0.0031494
7.4000	-0.0042145
7.4500	-0.0035135
7.5000	-0.0015187
7.5500	-5.6599e-05
7.6000	0.00069428
7.6500	0.0017456
7.7000	0.0030173
7.7500	0.0034498
7.8000	0.0021659
7.8500	0.0018294
7.9000	0.0024114
7.9500	0.0020550
8.0000	0.0015824
8.0500	0.00080422
8.1000	0.00037195
8.1500	0.00084576
8.2000	0.00025243
8.2500	-0.0010086
8.3000	-9.7989e-06
8.3500	0.00072112
8.4000	0.00025463
8.4500	0.00077665
8.5000	0.00049537
8.5500	-0.00094348
8.6000	-0.00087168
8.6500	0.0012149
8.7000	0.0012429
8.7500	-0.00055663
8.8000	-0.0012447
8.8500	-0.0013341
8.9000	-0.0016017
8.9500	-0.0023305
9.0000	-0.0024625
9.0500	-0.0020448
9.1000	-0.0019730
9.1500	-0.0011214
9.2000	-0.00019757
9.2500	-0.00077469
9.3000	-0.0010906
9.3500	-0.00074049
9.4000	-0.00016591
9.4500	2.8602e-06
9.5000	-1.6476e-05
9.5500	0.00043085
9.6000	0.00096817
9.6500	0.0012670
9.7000	0.00094514
9.7500	0.0011273
9.8000	0.0016746
9.8500	0.0013154
9.9000	0.00067551
9.9500	0.0010378
10.000	0.0019301
10.050	0.0019060
10.100	0.00066569
10.150	-8.8946e-05
10.200	-0.00021102
10.250	-0.00013918
10.300	0.00053326
10.350	0.00088069
10.400	0.00074169