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Modeling the attenuated total reflectance infrared (ATR-FTIR) spectrum of apatite

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Abstract

Attenuated Total Reflectance (ATR) infrared spectra were measured on a synthetic and a natural fluorapatite sample. A modeling approach based on the computation of the Fresnel reflection coefficient between the ATR crystal and the powder sample was used to analyze the line shape of the spectra. The dielectric properties of the samples were related to those of pure fluorapatite using an effective medium approach, based on Maxwell-Garnett and Bruggeman models. The Bruggeman effective medium model leads to a very good agreement with the experimental data recorded on the synthetic fluorapatite sample. The poorer agreement observed on the natural sample suggests a more significant heterogeneity of the sample at a characteristic length scale larger than the mid-infrared characteristic wavelength, i.e. about 10 micrometers. The results demonstrate the prominent role of macroscopic electrostatic effects over fine details of the microscopic structure in determining the line shape of strong ATR bands.

1. Introduction

Hard-mode infrared spectroscopy provides a detailed information on the short-range order in minerals (Salje and Bismayer 1997; Salje et al. 2000). It is a particularly relevant technique to investigate a variety of phenomena leading to structural heterogeneity, including the incorporation of molecular impurities (Yi et al. 2014), exsolution processes (Zhang et al. 1996), stacking variations in layered structures (Balan et al. 2010), phase transitions (Salje and Yagil 1996) and cationic ordering (Meyer et al. 2002).

However, the infrared spectra of minerals not only depend on their intrinsic vibrational properties, but are also a function of the sample micro-structure and experimental geometry. In many cases, the spectra are recorded on powder samples and the interplay of the various parameters involved may impede their straightforward interpretation in terms of microscopic structural parameters. This difficulty is particularly apparent when comparing absorption spectra recorded using the traditional pellet method with those obtained using the more recently developed attenuated total reflectance (ATR) infrared spectroscopy. In the first case, the sample consists of particles of the studied mineral diluted in a non-absorbing matrix, usually KBr, and the spectrum is recorded in a transmission geometry. In the second case, the absorption spectrum is recorded in a reflection geometry at the interface between the packed mineral powder and a highly refractive material. ATR thus appears as a very easy experimental procedure which facilitates the acquisition of "absorption-like" infrared spectra by avoiding the multi-step preparation of dry and well-dispersed sample pellets. In many circumstances, it is the only method available to record the infrared spectrum of a given sample. ATR measurements can also be used over a wide frequency range, as attested by measurement of electronic excitations in the near-infrared region (Salje and Güttler 1984).

However, the differences in sample nature and experimental configuration lead to significant differences in the signals recorded in transmission or ATR geometry.

In the present study, we adopt a modeling approach to investigate the line shape variations observed in the ATR infrared spectra of selected samples of fluorapatite. Apatite is a widespread calcium phosphate mineral with nominal composition $\text{Ca}_{10}(\text{PO}_4)_6(\text{OH},\text{F},\text{Cl})_2$. It occurs in many different geological contexts. For example, world phosphorus resources mainly consist of sedimentary deposits of carbonate-bearing fluorapatite (Knudsen and Gunter 2002), while carbonate-bearing hydroxyapatite is the major inorganic component of vertebrate skeletons (e.g. Elliott 2002; Reynard and Balter 2014). The flexibility of the apatite structure enables significant chemical variations, among which carbonate group incorporation is of peculiar importance (e.g. Leventouri et al. 2000; Pan and Fleet 2002; Antonakos et al. 2007). Infrared spectroscopy of apatite usually provides two different types of information. The crystalline quality of the samples is assessed from the width of absorption bands due to phosphate vibrational modes. This information is usually obtained from relative spectral intensities in the range of PO_4 bending vibrational modes ($\nu_4 \text{PO}_4$) and reported as an empirical "crystallinity" index (e.g. Shemesh 1990; Weiner and Bar-Yosef 1990; Surovell and Stiner 2001; Puc  at et al. 2004; Trueman et al. 2008; Roche et al. 2010). The other type of information is related to the concentration and location of molecular species, such as carbonate groups, that are detected by specific vibrational bands (e.g. Michel et al. 1995; Fleet and Liu 2008; Fleet 2009; Yi et al. 2014; Chevrin  ais et al. 2016). Although the majority of infrared spectroscopic studies of apatite were performed using transmission experiments, recent investigations have used ATR spectroscopy to unravel the modification of biologic apatite during fossilization (e.g. Stathopoulou et al. 2008; Lebon et al. 2014, Lang et al. 2016) and burning (Thompson et al. 2009). Systematic comparison between transmission and ATR

spectra have revealed small but significant differences in the corresponding "crystallinity" index and carbonate to phosphate ratio (Thompson et al. 2009; Beasley et al. 2014).

Our modeling strategy is similar to that previously used by Balan et al. (2002) to model the ATR-FTIR spectrum of aggregates of chrysotile nanotubes. It combines the computation of the frequency-dependent reflection coefficient at the interface between the ATR crystal and the sample, with an effective medium modeling of the dielectric properties of the composite powder sample. The obtained results make it possible to discuss the respective role of the various physical parameters determining the shape of ATR spectrum and to identify the most appropriate effective medium model that accounts for the properties of apatite powders.

2. Samples and methods

Two reference apatite samples have been investigated. The first one is a pure fluorapatite sample containing trace amounts of cadmium ($\text{Cd}_{0.005}\text{Ca}_{9.995}(\text{PO}_4)_6(\text{F},\text{OH})_2$) and synthesized at high temperature (Nounah and Lacout 1993). The transmission powder infrared spectrum of this sample has been investigated in details in Balan et al. (2011). The small linewidth ($\sim 4 \text{ cm}^{-1}$) of the weak $\nu_1 \text{ PO}_4$ stretching band measured at low temperature (10K) attests to the very high crystalline quality of this sample. Scanning electron microscope (SEM) observations display a significant porosity and sub-micrometric coalesced particles with a rounded shape (Fig. 1). The second one is a natural sample of carbonate-fluorapatite, originating from the sedimentary phosphate deposit of Taiba (Senegal) (Morin et al. 2002). It contains 90 wt% of highly crystalline fluorapatite with a few associated mineral impurities ($\sim 7 \text{ wt\%}$ quartz, $\sim 2 \text{ wt\%}$ crandallite, $<1 \text{ wt\%}$ goethite, $<1 \text{ wt\%}$ dioctahedral smectite). This sample has been previously investigated using transmission powder infrared spectroscopy and

solid-state nuclear magnetic resonance by Yi et al. (2013). Importantly, the line width of the ν_1 PO₄ stretching band is similar to that observed on the synthetic sample, indicating that this natural sample is also very well ordered despite the occurrence of substituted structural carbonate groups. SEM observations reveal that it is mostly composed of aggregated sub-micrometric prisms of apatite (Fig. 1). For this sample, ATR spectra have been recorded on the as-received sample and after hand-grinding in ethanol during 15 and 60 minutes.

Scanning electron microscope (SEM) observations were performed using a Zeiss ultra 55 SEM operating in the secondary electron mode with a 2kV accelerating tension and a 2.5mm working distance.

The ATR-FTIR spectra have been recorded using a Quest ATR device (Specac) and a Nicolet 6700 FTIR spectrometer equipped with an Ever-Glo source, KBr beamsplitter and DTGS-KBr detector. Pure powder samples have been packed at the surface of the ATR crystal and the spectra have been recorded between 400 and 4000 cm⁻¹ by averaging 200 scans with a resolution of 1 cm⁻¹. For each sample, two different spectra have been recorded using a diamond or a Ge ATR crystal. The ATR spectra are reported in absorbance units.

3. Modeling strategy

The modeling strategy is closely related to that successfully used to model the ATR spectrum of a pure chrysotile aggregate by Balan et al. (2002).

The theoretical unpolarized ATR spectrum is obtained by computing the complex Fresnel reflection coefficient for s- and p- polarized waves and an incidence angle of 45° at a perfectly plane interface between the ATR crystal and the sample (Born and Wolf 1980). The ATR crystal is defined as a homogeneous medium of constant real refractive index (4 for Ge,

2.4 for diamond); whereas the sample is defined as an infinite homogeneous medium characterized by a frequency-dependent dielectric function.

The porous powder sample corresponds to a binary composite made of the substance of interest, i.e. apatite, and air (here approximated by a homogeneous medium with a unity refractive index). Apatite displaying a crystalline structure with hexagonal symmetry, its dielectric properties are described by a diagonal uniaxial dielectric tensor with a parallel and two identical perpendicular components, noted $\varepsilon_{\parallel}$ and $\varepsilon_{\perp}$, respectively. Numerous models exist that provide the approximate dielectric function of a composite sample from its individual constituents in a quasi-static approach (e.g. Sihvola and Kong 1988; Spanier and Herman 2000; Kendrick and Burnett 2016). They neglect scattering losses and can be straightforwardly extended to frequency-dependent dielectric properties when the characteristic size of individual particles is much smaller than the wavelength of the electromagnetic radiation, a condition usually approximately satisfied in mid-infrared spectroscopy ($\lambda_{\text{IR}} > 10 \mu\text{m}$). One of the simplest model (here referred to as MG model) was proposed by J.C. Maxwell Garnett (Garnett 1904), in which the particles of one of the constituents are considered as embedded in a homogeneous medium, corresponding to the other constituent for binary samples. In the MG model, the two constituents are therefore not treated in a symmetrically equivalent way. The MG model is closely related to the Clausius-Mossotti derivation of the dielectric constant of molecular substances from the polarizability of molecules. For a uniform orientation of particles, the dielectric properties of the composite sample are isotropic. For spherical particles, the MG effective dielectric function ε_{MG} of a uniaxial substance embedded in an isotropic matrix is given by the following expression (Levy and Stroud 1997):

$$\varepsilon_{\text{MG}} = \varepsilon_h + 3f\varepsilon_h \frac{(\varepsilon_{\perp} + 2\varepsilon_h)(\varepsilon_{\parallel} - \varepsilon_h) - 2\varepsilon_h(\varepsilon_{\parallel} - \varepsilon_{\perp})}{(1-f)(\varepsilon_{\perp} + 2\varepsilon_h)(\varepsilon_{\parallel} + 2\varepsilon_h) + f\varepsilon_h(\varepsilon_{\perp} + 2\varepsilon_{\parallel} + 6\varepsilon_h)} \quad (1)$$

where f is the volume fraction occupied by the particles and ε_h is the dielectric constant of the host matrix. The MG model is well suited to describe the dielectric properties of a sample in which a low amount of one of the constituents is diluted in a non-absorbing matrix, such as the KBr pellets used for traditional transmission experiments. Note however that the MG model leads to average properties of the pure material for $f \rightarrow 1$. It also provides physically meaningful results when the sample micro-structure ensures that the dipolar interactions between neighboring particles cancel out, even at intermediate volume fractions. For example, this was the case of the closely packed nanotube aggregates corresponding to the chrysotile sample investigated by Balan et al. (2002). However, for arbitrary micro-structures and intermediate volume fractions, the particles of one of the constituents cannot be considered as fully isolated and embedded in a homogeneous matrix of the other constituent. Correspondingly, an alternative model was used in which each individual particle of a mixture is considered as embedded in an effective medium (Bruggeman 1935, Landauer 1978). At variance with the MG model, all constituents are treated in a symmetrically equivalent way. In this model, the effective dielectric properties of a binary sample consisting in a mixture of an isotropic phase and ellipsoidal particles of a uniaxial phase are defined by the self-consistent relation (Walker and Scharnberg 1990; Sushko 2009):

$$f \left[2 \frac{(1-g_{//})(\varepsilon_{Br} - \varepsilon_{\perp})}{(1+g_{//})\varepsilon_{Br} + (1-g_{//})\varepsilon_{\perp}} + \frac{g_{//}(\varepsilon_{Br} - \varepsilon_{//})}{(1-g_{//})\varepsilon_{Br} + g_{//}\varepsilon_{//}} \right] + 3(1-f) \frac{\varepsilon_{Br} - \varepsilon_h}{2\varepsilon_{Br} + \varepsilon_h} = 0 \quad (2)$$

In this expression, $g_{//}$ is the depolarization factor along the particle revolution axis and $(1-g_{//})/2$ corresponds to the depolarization factor in perpendicular directions (for spherical particles, $g_{//} = 1/3$). The corresponding polynomial equation displays three complex solutions

among which that with a positive imaginary part has to be selected. A detailed expression of the polynomial coefficients and solutions is given in the annexes.

The uniaxial frequency-dependent dielectric properties of fluorapatite have been modeled with a Drude-Lorentz expression of the coefficients of its diagonal dielectric tensor:

$$\varepsilon_{\alpha}(\omega) = \varepsilon(\infty) + \sum_i \frac{A_{i,\alpha}^2}{(\omega_i^2 - \omega^2 - i\omega\Gamma_i)} \quad (3)$$

where α refers to the Cartesian axis, $\varepsilon(\infty) = 2.65$ is the electronic dielectric tensor of apatite, here considered to be real, constant and isotropic, ω_i are the vibrational transverse optical (TO) frequencies of IR-active modes, $A_{i,\alpha}^2$ are effective mode oscillator strengths, and Γ_i is a damping parameter. The frequencies of vibrational modes have been experimentally determined on fluorapatite single-crystals in previous works (e.g. Klee 1970, Adams and Gardner 1974, Leroy et al. 2000). As slight differences occur between the previous experimental determinations, the frequencies used in the present work are listed in Table 1. The mode oscillator strengths used in the present work are those previously obtained by Balan et al. (2011) on fluorapatite using first-principles quantum mechanical calculations. The damping parameter Γ_i describes the homogeneous broadening of vibrational bands and has been arbitrarily sets to 8 cm^{-1} for all the modes.

3. Results

a- Comparative properties of the MG and Bruggeman effective models

As anticipated in their presentation, the two effective medium models used in the present work display very distinct properties as a function of the volume fraction of the constituents. This is particularly evident when considering a model isotropic substance displaying a single resonance in its frequency-dependent complex dielectric function $\epsilon_S = \epsilon_S' + i \epsilon_S''$ (Figure 2). In the limit of a vanishingly small damping coefficient, the maximum of ϵ_S'' defines the transverse optical (TO) frequency; whereas the longitudinal optical (LO) frequency is defined by $|\epsilon_S|=0$. The second medium of the binary composite is characterized by its dielectric constant ϵ_h . The relation (1) for the isotropic MG model becomes:

$$\frac{\epsilon_{MG} - \epsilon_h}{\epsilon_{MG} + 2\epsilon_h} = f \frac{\epsilon_S - \epsilon_h}{\epsilon_S + 2\epsilon_h} \quad (4)$$

The MG dielectric function (Figure 2) still presents a single resonance, which progressively shifts toward the resonance frequency of the pure material when the volume fraction increases. In the limit of low volume fraction and vanishingly small damping parameter, the resonance occurs when $\epsilon_S' = 2 \epsilon_h$. For this peculiar condition, the real part of the denominator in Eq. 4 cancels out and the resonance coincides with the surface mode of the isolated sphere. The intensity of the resonance is not linearly related to the volume fraction of the composite but displays a stronger than linear decrease when the volume fraction decreases. Note that an alternative cavity-Maxwell-Garnett model (cavity-MG) can be built by considering air cavities embedded in an isotropic solid. In this case (not shown), two discrete resonances are observed at the TO and LO frequencies of the pure bulk substance.

The symmetric Bruggeman model leads to the following isotropic relation:

$$(1-f) \frac{\epsilon_h - \epsilon_{Br}}{\epsilon_h + 2\epsilon_{Br}} + f \frac{\epsilon_S - \epsilon_{Br}}{\epsilon_S + 2\epsilon_{Br}} = 0 \quad (5)$$

In the limit cases $f \rightarrow 0$ or $f \rightarrow 1$, the Bruggeman dielectric function is identical to the MG one, with a single resonance at the characteristic frequency of the isolated sphere or of the pure bulk material, respectively. In contrast, the line shape of the imaginary part of the dielectric function at intermediate volume fractions significantly broadens and becomes asymmetric, instead of displaying a single resonance at intermediate frequencies. For volume fractions above 1/3, the lines extend between the TO and LO frequencies with a relatively higher intensity on the TO frequency side.

These peculiar shape and dependence on volume fraction can be understood by analyzing the mathematical properties of the model. More specifically, the Bruggeman dielectric function has the properties of a Herglotz function, i.e. it is analytic and maps the upper half complex plane ($(\epsilon_s/\epsilon_h)''$ is positive) to itself (ϵ_{Br}'' is also positive) (Cherkaeva and Golden 1998). In the present case, it can be shown that the imaginary part of the Bruggeman dielectric function consists in the sum of two terms (see Annexes). One is proportional to $(3f-1)\epsilon_s''$ and corresponds to the resonance at the TO frequency observed for $f > 1/3$. The second one mixes the imaginary and real parts of the dielectric function of the material and leads to the broad feature which extends to the LO frequency. For $f < 1/3$, $(3f-1)\epsilon_s''$ is negative. As the imaginary part of the Bruggeman dielectric function is positive, this term cancels out with the second one. This explains the significant change in the shape of the Bruggeman dielectric function at $f = 1/3$. This volume fraction coincides with the percolation threshold of the symmetric Bruggeman model. For conducting particles inserted in a dielectric matrix, such as cermet compounds, it corresponds to the volume fraction above which the sample properties change from the insulating to the conducting regime (e.g. Gadenne et al. 1989).

b- Experimental ATR spectra of apatite

The ATR spectrum of the synthetic fluorapatite sample (Fig. 3), reported in absorbance units, displays the usual absorption bands ascribed to the vibrational modes of phosphates groups. The bands associated with the ν_3 PO₄ vibrations (anti-symmetric stretch) are observed between 1000 and 1150 cm⁻¹. The weak and narrower ν_1 PO₄ band (symmetric stretch) is observed at ~966 cm⁻¹. The bands associated with the ν_4 PO₄ vibrations (bending modes) are observed between 540 and 620 cm⁻¹. As expected, the ATR spectra depend on the refractive index of the ATR crystal and significant variations are observed between Ge and diamond ATR spectra, with a much lower intensity for the most refractive material (here, Ge with $n_{\text{Ge}} = 4$). ATR spectra also differ from the transmission spectrum measured and modeled by Balan et al. (2011) using the traditional method i.e. dilution of apatite particles in a KBr matrix (Fig. 3). These variations not only consist in changes in spectral intensities but also affect the position and the width of the bands. Interestingly, the line shapes are significantly asymmetric. In the Ge spectrum, the band at 1095 cm⁻¹ extends to ~1130 cm⁻¹, whereas the intense ν_3 PO₄ band at 1030 cm⁻¹ extends to ~1090 cm⁻¹. A significant asymmetry is also observed on the ν_4 PO₄ band at ~600 cm⁻¹. In the diamond ATR spectrum, the intense ν_3 PO₄ band appears broadened toward lower wavenumbers and its maximum is shifted to 1017 cm⁻¹.

The ATR spectrum of the natural carbonate-fluorapatite displays similar features (Fig. 4). Additional small bands related to structural carbonates (ν_2 CO₃ bending mode at ~865 cm⁻¹) and associated quartz impurities (462, 780 and 799 cm⁻¹) are observed. The Ge and diamond ATR spectra of the carbonate-fluorapatite are significantly less intense than the respective spectra of synthetic fluorapatite. Sample grinding tends to increase the spectrum intensity. However, the dependence of the Ge and diamond spectrum as a function of grinding time is different. For diamond ATR, a moderate grinding provides a significant recovery of spectral intensity, which is not improved by further grinding; whereas an increase of the Ge ATR intensity is only observed for the longest grinding time (60 min). This observation can

be linked to a larger penetration depth for the diamond ATR than for the more refractive Ge one. The strong bands are also broader than in the pure fluorapatite spectrum. In the diamond ATR spectrum, the ν_3 PO₄ band extends from 900 to 1100 cm⁻¹. The overlap with the weak band related to the ν_1 PO₄ mode leads to a very asymmetric narrow resonance. Its shape shares some similarity with the derivative of a Lorentzian function, suggesting that the real part of the dielectric function related to this resonance significantly contributes to the spectrum. In contrast, this resonance appears with a standard peak shape in the Ge spectrum, in which ν_1 and ν_3 PO₄ bands do not overlap.

c- Theoretical ATR-FTIR spectra of apatite

Starting from a single dielectric function for pure fluorapatite, defined by the parameters specified in Table 1, and for a given effective medium model, the theoretical Ge and diamond ATR spectra are obtained by varying the dielectric constant of the ATR crystal. In this case, the only free parameter that can be used to optimize the agreement between the model ATR spectra and their experimental counterparts is the volume fraction. For the synthetic sample, a good agreement is achieved by using a fraction of 0.5 (Fig. 3) which is reasonable for an ordinary powder sample. For comparison, the ratio of the apparent density of quartz sand to that of pure quartz is 0.577 (Bisal and Hinman 1972).

For the two effective medium models, the shape of the theoretical Ge ATR spectra is very close to that of the imaginary part of the corresponding dielectric functions. The most significant variation are minor shifts (< 4 cm⁻¹) of the most intense ν_3 PO₄ band toward lower wavenumbers. It is noteworthy that the intensity of the modeled ATR spectra compares well with the raw experimental measurements. As anticipated, the MG model leads to a series of narrow peaks related to the various resonances of the apatite dielectric tensor, accounting

poorly for the experimental asymmetric line shapes. In contrast, the specific line shapes observed for intermediate volume fractions in the Bruggeman model compare well with their experimental counterparts (Fig. 3). More specifically, they reproduce the peak maximum at low frequency, corresponding to the TO frequencies of the apatite vibrational modes, and the broad absorption feature observed on the high-frequency side. As shown in the analysis of the Bruggeman model, this feature extends up to the LO frequency of apatite vibrational modes. In the ν_4 PO_4 range, the Bruggeman model also reproduces the asymmetric line shape of the band at 600 cm^{-1} and the line splitting observed at 564 and 578 cm^{-1} . Considering the diamond ATR spectrum, its shape differs from that of the imaginary part of dielectric functions by the fact that the most intense ν_3 PO_4 band is broadened toward the lower wavenumbers for the two effective medium models. This effect on the line shape is therefore specifically ascribed to the ATR experimental configuration. As for the Ge ATR spectrum, the best agreement on the line positions and spectral shape is obtained by using the Bruggeman dielectric model, both in the ν_3 PO_4 and ν_4 PO_4 ranges. The experimentally observed position of the bands, and more importantly, the relative changes between the spectra and the relative shifts with respect to the resonances of the dielectric tensor of pure bulk apatite are well-reproduced by the model. The slight but systematic under-estimation of the position of the weak ν_1 PO_4 band (by 4 cm^{-1}) suggests that the TO frequency used to model the low-frequency dielectric tensor of apatite (Table 1) was slightly underestimated. Note that attempts using the cavity-MG model did not provide a better modeling of the observed peculiar line shapes than the standard MG model. Indeed, this model also leads to a series of discrete peaks instead of the broad features extending between TO and LO frequencies for intermediate volume fractions.

The ATR spectrum of the natural sample of carbonate-fluorapatite was modeled using the same apatite dielectric tensor as for the synthetic one (Fig. 5). No attempt has been made

to take into account the small spectroscopic signal related to the structural carbonate ions. Importantly, the line width was not modified, which is consistent with the small ν_1 PO₄ linewidth measured in previous transmission IR spectroscopy studies (Balan et al. 2011; Yi et al. 2013). This small linewidth attests to the weakness of the inhomogeneous broadening related to microscopic disorder parameters. A moderate increase in the apatite volume fraction significantly increases the asymmetric broadening of the intense ν_3 PO₄ band in the diamond ATR spectrum. It overlaps with the narrow ν_1 PO₄ band, leading to a shape similar to the derivative of a Lorentzian, consistent with the experimental observations. However, the theoretical ATR spectra do not account for the significant decrease in spectral intensity observed from the synthetic to the natural sample. The modeled diamond and Ge spectra consistently overestimate their experimental counterparts by a factor of approximately two (Fig. 5). This points to a failure in the model of the sample effective dielectric properties. A microscopic sample heterogeneity, e.g. related to carbonate substitutions, cannot explain such discrepancy because it is not supported by the observed ν_1 PO₄ linewidth. In contrast, heterogeneities occurring at a length scale larger than the characteristic length scale of the spectroscopic measurements, typically the IR wavelength, could account for the observed discrepancy. We particularly suspect spatial fluctuations in the solid/air fraction to account for the observed discrepancy.

4. Concluding remarks

The present modeling of apatite ATR spectra demonstrates the prominent role of macroscopic electrostatic effects over fine details of the microscopic structure in the line shape of strong ATR bands. A relatively simple effective medium modeling of the dielectric properties of the sample, based on the Bruggeman approximation, already provides important

insights into these respective contributions. It points to the key role of the volume fraction of the constituents, which is the only free parameter of the model. More sophisticated models would most probably improve the agreement between theory and experiment (e.g. Yagil et al. 1995; Spanier and Herman 2000) but at the price of a corresponding increase in the number of free parameters.

As previously discussed in Balan et al. (2011), macroscopic electrostatic effects have a weaker influence on the shape and position of the weak absorption bands of apatite. These bands, although often neglected and sometimes more difficult to record, are expected to provide a better insight into the short-range order in crystal structures. However, the present modeling indicates that their overlap with stronger resonances in the diamond ATR spectrum significantly affects their shape by mixing the real and imaginary parts of the dielectric function.

Finally, it appears that the relation between ATR and transmission spectra is not straightforward even though commercial softwares provide ATR correction algorithms to facilitate the comparison. The different composite nature (substance/air vs substance/KBr) and more importantly the different dilution state (pure powder vs isolated particles) of the measured samples affect their macroscopic electrostatic properties and result in changes in the position, shape and relative intensity of strong absorption bands. This can significantly bias the comparison between ATR and transmission spectra despite the fact that the analyzed bulk substance is characterized by a single dielectric function. Importantly, the differences observed in the analyzed samples are explained by the long-range character of electrostatic interactions affecting (sub-)micronic particles. They differ in essence from potential surface-induced relaxation effects which occur at the atomic scale and can become dominant in nanometer-sized particles (Waychunas and Zhang 2008).

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Annexes:

1- Complex solutions of the uniaxial Bruggeman model

The Bruggeman relation is (Eq. 2):

$$f \left[2 \frac{(1-g_{//})(\varepsilon_{Br} - \varepsilon_{\perp})}{(1+g_{//})\varepsilon_{Br} + (1-g_{//})\varepsilon_{\perp}} + \frac{g_{//}(\varepsilon_{Br} - \varepsilon_{//})}{(1-g_{//})\varepsilon_{Br} + g_{//}\varepsilon_{//}} \right] + 3(1-f) \frac{\varepsilon_{Br} - \varepsilon_h}{2\varepsilon_{Br} + \varepsilon_h} = 0$$

The solutions of the equation are the roots of a cubic polynomial with complex coefficients:

$$ax^3 + bx^2 + cx + d = 0$$

with

$$\begin{aligned} a &= -3 - f + 6fg_{//} + 3g_{//}^2 - 9fg_{//}^2 \\ b &= 3(\varepsilon_h - \varepsilon_{\perp}) + (7\varepsilon_{\perp} - 5\varepsilon_h)f + (6\varepsilon_{\perp} - 3\varepsilon_{//})g_{//} + fg_{//}(3\varepsilon_h + \varepsilon_{//} - 16\varepsilon_{\perp}) - 3g_{//}^2(\varepsilon_{//} + \varepsilon_{\perp} + \varepsilon_h) + 9(\varepsilon_{//} + \varepsilon_{\perp})fg_{//}^2 \\ c &= \varepsilon_h\varepsilon_{\perp}(3-f) + (3\varepsilon_h\varepsilon_{//} - 6\varepsilon_h\varepsilon_{\perp} - 3\varepsilon_{//}\varepsilon_{\perp})g_{//} + (9\varepsilon_{//}\varepsilon_{\perp} + \varepsilon_h\varepsilon_{\perp} - 4\varepsilon_h\varepsilon_{//})fg_{//} \\ &\quad + 3g_{//}^2(\varepsilon_h\varepsilon_{//} + \varepsilon_h\varepsilon_{\perp} + \varepsilon_{//}\varepsilon_{\perp}) - 9\varepsilon_{//}\varepsilon_{\perp}fg_{//}^2 \\ d &= 3\varepsilon_h\varepsilon_{//}\varepsilon_{\perp}(g_{//} - g_{//}^2) \end{aligned}$$

The three roots are:

$$\begin{aligned} x_1 &= -\frac{b}{3a} - \frac{2^{1/3}(-b^2 + 3ac)}{3a(-2b^3 + 9abc - 27a^2d + \sqrt{-4(b^2 - 3ac)^3 + (2b^3 - 9abc + 27a^2d)^2})^{1/3}} \\ &\quad + \frac{(-2b^3 + 9abc - 27a^2d + \sqrt{-4(b^2 - 3ac)^3 + (2b^3 - 9abc + 27a^2d)^2})^{1/3}}{3a2^{1/3}} \\ x_2 &= -\frac{b}{3a} + \frac{(1+i\sqrt{3})(-b^2 + 3ac)}{3a2^{2/3}(-2b^3 + 9abc - 27a^2d + \sqrt{-4(b^2 - 3ac)^3 + (2b^3 - 9abc + 27a^2d)^2})^{1/3}} \\ &\quad - \frac{(1-i\sqrt{3})(-2b^3 + 9abc - 27a^2d + \sqrt{-4(b^2 - 3ac)^3 + (2b^3 - 9abc + 27a^2d)^2})^{1/3}}{6a2^{1/3}} \\ x_3 &= -\frac{b}{3a} + \frac{(1-i\sqrt{3})(-b^2 + 3ac)}{3a2^{2/3}(-2b^3 + 9abc - 27a^2d + \sqrt{-4(b^2 - 3ac)^3 + (2b^3 - 9abc + 27a^2d)^2})^{1/3}} \\ &\quad - \frac{(1+i\sqrt{3})(-2b^3 + 9abc - 27a^2d + \sqrt{-4(b^2 - 3ac)^3 + (2b^3 - 9abc + 27a^2d)^2})^{1/3}}{6a2^{1/3}} \end{aligned}$$

Among these three roots, only that with a positive imaginary part has to be selected.

2- Some mathematical properties of the Bruggeman model

The solutions of Eq. 5 are the roots of a quadratic polynomial equation with complex coefficients:

$$x^2 + 2bx + c = 0$$

with $x = \frac{\varepsilon_{Br}}{\varepsilon_h}$, $b = -\frac{(3f-1)\frac{\varepsilon_s}{\varepsilon_h} + (2-3f)}{4}$ and $c = -\frac{\varepsilon_s}{2\varepsilon_h}$. ε_h is the dielectric constant of the

host and $\varepsilon_s = \varepsilon_s' + i\varepsilon_s''$ is the complex dielectric function of the substance.

The corresponding solutions are: $x_{\pm} = -b \pm \sqrt{\Delta}$ with $\Delta = b^2 - c$. We will now show that the imaginary part of the solution x_+ is positive.

The imaginary part of $\sqrt{\Delta}$ is equal to $\sqrt{(|\Delta| - \Delta_r)}/2$ where $\Delta = \Delta_r + i\Delta_i$. The imaginary part of x_+ is thus:

$$\text{Im}(x_+) = -b_i + \sqrt{\frac{(|\Delta| - \Delta_r)}{2}} \quad \text{Eq. A1}$$

where $b = b_r + ib_i$

We now show that the absolute value $|b_i|$ is smaller than $\sqrt{\frac{(|\Delta| - \Delta_r)}{2}}$. This is equivalent to

determining that $|\Delta|^2 - (2b_i^2 + \Delta_r)^2$ is positive. Replacing the coefficients Δ and b_i by their expressions as a function of the dielectric function of the substance and matrix, we obtain:

$$|\Delta|^2 - (2b_i^2 + \Delta_r)^2 = \frac{9}{8} \left(\frac{\varepsilon_s''}{\varepsilon_h} \right)^2 f(1-f) \text{ which is indeed always positive for } 0 \leq f \leq 1.$$

As a consequence, $\text{Im}(x_+)$ is also positive. This property is even stronger than the Herglotz property because ε_s'' is not assumed to be positive. In particular for $f = 0$, $\text{Im}(x_+) =$

$(\varepsilon_s'' + |\varepsilon_s''|)/2 \geq 0$ even if ε_s'' is negative. The condition that ε_h is real and positive thus implies that $\text{Im}(\varepsilon_{Br})$ is positive, for any function ε_s .

As $-b_i = (3f - 1) \frac{\varepsilon_s'}{4\varepsilon_h}$ and $\varepsilon_s'' > 0$, the first term in Eq. A1 is positive when $f > 1/3$. This contribution is thus proportional to the dielectric function of the pure bulk substance and always present. For $f < 1/3$, this contribution becomes negative but since $\text{Im}(x_+) \geq 0$, this negative contribution is compensated for by the second term in Eq. A1. The ε_s'' peak then disappears completely from the spectrum. Equation A1 has another interesting consequence. In a region where $|\Delta_i|$ is smaller than $|\Delta_r|$, there are two regimes: if $\Delta_r \geq 0$, then $|\Delta| - \Delta_r \approx \Delta_i^2 / (2\Delta_r)$ is small and the first term of Eq. A1 is dominant for $f > 0.3$; if $\Delta_r \leq 0$, then $|\Delta| - \Delta_r \approx 2|\Delta_r|$ and the deformation of the spectrum due to the second term of Eq. A1 becomes large. In other words, the deformation of the spectrum of apatite for $f > 0.3$ occurs when Δ has a large negative real part.

We now examine the properties of the imaginary part of the solution for $f=0.5$ and $\varepsilon_h=1$, i.e. the parameters used to model the spectrum of synthetic fluorapatite (Fig. 6). In this case, $b^2 - c = \varepsilon_s(\varepsilon_s + 34) + 1$. For a vanishingly small damping parameter and out of the TO frequency, ε_s is real: $\varepsilon_s \approx \varepsilon_s'$. Therefore, only the second term contributes to the imaginary part of the solution when $b^2 - c$ is negative. This condition is met for ε_s' approximately ranging between -34 and zero, i.e. on the high energy side of the TO frequency and up to about the LO frequency. For ε_s' below -34, the imaginary part of ε_s becomes dominant and the spectrum is proportional to the dielectric function of the pure bulk substance.

Figures:

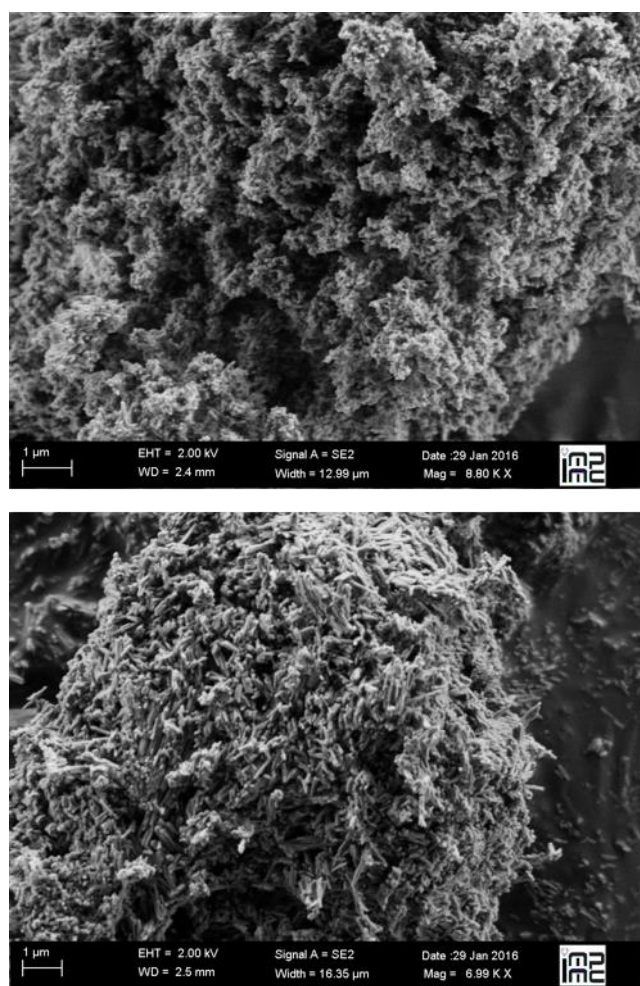


Figure 1: SEM micrographs of synthetic fluorapatite (top) and natural carbonate-fluorapatite (bottom).

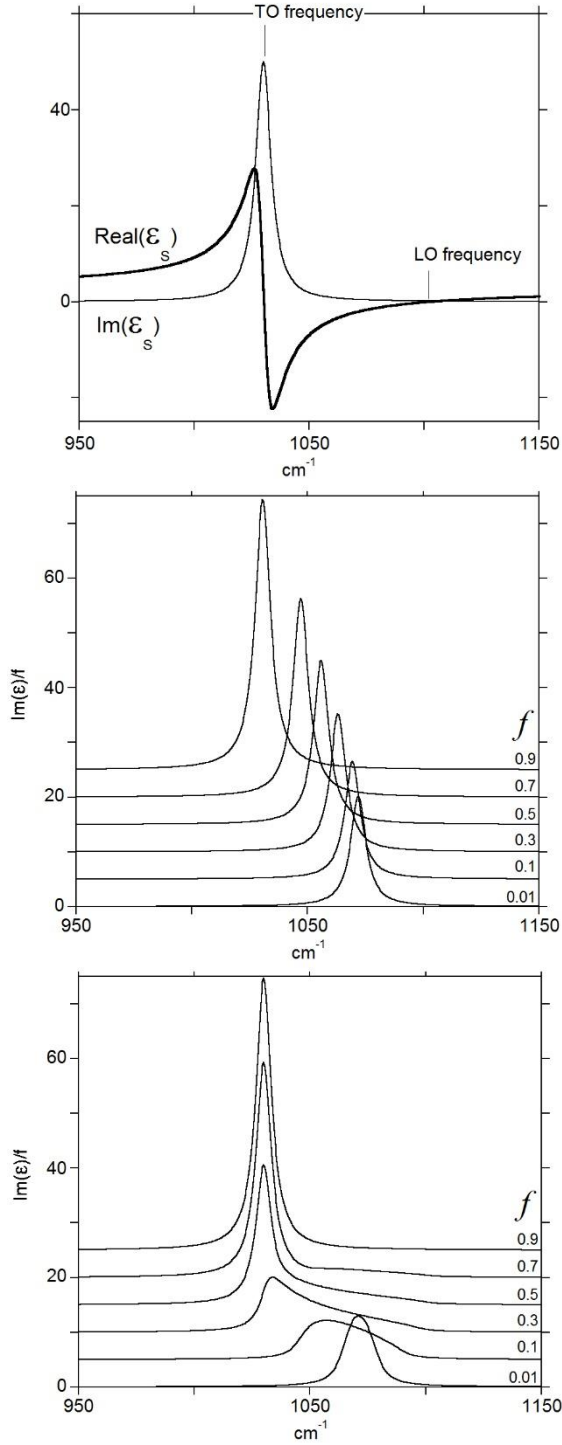


Figure 2: Real and imaginary parts of a model single-resonance dielectric function (top) and corresponding imaginary part of the effective dielectric function in the Maxwell-Garnett (middle) and Bruggeman (bottom) models. The parameters of the model dielectric function (Eq. 3) were $\omega_0=1030 \text{ cm}^{-1}$, $A_\alpha=642$, $\Gamma=8 \text{ cm}^{-1}$, $\epsilon(\infty)=2.65$. The effective dielectric functions have been divided by the corresponding volume fraction f value, reported on the right side.

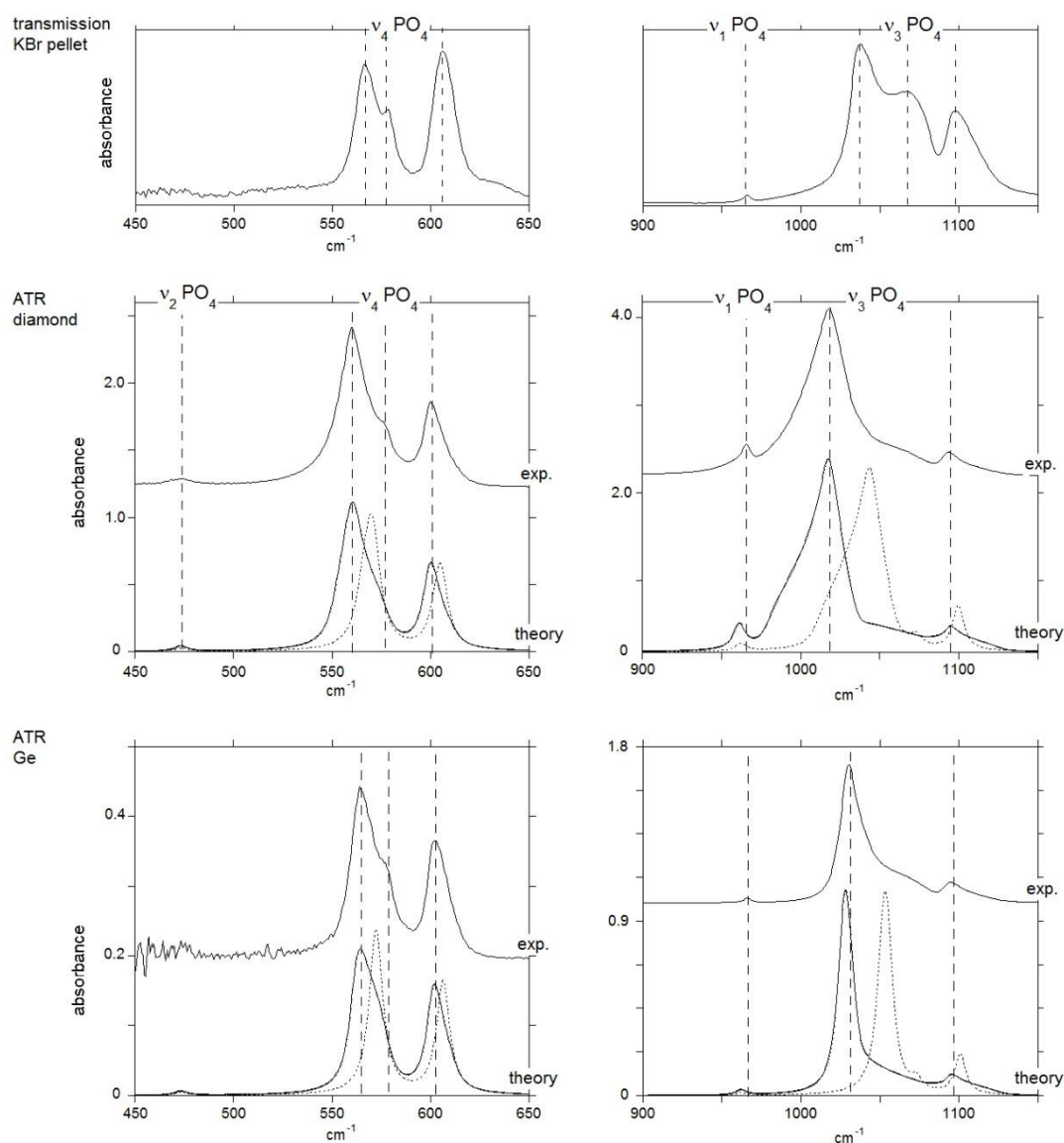


Figure 3: Experimental and modeled diamond (middle) and Ge (bottom) ATR spectra of synthetic fluorapatite. Experimental spectra have been vertically shifted for clarity. Maxwell-Garnett and Bruggeman modeled ATR spectra are displayed as dotted and solid lines respectively. The transmission spectrum recorded by Balan et al. (2011) on the same sample after dilution in KBr is reported for comparison (top). Note the significant changes affecting strong absorption bands. Vertical dotted lines serve as guide to the eye.

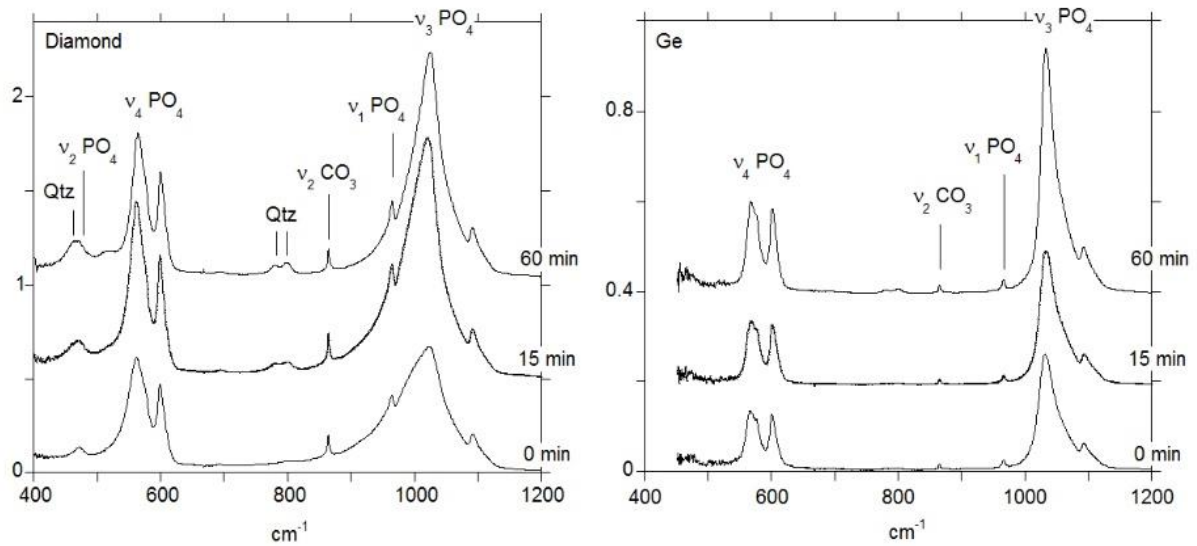


Figure 4: Experimental ATR spectra of natural carbonate-fluorapatite samples. Grinding times are reported on the right side for each spectrum. Grinding time affects the intensity of Ge and diamond spectra differently, suggesting an interplay between the sample spatial heterogeneity and characteristic IR penetration depth in the ATR crystals.

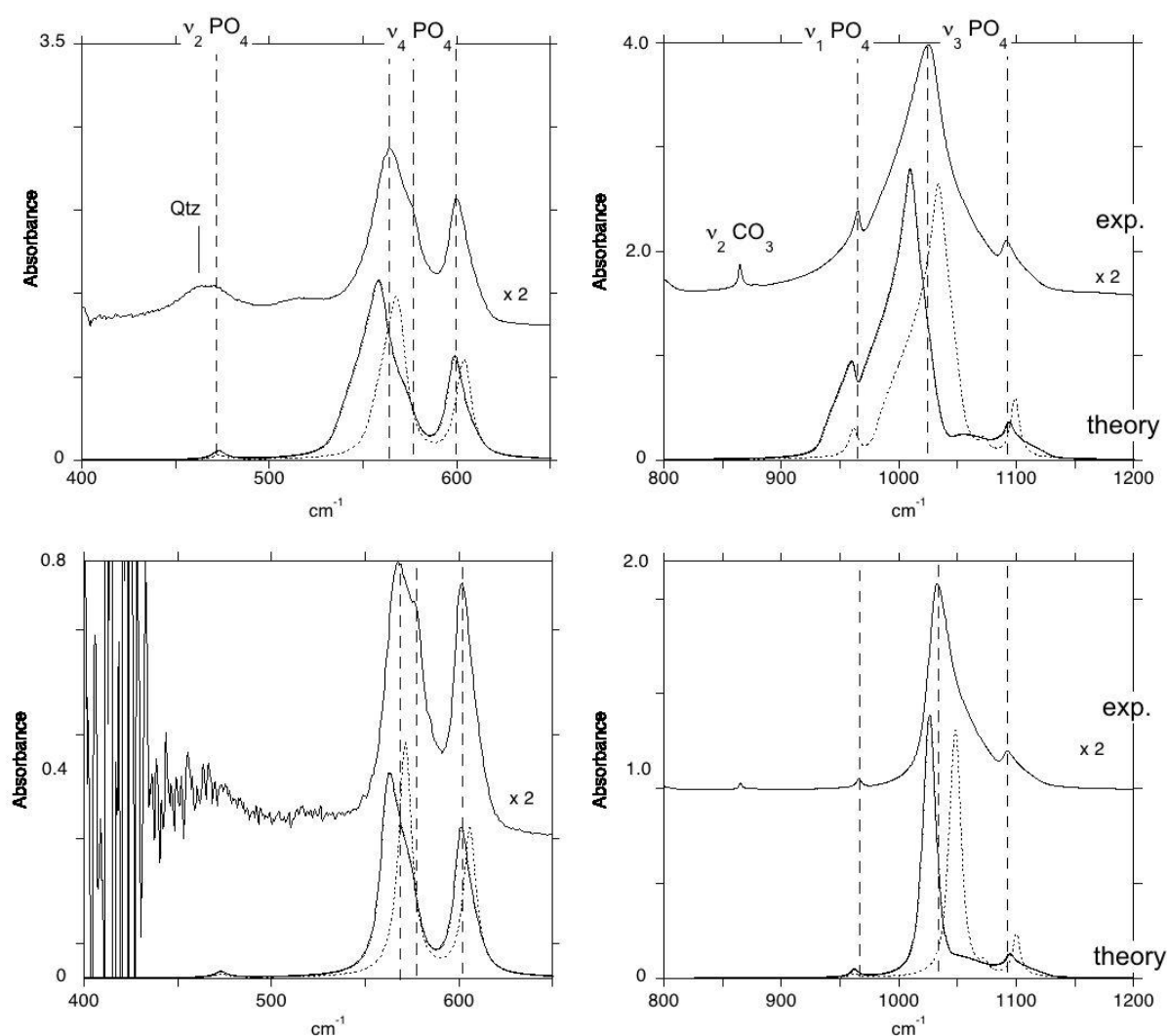


Figure 5: Experimental and modeled diamond (top) and Ge (bottom) ATR spectra of natural carbonate-fluorapatite. Experimental spectra have been vertically shifted for clarity and experimental intensities were multiplied by 2. Note that the wavenumber ranges slightly differ from those on figure 3. Maxwell-Garnett and Bruggeman modeled ATR spectra are displayed as dotted and solid lines respectively.

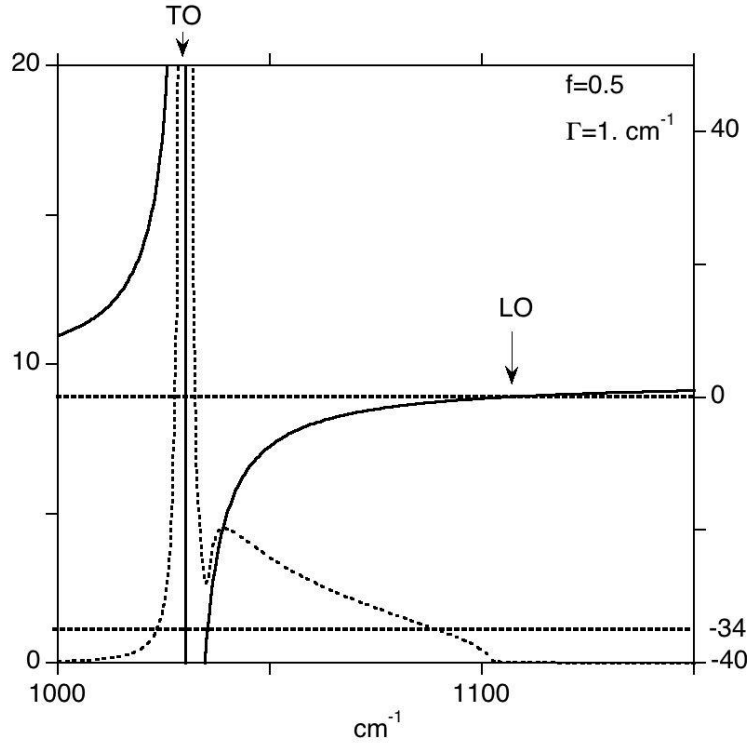


Figure 6: Real part of the model dielectric function (bold line, right axis) and imaginary part of the corresponding effective dielectric function (dotted line, left axis). The small damping parameter (1 cm^{-1}) makes it possible to discriminate the two superposed contributions for a volume fraction of 0.5. The thin contribution at TO frequency displays an almost Lorentzian shape proportional to the imaginary part of the model dielectric function. In contrast, the broad contributions at higher frequency is observed when the real part of the model dielectric function takes values between ~ -34 and ~ 0 (LO frequency), indicated by the horizontal dotted lines.

Table 1: Parameters of the model dielectric tensor of apatite (Eq. 3).

Symmetry	Mode	ω_i (cm ⁻¹)	$A_{i,a}$ (arb. units) ⁴	Γ_i (cm ⁻¹)
A_u	ν_3	1030 ¹	665	8
	ν_4	560 ²	300	8
	ν_2	473 ³	45	8
E_{1u}	ν_3	1094 ¹	225	8
	ν_3	1034 ¹	630	8
	ν_1	962 ²	90	8
	ν_4	601 ²	215	8
	ν_4	575 ²	120	8

¹Adams and Gardner (1974) ²Klee (1970) ³Leroy et al. (2000) ⁴Balan et al. (2011)