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Rietveld crystal structure refinement of a natural rhombohedral grossular-andradite garnet from Serbia

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Four macroscopically visible B–E (rim) zones, within a large natural ferric grossular garnet of $\text{Gr}_{58-64}\text{Adr}_{36-42}\text{Sps}_2$ composition, are described by means of Rietveld refinements of the crystal structures in a series of six space groups, followed by comparative analysis of the R -values, site occupancy factors, and bond lengths and angles. The garnet crystallized in the rhombohedral $R\bar{3}c$ space group. Various polyhedral distortions and structural order-disorder variations between the zones studied are also described and discussed. The rhombohedral symmetry of the ferric grossular garnet analysed can be regarded as primary, residual strain being a secondary cause for its slight optical anisotropy.

Key words: ferric grossular garnet, XRPD, Rietveld method, crystal structure, $R\bar{3}c$ space group, order-disorder.

INTRODUCTION

Garnets are among the most widely occurring minerals on Earth. They are an important group of rock-forming minerals that are stable over a wide range of temperatures (up to 2000°C), pressures (up to ~25 GPa), and chemical composition. According to the International Mineralogical Association (IMA) nomenclature, the garnet supergroup includes all minerals isostructural with garnet regardless of what elements occupy the four atomic sites, i.e., the supergroup includes several chemical classes. The general formula for the garnet supergroup minerals is $\{X_3\}[Y_2](Z_3)\varphi_{12}$, where X , Y and Z refer to dodecahedral, octahedral, and tetrahedral sites, respectively, and φ is O, OH, or F. Most garnets are cubic, of space group $Ia\bar{3}d$ (no. 230), but two OH-bearing species (henritermierite and holtstamite) have tetragonal symmetry, space group, $I4_1/acd$ (no. 142), and their X , Z , and φ sites are split into more symmetrically unique atomic positions. The total charge at the Z site and symmetry are criteria for distinguishing groups, whereas the dominant-constituent and dominant-valency rules are critical in identifying species. Twenty-nine species belong to one of five groups: the tetragonal henritermierite group and the isometric bitikleite, schorlomite, garnet, and

berzeliite groups with a total charge at Z of 8 (silicate), 9 (oxide), 10 (silicate), 12 (silicate), and 15 (vanadate, arsenate), respectively (Grew et al., 2013).

Species of the grossular-andradite $\{\text{Ca}_3\}[\text{Al,Fe}^{3+}]_2(\text{Si}_3)\text{O}_{12}$ solid-solution binary series, occurring in a variety of geological environments, are known to appear in the form of zoned crystals (Deer et al., 2013; Tančić et al., 2012a and references therein). A substantial number of previously-observed grossular-andradite garnets exhibit optical anisotropy and non-cubic low symmetry in various space groups, such as: $\Gamma 1$ (Takéuchi et al., 1982; Allen and Buseck, 1988; Kingma and Downs, 1989; Frank-Kamenetskaya et al., 2007; Kobayashi et al., 2013; and Nakamura et al., 2016); $Fddd$ (Takéuchi et al., 1982; Gali, 1984; Frank-Kamenetskaya et al., 2007; Badar et al., 2016); $Fddd$ or $C2/c$ (Hirai and Nakazawa, 1986) and $R\bar{3}c$ or $Fddd$ (Tančić et al., 2020).

To the best of our knowledge, only three optically anisotropic and oscillatory zoned grossular-andradite garnet samples from Serbia have been characterized in more or less detail by Tančić et al. (2012a, 2020), Srećković-Batočanin et al. (2014) and Kostić et al. (2021), i.e. from skarns of Kopaonik Mt., Rogozna Mt. and Rudnik Mt., respectively.

The macroscopically-zoned garnet from Meka Presedla (Kopaonik Mt., Serbia) studied in this paper was previously characterized by Tančić et al. (2012a). It is slightly anisotropic throughout, without the presence of isotropic areas. Based on electron microprobe analysis (EMPA) median chemical compositions, considerable inversely related variation is observed only by the $\text{Al}^{3+}\text{--Fe}^{3+}$ substitutions in the octahedral site, and a $\text{Gr}_{58-64}\text{Adr}_{36-41}\text{Sps}_2$ composition was determined [mineral

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name symbols such as Adr, Grs and Sps refer to andradite, grossular, and spessartine, respectively (Warr, 2021). Such composition therefore defines this garnet more specifically as ferric grossular]. FT–IR and Raman bands are consistent with the chemical composition. All of the zones should be considered as anhydrous, formed under conditions of miscibility and without gaps. They represent solid solutions, rather than mixtures. Formation temperatures from ~600 to 720°C, and pressures of ~2–3 kbars, are derived. Among five possible causes for the slightly optical anisotropy, three were rejected, because: (1) the crystal is microscopically untwinned; (2) no rare earth elements (REE) were detected; and (3) there was no distribution of OH⁻ groups due to hydro-garnet substitution.

The ferric grossular core (zone A) with Grs_{64±1}Adr_{36±1}Sps₂ composition (Tančić et al., 2012a) was further crystallographically studied by Tančić et al. (2020). The procedure applied included detailed analysis of the powder diffraction patterns, and Rietveld refinements of the crystal structures in a series of 18 space groups and two mixtures, followed by the comparative analysis of the *R*-values, site occupancy factors (hereinafter *sof*'s), and bond lengths and angles. It was shown to be not cubic, neither as mono phase, nor as multiple phases in a mixture. Structural dissymmetrization occurred, and it crystallized in disordered rhombohedral *R*⁻3c or orthorhombic *Fddd* space groups, the first one being more probable. Besides establishing lower symmetry (5), which could be treated as primary cause, residual strain (4) was also not excluded as a possible secondary cause for its slight optical anisotropy.

In the present study, the four remaining, B–E, zones of the Grs_{58–64}Adr_{36–41}Sps₂ ferric grossular (Tančić et al., 2012a) with Grs_{62±1}Adr_{38±1}Sps₂ (B), Grs_{59±2}Adr_{41±2}Sps₂ (C), Grs_{58±2}Adr_{42±2}Sps₂ (D) and Grs_{58±1}Adr_{42±1}Sps₂ (E, rim) compositions, were analysed. Our primary goal was to perform an X-ray powder diffraction (XRPD) study of the crystal structure by the Rietveld method in the *R*⁻3c and *Fddd* space groups, i.e. to check what could be the true space group of its crystallization between these two. These results were compared with those for the *Ia*⁻3d, *R*⁻3, *C2/c* and *Γ*1 space groups. All of the results obtained were also further compared with those for zone A

(Tančić et al., 2020). For the space group definition it was mainly the chosen distances, *sof*'s and lower comparative *R*-values which were mostly, as argued in our previous studies, the decisive factors not only in vast majority of the low-symmetry grossular-andradite garnets (Tančić et al., 2020), but also in other characteristic natural mineral solid solutions with Al³⁺–Fe³⁺ substitutions over the octahedral sites (Tančić et al., 2012b). The XRPD method was used in order to avoid any kind of “Umweganregung” and/or “Aufhellung” (Rossmannith and Armbruster, 1995).

MATERIALS AND METHODS

As described by Tančić et al. (2012a), the macroscopically-zoned ferric grossular garnet investigated from the Meka Presedla locality (Kopaonik Mt., Serbia) was initially a spherical mega-crystal with a diameter of ~8 cm. It was cut into parallel slices ~0.5 cm thick. The largest slice, representing the central part, is shown in Figure 1. In this figure, the macroscopically-visible, different zones marked with solid lines and A–E signs are parallel to the rhombic dodecahedron (110), with a characteristic angle of 120°. Marked zones were carefully cut down, and then powdered and homogenized under alcohol in an agate mortar for several hours, in order to ensure good statistical distribution of the grains.

The samples from the B–E (rim) zones were taken in reflection mode in a bottom-loaded aluminum carrier. Diffraction data were collected on a Phillips PW1710 diffractometer equipped with a graphite monochromator (Cu Kα₁, λ = 1.540562 Å; Cu Kα₂, λ = 1.544390 Å) and a Xe-filled proportional counter, at room temperature (23°C). Divergence and receiving slits were fixed to 1° and 0.1 mm, respectively, and the generator was set up at 40 kV and 30 mA. The diffractometer alignment was properly checked using a reference material of powdered crystalline silicon, according to the manufacturer's user manual. Data were collected in scan-step mode, with 0.02° 2θ step, and between 4° and 135° 2θ and 12.9s step⁻¹ for collecting data for Rietveld structure refinements.

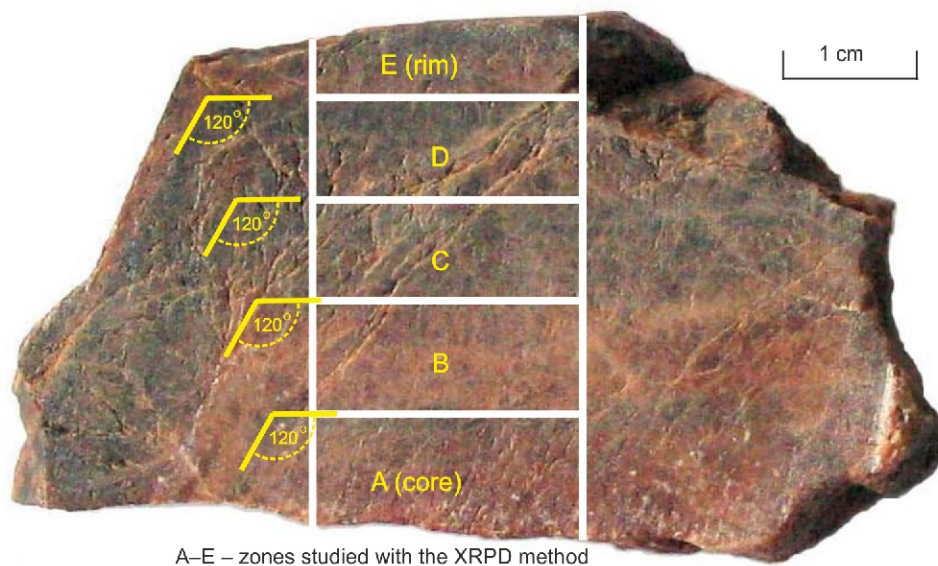


Fig. 1. The macroscopically-zoned large ferric grossular crystal from the Meka Presedla locality studied (Kopaonik Mt., Serbia)

Note that the macroscopically-visible different zones are of the rhombic dodecahedron (110) shape, with a characteristic 120° angle (modified from Tančić et al., 2012a)

RESULTS AND DISCUSSION

XRPD PATTERNS ANALYSIS

Checking what could be the true space group of the ferric grossular garnet studied started with the comparison and close inspection of the presence and nature of the reflections with Miller's indices 12 2 2 and 888 in the A–E zones. These clearly have diffraction peaks of different intensities and with more or less visible doublets and/or extra-broadening (Fig. 2B, C). On the other hand, other diffraction peaks lack visible doublets, such as the strongest 420 reflection (Fig. 2A), for example. According to the previous observations by Tančić et al. (2020), this seems to be of considerable importance, because it indicates that the garnet studied did not crystallize in the cubic system at all; neither as mono, nor as multiple phases as a mixture, because in this case the observed intensities (Y_{obs}) do not match with the calculated Bragg's positions (Y_{calc}) and *vice versa*. Also, among 18 space groups and two mixtures studied it was more specifically demonstrated that it cannot crystallize in the cubic $la\bar{3}d$, $\Gamma 43d$, $I4_132$, $P4_132$, $la\bar{3}$ and $I2_13$; tetragonal $I4_1acd$ and $I4_1/a$; and orthorhombic $Ibca$ and $Pbca$ space groups; or as cubic $la\bar{3}d$ & $la\bar{3}d$ and $P4_132$ & $la\bar{3}d$ mixtures. The remaining eight possibilities were among the rhombohedral $R\bar{3}c$ and $R\bar{3}$; orthorhombic $Fddd$; monoclinic $I2/a$, $I2/b$, $I2/c$ and $C2/c$; and triclinic $\Gamma 1$ space groups. Geometrical–mathematical calculations could initially indicate only the following six solutions: $R\bar{3}c$, $R\bar{3}$, $Fddd$, $I2/a$, $I2/c$ and $C2/c$ space groups, in very close agreement with previous observations. However, because it was shown by Rietveld crystal structure refinements that there were no significant differences between Al *sof's* and their corresponding Y–O distances over the octahedral Y sites, the $R\bar{3}$, $I2/a$, $I2/b$, $I2/c$, $C2/c$ and $\Gamma 1$ space groups were also excluded as possibilities, leaving only the $R\bar{3}c$ and $Fddd$ space groups (Tančić et al., 2020).

CRYSTAL STRUCTURE REFINEMENTS

Two types of crystal structure were determined using the Rietveld profile method and *Fullprof* program (Rodríguez-Carvajal, 1990): (1) zones B–D were refined as ferric grossular monophases; and (2) zone E was refined as a mixture of two phases, i.e. ferric grossular and scapolite, because in this zone scapolite is most abundant (Tančić et al., 2012a). Crystal structure refinements of the ferric grossular garnet were performed following the procedure described by Tančić et al. (2020), in order to achieve all the necessary conditions for data comparison between the A–E zones. It started from the cubic $la\bar{3}d$ space group, which was nominally assigned for garnet group crystallization (Grew et al., 2013). The pseudo-Voigt peak shape function was assumed. In the last cycle of the refinements, 117 parameters were varied: 100 parameters for the background description, 11 profile parameters (one scale factor, one Eta ($p-v$) parameter, one overall temperature parameter, three parameters for the description of the halfwidths, one parameter for preferred orientation, two asymmetry parameters, one mixing parameter and one zero point), one parameter for the a_0 unit cell dimension, and five parameters for the three free oxygen atom coordinates, and two *sof's* at the dodecahedral and octahedral sites. As is well known, symmetry lowering from the $la\bar{3}d$ space group to the other five space groups studied increases the number of ions in special and general positions, and the parameters of their atom coordinates, unit cell dimensions and *sof's*. For zone E, an additional 23 parameters were varied for scapolite in the $I4/m$ space group

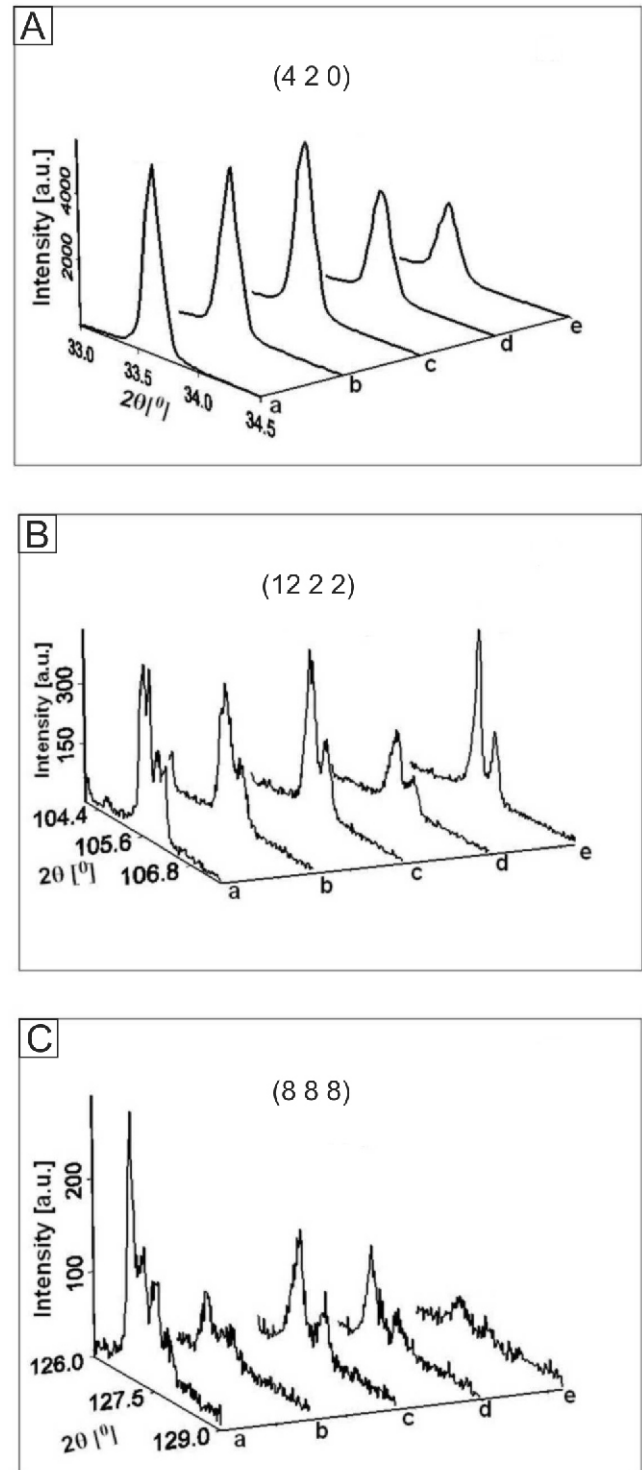


Fig. 2. Comparative powder diffraction patterns of the A–E zones for the three different angle regions and with reflections marked with hkl 's belonging to the $la\bar{3}d$ space group: A – 420; B – 12 2 2; C – 888

Marks: a-zone A; b-zone B; c-zone C; d-zone D; e-zone E

(Sherriff et al., 2000): one scale factor, one mixing parameter, two lattice constant parameters, atomic parameters for two free coordinates of the Ca^{2+}/Na^+ cations, five free coordinates of the Si^{4+}/Al^{3+} cations, ten free coordinates of the oxygen ions, and two *sof's* for the Ca^{2+}/Na^+ and Cl^-/C^{2-} positions.

Table 1

Profile and structural parameters, unit cell dimensions, and reliability factors for the $R\bar{3}c$ space group without constraints

Zone	A (core) ¹	B	C	D	E (rim)	
SCOR	2.1962	2.3374	1.8919	1.8017	1.6553	
Overall scale f. ($\times 10^{-7}$)	52(3)*	80(4)	66(3)	170(2)	29(2)	
Eta (p-v) or m(p-vii)	0.32(7)	0.29(9)	0.09(6)	0.27(7)	0.12(8)	
Overall temp. factor	0.1(1)	0.3(1)	0.34(8)	0.7(1)	0.40(9)	
Halfwidth parameters	U	0.029(6)	0.06(1)	0.000(5)	-0.04(1)	0.013(5)
	V	-0.007(8)	-0.04(1)	0.024(7)	0.02(1)	-0.015(8)
	W	0.012(2)	0.026(4)	0.015(2)	0.014(4)	0.030(3)
Preferred orientation	0.72(5)	0.31(6)	0.43(4)	-1.0(1)	0.35(4)	
Asymmetry parameters	As 1	0.05(5)	0.18(4)	0.05(4)	0.08(5)	0.05(4)
	As 2	0.051(7)	0.067(7)	0.049(6)	0.044(6)	0.039(5)
X parameter	0.004(1)	0.003(2)	0.007(1)	0.003(1)	0.007(4)	
Zero-point	0.013(6)	0.032(7)	-0.003(6)	0.018(9)	-0.042(6)	
a_0 (Å)	16.852(1)	16.861(1)	16.8824(9)	16.873(2)	16.8615(9)	
c_0 (Å)	10.3252(9)	10.341(1)	10.3308(8)	10.338(1)	10.3386(7)	
V_0 (Å ³)	2539.4(3)	2547.8(5)	2549.9(3)	2548.7(5)	2545.6(2)	
N-P+C	6417	6417	6417	6417	6394	
R_p	14.4	14.1	12.0	13.3	12.8	
R_{wp}	19.3	19.2	16.3	18.5	17.5	
R_{exp}	11.48	11.41	11.63	14.08	14.04	
Conventional R_p	22.0	21.9	18.8	21.5	24.2	
Conventional R_{wp}	25.1	25.2	21.4	25.0	26.5	
Conventional R_{exp}	14.95	14.98	15.28	19.00	21.20	
Chi2	2.83	2.83	1.97	1.73	1.56	
Chi2 (Bragg contr.)	3.25	3.16	2.21	1.94	1.64	
DW-stat.	0.9079	0.8863	1.2388	1.3483	1.4249	
DW-exp.	1.9651	1.9651	1.9651	1.9651	1.9724	
GoF	1.6	1.7	1.4	1.3	1.2	
R_B	8.17	8.64	6.07	6.78	6.02	
R_F	6.80	7.42	6.31	6.94	6.24	

* – the numbers in parentheses are the estimated standard deviations multiplied with SCOR and refer to the last significant number; ¹ – Tančić et al. (2020)

Crystal structure studies were performed without any constraints in the $la\bar{3}d$, $R\bar{3}c$, $R\bar{3}$, $Fddd$, $C2/c$ and $\Gamma 1$ space groups (Table 1 and Appendix 1). Namely, as was shown by Tančić et al. (2020), two of these ($R\bar{3}c$ and $Fddd$) are of special interest, whereas other four space group refinements were made in this paper mostly for comparison, additional checking and other necessary data collecting. Also, among these space groups, five of them are very significant because of the possibility of the T_{2g} route, i.e. the phase transition from the $la\bar{3}d$ to the $R\bar{3}c$, $Fddd$, $C2/c$ and $\Gamma 1$ space groups (Hatch and Griffen, 1989), which is consistent with the chemical composition of the ferric grossular garnet studied (Tančić et al., 2012a, 2020). For the $R\bar{3}c$ and $Fddd$ space groups, structure studies with applied constraints were also performed (Appendix 2), which were appropriate to the observed chemical compositions of the zones (Tančić et al., 2012a), and calculated from the data given by Novak and Gibbs (1971). All angles were treated without constraints. Standard deviation values were multiplied with SCOR factors (Berar and Lelann, 1991).

Unit cell dimensions, calculated Ca and Al *sof*'s, and R_B and R_F reliability factors for the six space groups studied are listed in Appendix 1. At the first sight, it is obvious that the zones studied more or less differ from each other. Namely, unit cell dimen-

sions and volumes (V_0), which were in our previous studies were already shown as useful tools for approximate estimation of the Al-Fe³⁺ substitutions in the octahedral site (Tančić et al., 2012b; Cvetković and Tančić, 2019), mostly increase from zone A (core) to zones C and/or D, and then decrease in zones D and/or E (rim). For example, in the $la\bar{3}d$ space group there were calculated Al *sof*'s (Al_{calc} ; from 59 to 66 at.%) on the basis of the determined V_0 , and according to the starting data for Grs and Adr by Novak and Gibbs (1971). These are in concordance with their chemical compositions and with the Al-Fe³⁺ substitutions in the octahedral site. Namely, the Al³⁺ cation content decreases from zone A to D, with a difference of ~6 at. %, and it slightly increases in zone E (Tančić et al., 2012a). However, in most of the space groups studied, calculated Al (and Ca, as well) *sof*'s are more or less inconsistent with the ones expected (Appendix 2), and they are most well expressed in zones A and C. Differences between A and C by comparison with the other zones are also visible from variations in the cubic a_0 axis (Appendix 1) by the Raman shift of band VII detected (Tančić et al., 2012a), shown in Appendix 3.

In the triclinic $\Gamma 1$ space group, β_0 angles of 90.081(9)°, 90.111(1)° and 90.12(5)°, in the A, D and E zones, respectively; and γ_0 angles of 89.89(1)° and 89.922(4)°, in the B and C zones,

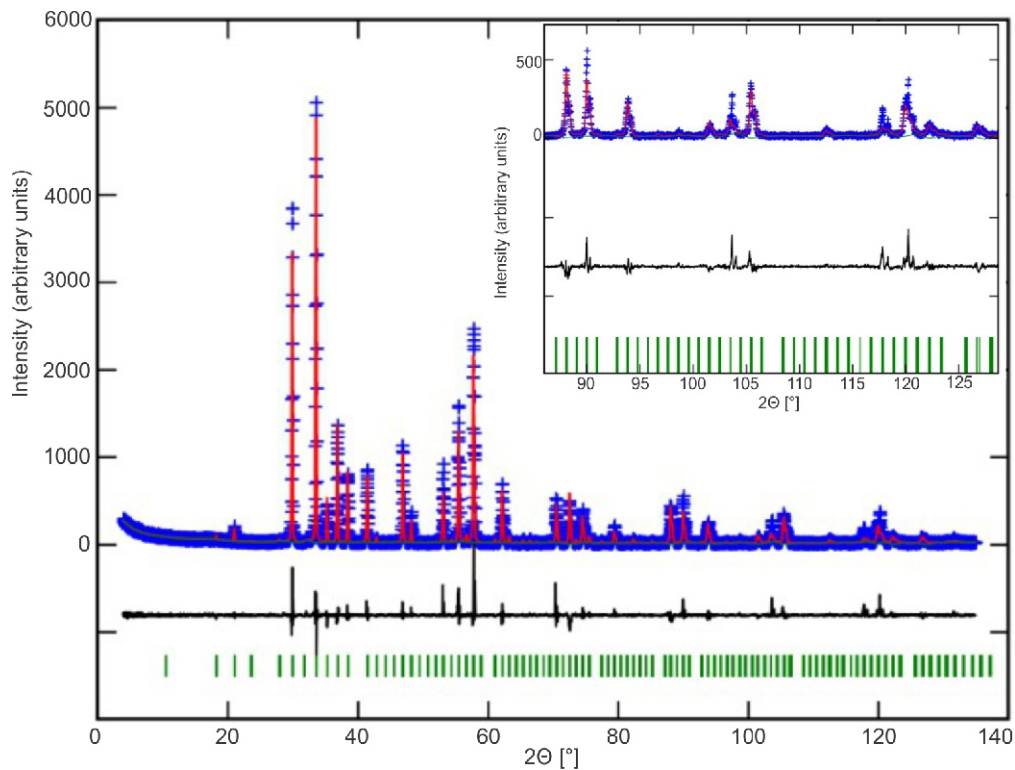


Fig. 3. Final Rietveld plot for the $R\bar{3}c$ space group of zone B

The area of $\sim 86\text{--}129^\circ$ 2θ angle range is magnified for a better view. The computer program plotCIF was used (<http://pubcif.iucr.org/services/tools/pdcifplot.php>). Observed intensities (Y_{obs}) are shown in blue, calculated intensities (Y_{calc}) are shown in red, and differences between observed and calculated intensities ($Y_{\text{obs}} - Y_{\text{calc}}$) were presented with black colour, whereas reflection (Bragg) positions were presented with green vertical bars

respectively, also clearly reveal violation of the cubic symmetry (Appendix 1). Therefore, differences between these angles and 90° are considerable (from $\sim 0.08^\circ$ in the A and C zones, to $\sim 0.12^\circ$ in the B, D and E zones), and increase in the following order: $C < A < B = D < E$. Average $\langle a_0 \rangle$ crystallographic unit cell parameters are mostly similar to the a_0 determined in the $Ia\bar{3}d$ space group. Also, among three crystallographic unit cell parameters in this space group, two of them are mainly quite similar, whereas the third is significantly different, strongly indicating the rhombohedral crystal system. The same observations may be likewise noticeable in the $Fddd$ and $C2/c$ space groups, after adequate transformation of their c_0 and a_0 crystallographic axes, respectively.

Furthermore, all of the rhombohedral distortion angles (α) calculated depart significantly by more than 0.004° from 60° , i.e. from 0.024° in the A and D zones, to 0.068° in the B zone in the $R\bar{3}c$ space group; and from 0.026° in the A and D zones, to 0.054° in the B zone in the $R\bar{3}$ space group (Appendix 1), which shows that the ferric grossular garnet studied is not cubic (Ishizawa and Inagaki, 2008).

Finally, because there are no significant differences between Al *sof*'s over the 4, 4 and 8 octahedral Y sites, which are the required conditions in the $R\bar{3}$, $C2/c$ and $I\bar{1}$ space groups, respectively (Appendix 1), it was shown that these possibilities were previously justifiably excluded (Tančić et al., 2020). Similarly, it is well known that if the single octahedral Y site in the $Ia\bar{3}d$ space group is split into two differently occupied sites Y1 and Y2, the symmetry reduces to the orthorhombic $Fddd$ or rhombohedral $R\bar{3}c$ space groups, which are of most significance in our case. However, here it can be seen that there are

two identical Al *sof*'s over the octahedral Y sites already in the zone B for the $Fddd$ space group, which could also reasonably indicate that ferric grossular garnet studied cannot crystallize in this space group. Accordingly, the results obtained for this space group will not be provided in detail, but only those which were necessary for required additional analysis and adequate data comparison with the $R\bar{3}c$ space group.

For the $R\bar{3}c$ space group, the final Rietveld plots are shown in Figures 3–6; while profile and structural parameters, unit cell dimensions, reliability factors, atom parameters, calculated *sof*'s of Ca and Al, and selected distances and angles, are shown in Table 1 and Appendices 4–7.

By comparison of the Rietveld plots (Figures 3–6), it could be seen that, generally, most of the intensities decrease from zones A–C to zones D–E, i.e. from core to rim. Another characteristic is that some peak intensities differ for zones A (Tančić et al., 2020) and C in comparison with zones B, D and E, especially in the area of the $\sim 87\text{--}107^\circ$ 2θ angle range (see also Figure 2). The magnified angle area in Figures 3–6 is further analyzed to the number of calculated Bragg's positions (Y_{calc}) and differences between starting and ending corresponding Y_{calc} 's (ΔY_{calc}), and shown in Appendix 8. Namely, Tančić et al. (2020) demonstrated that in zone A there is the simultaneous existence of diffraction peaks which include more or less clearly visible doublets and/or extra-broadening, whereas some others are without doublets. Although the $R\bar{3}c$ and $Fddd$ space groups have at least two calculated Bragg's positions for each diffraction peak in these selected areas, it is obvious that diffraction peaks without doublets have significantly smaller differences (ΔY_{calc}) between starting and ending corresponding Y_{calc} ,

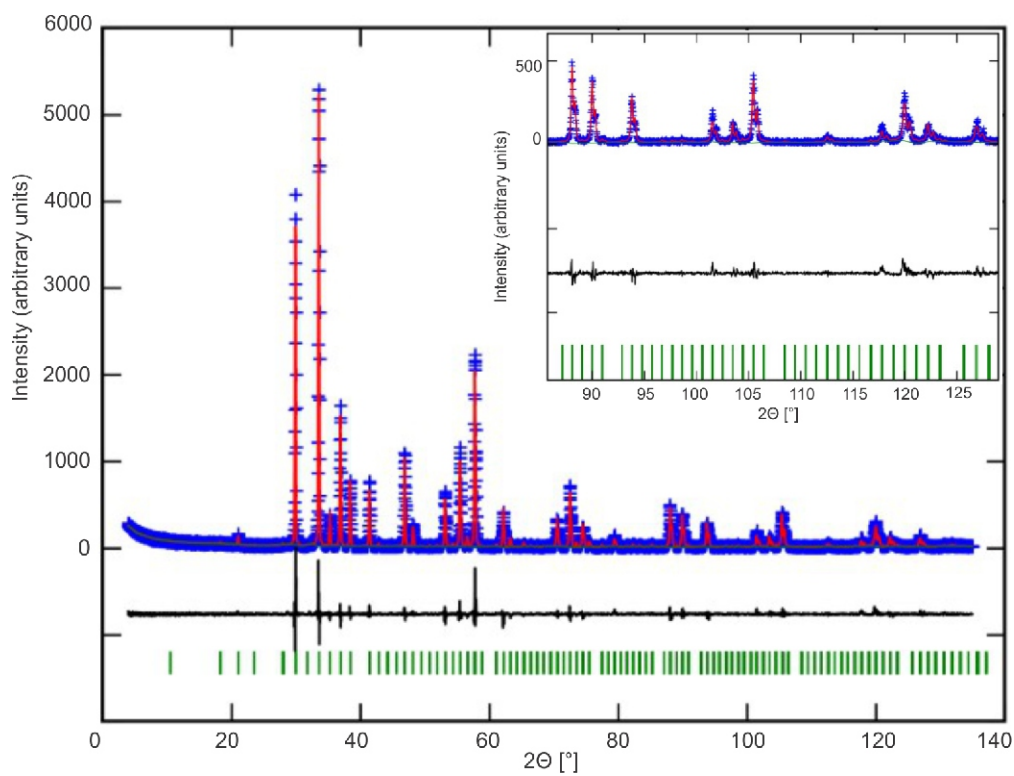


Fig. 4. Final Rietveld plot for the $R\bar{3}c$ space group of zone C

Symbols are the same as in [Figure 3](#)

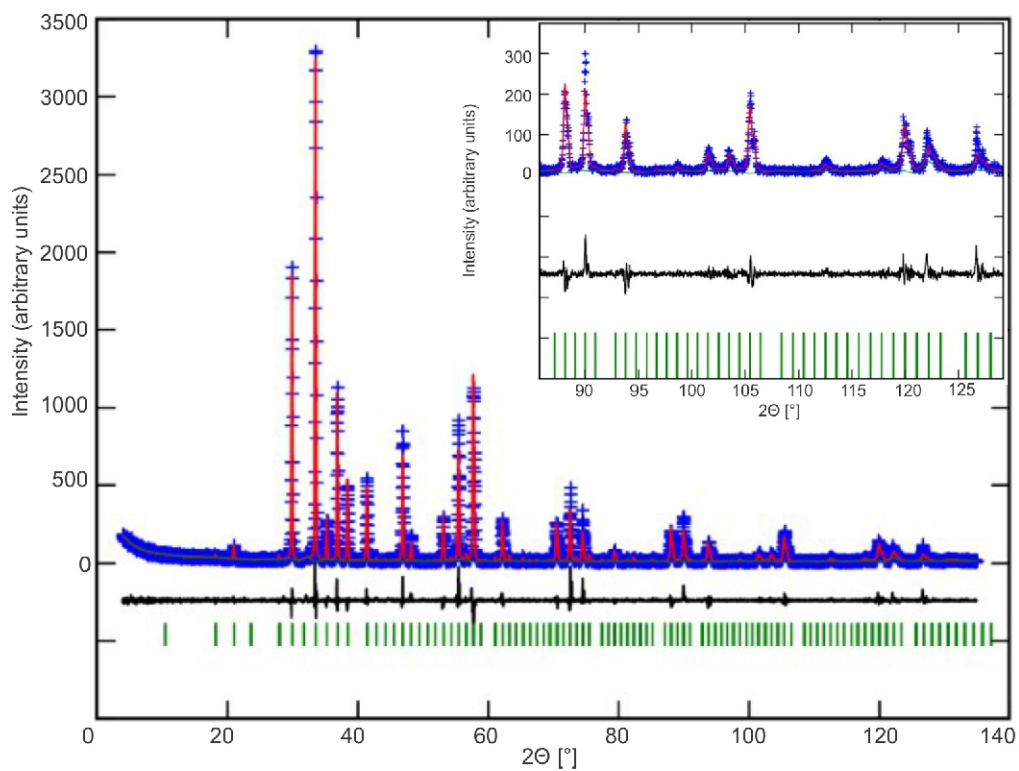


Fig. 5. Final Rietveld plot for the $R\bar{3}c$ space group of the zone D

Symbols are the same as in [Figure 3](#)

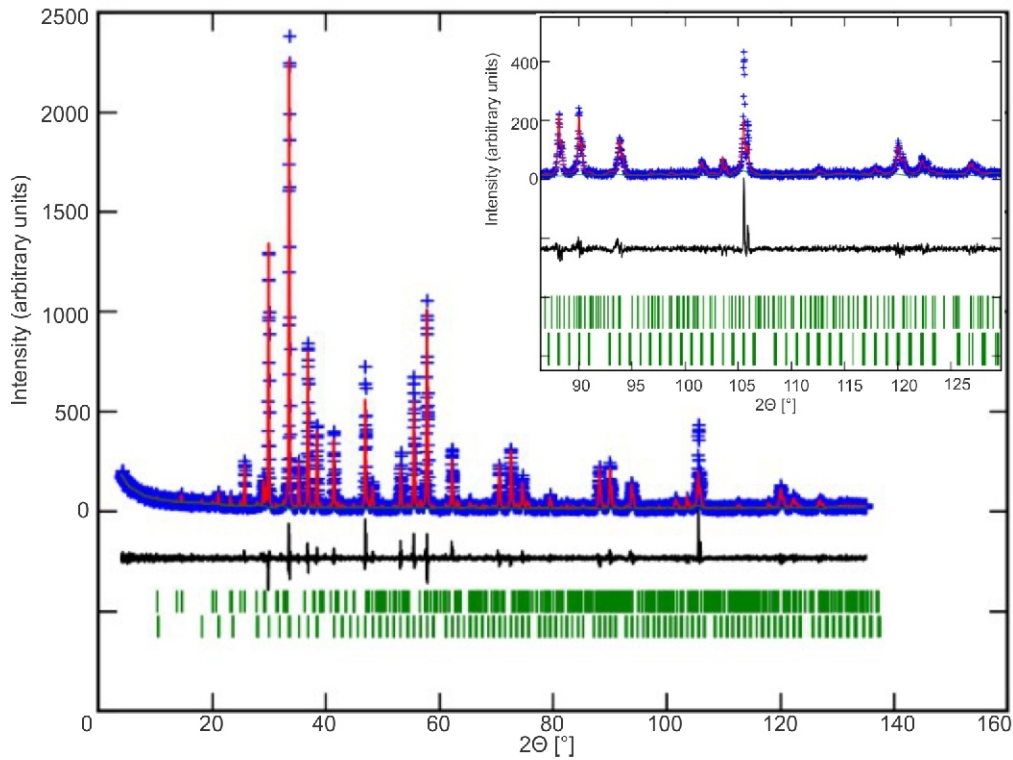


Fig. 6. Final Rietveld plot for the $R\bar{3}c$ space group of zone E

Symbols are the same as in Figure 3. First and second rows with green vertical bars represent minor scapolite and major ferric grossular phases, respectively

i.e. they are mostly within a narrower area than those with doublets in each of the zones studied. The clearest example of such a case is where one single diffraction peak (12 2 0) is in close vicinity to and between two pronouncedly diffraction doublets (884 & 12 2 2).

With refinement of the structure in zone E it was also established that the quantitative contents of ferric grossular is 92(5)%

and of scapolite 7.8(7)%, consistent with the previous microscopic estimation of ~5–6 % (Tančić et al., 2012a). The scapolite has the following unit cell dimensions: $a_0 = 12.130(2)$ Å, $c_0 = 7.568(1)$ Å, and $V_0 = 1113.5(3)$ Å³. The ratios of Ca : Na = 77(1) : 23(1) and Cl : C = 99(2) : 1(2) correspond to about 77% of the meionite composition. Structure refinement with applied constraints shows almost identical results: the quantitative con-

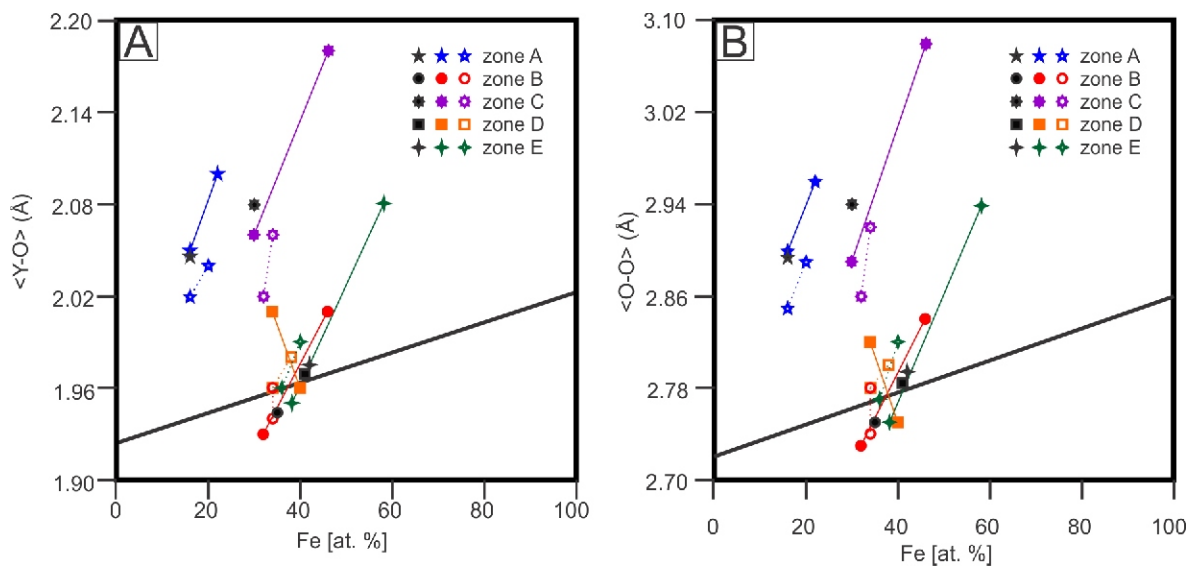


Fig. 7. Variations in Fe^{3+} and Al^{3+} over the octahedral sites in $Ia\bar{3}d$ (marked with black filled symbols), $R\bar{3}c$ (marked with coloured filled symbols and full lines) and $Fddd$ (marked with coloured empty symbols and dotted lines) space groups, by average: a) $\langle Y-O \rangle$; and b) $\langle O-O \rangle$ distances without constraints

Linear Grs-Adr joins (Novak and Gibbs, 1971) were also plotted for comparison

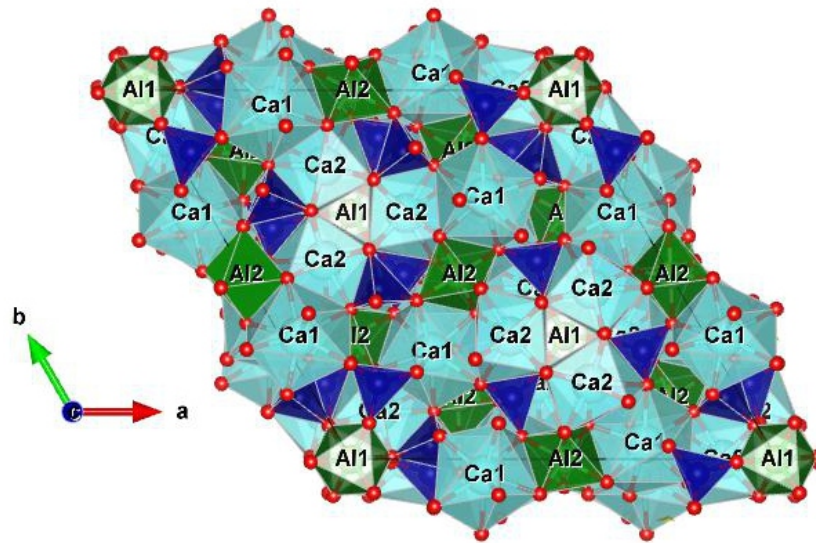


Fig. 8. Polyhedral crystal structure of the ferric grossular garnet studied in the rhombohedral $R\bar{3}c$ space group viewed down the c -axis

Symbols: Ca1 and Ca2 dodecahedra – light blue; Al1-Fe1 and Al2-Fe2 octahedra – green; Si–O tetrahedra – dark blue; oxygen anions – red

tent of ferric grossular is 92(4)% and of scapolite 7.5(6)%; unit cell dimensions are $a_0 = 12.130(2)$ Å, $c_0 = 7.568(1)$ Å, and $V_0 = 1113.7(3)$ Å³; and ratios are Ca : Na = 78(1) : 22(1) and Cl : C = 98(2) : 2(2). In order to additionally check these compositions, the equation given by Wehrenberg (1971) was used. Namely, taking into account the previously determined characteristic FT–IR band of 609 cm⁻¹ of scapolite (Tančić et al., 2012a), ~71 % of the meionite component was calculated, which correlates relatively well with the structure refinement.

The most important data for comparison between the $R\bar{3}c$ and $Fddd$ space groups is given in Appendix 9. Taking into account all of the studied zones in the $Fddd$ space group without constraints, it can be seen that the differences between Al *sof*'s (hereinafter Δ Al's) are insignificant, i.e. from 0 to 4 at.% (average 2.8 at.%), and with their estimated standard deviations (hereinafter *esd*'s) which exceed these values. On the other hand, Δ Al's over the octahedral Y sites in the $R\bar{3}c$ space group are much higher, i.e. from 6 to 20 at.% (average 12.4 at.%), and with mostly smaller *esd*'s. These Δ Al's are also proportional to the ΔY_{calc} values (Appendix 8). Differences of the average octahedral $\langle Y-O \rangle$ and distances are in very good agreement between each other and mostly proportional to the Fe³⁺ and Al³⁺ contents (Fig. 7) and Δ Al's. These distances are: for $\langle Y-O \rangle$ from 0.05 to 0.14 Å (average 0.09(2) Å) and from 0.02 to 0.04 Å (average 0.03(3) Å), and for from 0.06 to 0.19 Å (average 0.12(3) Å) and from 0.02 to 0.06 Å (average 0.04(4) Å) for the $R\bar{3}c$ and $Fddd$ space groups, respectively. Further, calculated degrees of ordering (m) and intersite distribution coefficients (K_{1-2}) are also concordant with each other and proportional to the other previously determined parameters. These parameters are: for m from 0.037 to 0.192 (average 0.104) and from 0 to 0.032 (average 0.020), and for K_{1-2} from 1.294 to 2.253 (average 1.765) and from 1 to 1.312 (average 1.156) for the $R\bar{3}c$ and $Fddd$ space groups, respectively. Variations in the Δ Al, m and K_{1-2} do not correlate well with the compositions of the zones studied for the $Fddd$ space group, in concordance with the observations by Shtukenberg et al. (2005). However, it is interesting that such variations are mostly proportional (except for zone D) for the $R\bar{3}c$ space group (Appendix 10). Finally, the R_B

and R_F factors are mostly smaller in $R\bar{3}c$ than for the $Fddd$ space group (Appendix 1). Therefore, all of the results given reasonably indicate that ferric grossular garnet studied crystallized in the rhombohedral $R\bar{3}c$ space group (Fig. 8), supporting our previous results (Tančić et al., 2020).

POLYHEDRAL DISTORTIONS AND STRUCTURAL DISORDERING BETWEEN THE ZONES STUDIED

Average determined selected distances without constraints (Appendix 6) were analyzed and compared with the expected ones, i.e. according to their chemical compositions (Appendix 2). It can be seen as follows: (1) $\langle \text{Si-O} \rangle$ distances are shorter in zones A and C, longer in zone B, and adequate in zones D and E; (2) $\langle Y-O \rangle$ distances are longer in zones A, C and E, and adequate in zones B and D; and (3) $\langle X-O \rangle$ distances are shorter only in zone B, and adequate in all of the other zones.

Also, as shown by Tančić et al. (2012b), the shift of the cation and oxygen sites could be of special interest. For example, oxygen O4 is very significant in our case, because it is common to the Si tetrahedron, Y1 octahedron and X2 dodecahedron. From Appendix 11 it can be seen that the Si–Y1 and Si–X2 distances determined mostly increase from zone A to zone D, and then decrease in zone E. This is in a very good agreement with the chemical compositions of the zones, because they follow the Al–Fe³⁺ substitutions at the octahedral site (Tančić et al., 2012a). On the other hand, Y1–X2 distances are in disagreement with previous observations, because they decrease as A>C>E>D>B. Also, from Appendix 6 it can be seen that the oxygen O4 site is most shifted towards Si and away from Y1 and X2 sites in zones A and C. Namely, $\langle \text{Si-O4} \rangle$ distances increase as C<A<E<B<D, whereas $\langle \text{Y1-O4} \rangle$ and $\langle \text{X2-O4} \rangle$ distances mostly decrease in the opposite direction, i.e. as C>A>E>B=D and C>A>E>D>B, respectively.

Therefore, the largest deviations were determined in the A and C zones, which also have the highest deviations from the Grs–Adr joins (Fig. 7). As argued by Tančić et al. (2020), such discrepancies occurred between the determined and expected

Table 2

Comparison of the Diamantina, Meka Presedla and Munam ferric grossular garnet specimens

	Space group	<Si–O>	<Y–O>	Al <i>sof</i>	<X–O>	Ca <i>sof</i>	<D–O>	< $r_{(x)}$ >/< $r_{(y)}$ >	
Diamantina	$la\bar{3}d$	1.642(2)	1.970(2)	66.9(8)	2.410(2) ^a	100	2.108(2) ^a	n. c.	
Meka Presedla	$R\bar{3}c$	A	1.637(6)	1.964(4)	64(2)	2.416(5)	98	2.108(5)	1.942
		B	1.658(6)	1.960(4)	61(6)	2.405(5)	98	2.107(5)	1.938
		C	1.631(6)	1.975(4)	60(3)	2.417(5)	98	2.110(5)	1.927
		D	1.644(6)	1.973(4)	58(2)	2.414(5)	98	2.111(5)	1.921
		E	1.643(6)	1.971(4)	58(6)	2.412(5)	98	2.110(5)	1.923
Munam	$\Gamma 1$	1.647(5) ^a	1.959(5) ^a	67	2.413(5) ^a	97	2.108(5) ^a	1.948	

Distances are given in Å; Al and Ca *sof*'s in at. %; <D–O> = {<Si–O> + <Y–O> + 2 × <X–O>}/4; ^a – Tančić et al. (2020); n. c. – not calculated

distances, and also Al and Ca *sof*'s, and should be treated as a consequence of a non-cubic space group of crystallization, polyhedral distortions and/or crystal disordering, because the <D–O> distances determined are within the expected ones in all of the zones studied. That being true, these distances become more adequate after constraints are applied and they return to the more “regular” positions (Appendices 2, 6 and 12). On the other hand, all of the average determined selected <O–Si–O>, <O–Y–O> and <O–X–O> angles (with and without constraints) are almost the same throughout the zones, and consistent with the expected ones (Appendix 7).

Because of the previously observed discrepancies, which deviate from the expected ones, especially in zones A and C, polyhedral volumes, quadratic elongations and angle variances in the tetrahedron and two octahedra (Robinson et al., 1971), were calculated and are given in Appendix 13. From the results, the discrepancies are generally more explicit without than those with constraints, as expected. Also, octahedron-1 is mostly more distorted than octahedron-2, except for zone B. Polyhedral volumes of the tetrahedron and two octahedra are in concordance with the average <Si–O>, <O–O>_{tet}, <Y–O> and <O–O>_{oct} distances throughout the zones (Appendix 6). Other results obtained also strongly indicate that distortions of the tetrahedron and octahedra indeed occurred to varying degrees, and influenced the deviations within the crystal structure. Namely, quadratic elongations and angle variances in the tetrahedron decrease in the following order among the zones: C>A>D>B>E; whereas, in octahedron-1, they decrease as C>A>D>E>B. Such orders are inversely proportional to the differences determined between β_0 (or γ_0) and 90° in the $\Gamma 1$ space group (Appendix 1). In this way, it is demonstrated that the A and C zones can be treated as disordered, whereas B, D and E may be treated as more or less ordered. Such variations between the zones could be further presented as two different hypothetical models, i.e. as (pseudo)cyclical or an oscillatory nature, or even as some kind of (pseudo)rotational or spiral (Appendix 14; see also Milke, 2004).

Notably, the *R*-values obtained are obviously not in concordance with the order-disorder variations determined, because their highest values are in the A, B and D, and lowest in the C and E zones (Table 1, Appendices 1 and 4). To explain this, further experiments are necessary, which is beyond the scope of this paper. Likewise, it is obvious that such order-disorder variations are not caused by the chemical compositions of the zones and Al³⁺–Fe³⁺ substitutions at the octahedral sites, but most probably to the formation conditions, which will be further analysed here.

As previously shown by Tančić et al. (2020), there are only two ferric grossular samples found in the available literature, which have very similar chemical compositions to our sample. These are Grs₆₀Adr₃₈–Grs₆₇Adr₃₃ from Diamantina, Brazil (cubic $la\bar{3}d$ space group, Resende and Fernandes, 2005), and Grs₆₇Adr₃₃ from Munam, North Korea (triclinic $\Gamma 1$ space group, Takéuchi et al., 1982). However, these three ferric grossular garnets are obviously structurally very different between each other, as all of them crystallize not only in different space groups, but also in different crystallographic systems. The deviations between the Diamantina, Meka Presedla and Munam specimens are also visible by comparison of their <Si–O>, <Y–O>, <X–O> and <D–O> distances, shown in Table 2 and Figure 9. From these results, the Diamantina and Meka Presedla specimens seem more disordered than that from Munam. This is not surprising, as symmetry increase, whereas ordering and birefringence decrease, with temperature increase (Allen and Busseck, 1988). The Meka Presedla and Munam specimens are from skarns, whereas for Diamantina the garnet's origin was not stated by Resende and Fernandes (2005); this limited our opportunities for some additional comparisons between them.

Finally, if we consider that heating of Grs up to 750°C causes the average <X–O> and <Al–O> distances to increase (Meagher, 1975), and implement this possibility to the determined average <X₂–O₄₁–O_{4\beta_0 (or γ_0) and 90°. Such temperature variations between the zones may also be inferred as cyclical or oscillatory (see Appendix 14A).}

However, if the zones C and A were formed at higher temperatures, beside of the polyhedral distortions and order-disorder variations already established, the hypothetical possibility that some part of the thermally generated vacancies replaced Si in the tetrahedron cannot be excluded. Namely, the lack of boron (Tančić et al., 2012a), and shorter <Si–O> and <O–O>_{tet} distances, and smaller polyhedral volumes in the tetrahedron (even with applied constraints), could reasonably indicate such a possibility. Similarly, some part of Al³⁺ or Fe³⁺ could replace Si in the tetrahedron at zone B, because of the longer <Si–O> and <O–O>_{tet} distances, and larger polyhedral volumes in the tetrahedron (Table 2 and Appendices 2, 6 and 13), which were not previously established with the EMPA measurements and calculations applied (Tančić et al., 2012a).

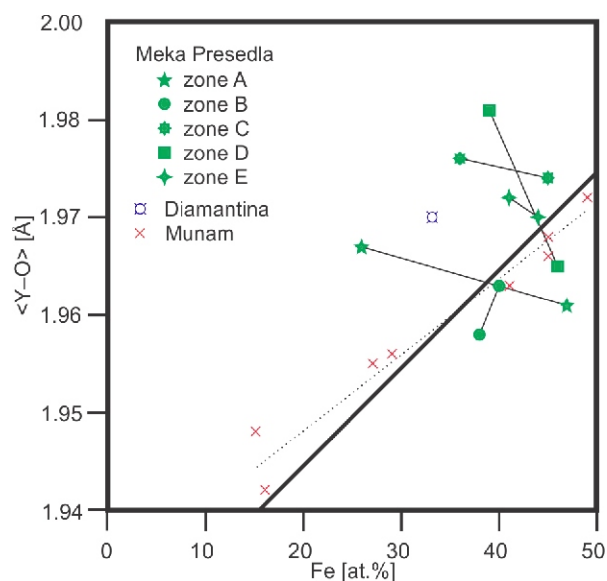


Fig. 9. Comparison of Fe^{3+} and Al^{3+} variations over the octahedral sites by the octahedral average $\langle Y-O \rangle$ distances for the Meka Presedla rhombohedral $R\bar{3}c$ space group (green), Diamantina cubic $Ia\bar{3}d$ space group (blue), and Munam triclinic $\Gamma 1$ space group (red) ferric grossular garnets

The linear Grs-Adr join (Novak and Gibbs, 1971) was also plotted for comparison

CONCLUSIONS

Although Rietveld crystal structure refinement has some disadvantages in comparison with single-crystal X-ray diffraction (SCXRD) and other methods, the results described here and by Tančić et al. (2020) indicate that there is sufficient reliable evidence showing that the ferric grossular garnet crystallized in the rhombohedral $R\bar{3}c$ space group. However, one could reasonably ask if this is possible. To the best of our knowledge, there is no published physical or other reliable evidence for grossular-andradite garnet that crystallizes in the rhombohedral crystal structure. Furthermore, there is no such evidence for any of the natural garnet subgroup member (Grew et al., 2013).

On the other hand, much evidence supports a rhombohedral crystal structure for this ferric grossular garnet, such as:

1. nikmelnikovite ($\text{Ca}_{12}\text{Fe}^{2+}\text{Fe}^{3+}_3\text{Al}_3(\text{SiO}_4)_6(\text{OH})_{20}$), is the first IMA-approved member of the garnet supergroup which crystallizes in the rhombohedral $R\bar{3}$ space group (Krivovichev et al., 2021);
2. there are some already known synthetic garnets which crystallize in the rhombohedral $R\bar{3}c$ and $R\bar{3}$ space groups, such as $\text{Ho}_3\text{Fe}_5\text{O}_{12}$ (Guillot et al., 1984), $\text{Y}_3\text{Fe}_5\text{O}_{12}$ (Rodić et al., 1999) and $\text{R}_3\text{Fe}_5\text{O}_{12}$ with $R = \text{Y}$, Eu and Dy (Vandormael et al., 2001);
3. $R\bar{3}c$ is a subgroup of the $Ia\bar{3}d$ space group;
4. $R\bar{3}c$ is one of the possibilities of the T_{2g} route, i.e. a phase transition from $Ia\bar{3}d$ to the $R\bar{3}c$, $Fddd$, $C2/c$ and $\Gamma 1$ space groups (Hatch and Griffen, 1989), which

is consistent to the chemical composition of the ferric grossular garnet studied (Tančić et al., 2012a, 2020);

5. only the rhombohedral $R\bar{3}c$ and $R\bar{3}$ space groups satisfied both possible geometrical-mathematical conditions, i.e. following transformations of the cubic symmetry axis: (a) $a_{\text{orthomb}} = a_{\text{0cubic}} \times \sqrt{3} / 2 = \sim 10.32 \text{ \AA}$ and (b) $c_{\text{orthomb}} = a_{\text{0cubic}} \times \sqrt{2} = \sim 16.86 \text{ \AA}$ (Tančić et al., 2020);
6. among three determined or calculated crystallographic axes, two are quite similar, whereas the third is significantly different, indicating the rhombohedral crystal system (Appendix 1);
7. observed departures of the calculated rhombohedral distortion angles α from 60° (Appendix 1);
8. the degree of ordering (m) and intersite distribution coefficient (K_{1-2}) determined for the $Fddd$ space group (Appendix 9) are the lowest among all those determined for the $Fddd$ and $\Gamma 1$ space groups (Shtukenberg et al., 2005);
9. differences in sof 's decrease with an increase in temperature (Shtukenberg et al., 2005). Accordingly, the crystallization temperature of the ferric grossular garnet studied should be higher than required for the $Fddd$ space group, because of their insignificant ΔAl 's;
10. it is well known that symmetry increases with increasing temperature, and the preferred crystal structure symmetry is governed by the crystallization temperature;
11. Milke (2004) suggests that an ordering process which relates to the birefringence in intermediate and Grs-rich grossular-andradites seems to be largely controlled by the growth temperature, whereas chemical composition, growth medium, and pressure are only minor factors. He obtained a temperature of 730°C between birefringent and isotropic $\text{Gr}_{76-99}\text{Adr}_{1-24}$ samples. Also, Hariya and Kimura (1978) found a transition from anisotropic to isotropic $\text{Gr}_{50}\text{Adr}_{50}$ grossular-andradite at temperatures slightly below 700°C , with a tendency to lower temperatures at higher pressure. According to Shtukenberg et al. (2005), the transformation from the orthorhombic to the triclinic crystal structure occurs in the range $150\text{--}350^\circ\text{C}$. Crystallization at elevated temperatures leads to a more symmetric orthorhombic crystal structure. Therefore, it can be concluded that a rhombohedral crystal structure could be possible in the temperature range somewhere between the orthorhombic and cubic systems. Previously obtained formation temperatures between ~ 600 and 720°C (Tančić et al., 2012a) of the $\text{Gr}_{58-64}\text{Adr}_{36-41}\text{Sp}_2$ ferric grossular garnet studied in this paper, just below those required for cubic symmetry, could sufficiently satisfy such a condition.

Besides the rhombohedral symmetry determined, which could be treated as a primary cause for its slight optical anisotropy, residual strain may be also treated as the second possible cause, due to the chemical, structural, order-disorder and formation differences recognized between the zones.

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APPENDIX 1

Unit cell dimensions, calculated Ca and Al *sof*'s (in at.%), and R_B and R_F reliability factors (in %) for the studied six space groups without constraints

Space g.		A (core) ¹	B	C	D	E (rim)
$Ia\bar{3}d$	SCOR	2.3375	2.4607	2.0808	2.0659	1.7162
	a_0 (Å)	11.9188(6)	11.9320(8)	11.9342(5)	11.9342(8)	11.9287(6)
	V_0 (Å ³)	1693.2(1)	1698.8(2)	1699.7(1)	1699.7(2)	1697.4(1)
	$a_0 \times \sqrt{3}/2$ (Å)	10.3220	10.3334	10.3353	10.3353	10.3306
	$a_0 \times \sqrt{2}$ (Å)	16.8557	16.8744	16.8775	16.8775	16.8697
	Ca	78(5)	96(1)	79(1)	98(4)	97(4)
	Al	84(1)	65(1)	70(1)	59(1)	58(1)
	Al _{calc}	66	60	59	59	61
	R_B	9.23	8.87	6.57	7.98	6.82
	R_F	7.69	7.20	6.51	6.89	5.83
$R\bar{3}c$	SCOR	2.1962	2.3374	1.8919	1.8017	1.6553
	a_0 (Å)	16.852(1)	16.861(1)	16.8824(9)	16.873(2)	16.8615(9)
	c_0 (Å)	10.3252(9)	10.341(1)	10.3308(8)	10.338(1)	10.3386(7)
	V_0 (Å ³)	2539.4(3)	2547.8(5)	2549.9(3)	2548.7(5)	2545.6(2)
	$4 \times c_0/a_0$	2.45080	2.45324	2.44771	2.45078	2.45259
	α (°)	59.976	59.932	60.032	59.976	59.944
	Ca1	100(4)	94(6)	100(4)	102(2)	98(4)
	Ca2	80(20)	96(4)	78(12)	96(2)	96(14)
	<Ca>	90(12)	95(5)	89(8)	99(2)	97(9)
	Al1	78(12)	54(18)	54(12)	66(6)	42(12)
	Al2	84(4)	68(6)	70(4)	60(2)	62(4)
	<Al>	81(8)	61(12)	62(8)	63(4)	52(8)
	R_B	8.17	8.64	6.07	6.78	6.02
	R_F	6.80	7.42	6.31	6.94	6.24
$R\bar{3}$	SCOR	2.1321	2.2775	1.8849	1.7886	1.6468
	a_0 (Å)	16.852(1)	16.867(1)	16.8713(8)	16.872(2)	16.874(1)
	c_0 (Å)	10.3255(8)	10.341(1)	10.3416(7)	10.338(1)	10.326(1)
	V_0 (Å ³)	2539.4(3)	2547.7(4)	2549.2(2)	2548.7(5)	2546.4(2)
	$4 \times c_0/a_0$	2.45087	2.45236	2.45188	2.45092	2.44779
	α (°)	59.974	59.946	59.956	59.974	60.030
	Ca1	110(20)	94(7)	110(20)	100(20)	110(20)
	Ca2	70(20)	96(5)	80(20)	96(3)	100(20)
	<Ca>	90(20)	95(6)	95(20)	98(12)	105(20)
	Al1	78(36)	60(36)	54(24)	60(42)	42(36)
	Al2	78(36)	48(36)	66(36)	66(36)	60(36)
	Al3	80(20)	70(14)	66(16)	58(18)	58(18)
	Al4	80(20)	70(14)	72(14)	66(18)	72(16)
	<Al>	79(28)	62(25)	64(22)	62(28)	58(26)
	R_B	7.69	8.48	6.23	6.75	6.12
R_F	6.58	7.69	6.64	7.11	5.77	
$Fddd$	SCOR	2.2850	2.3749	1.8482	1.8893	1.6218
	a_0 (Å)	16.874(1)	16.895(2)	16.895(1)	16.861(2)	16.892(1)
	b_0 (Å)	16.849(1)	16.863(2)	16.873(1)	16.8919(9)	16.860(1)
	c_0 (Å)	11.9137(7)	11.927(1)	11.9292(7)	11.922(1)	11.9253(7)
	V_0 (Å ³)	3387.2(4)	3398.1(7)	3400.7(4)	3395.6(5)	3396.4(4)
	$c_0 \times \sqrt{2}$ (Å)	16.8485	16.867	16.8704	16.860	16.8649
	Ca1	88(8)	96(8)	92(8)	92(8)	112(8)
	Ca2	104(12)	88(8)	84(8)	104(4)	108(8)
	Ca3	109(3)	98(3)	113(2)	98(2)	95(2)
	<Ca>	100(8)	94(6)	96(6)	98(5)	105(6)
	Al1	80(8)	66(10)	66(8)	66(6)	64(8)
	Al2	84(8)	66(10)	68(8)	62(4)	60(8)
	<Al>	82(8)	66(10)	67(8)	64(5)	62(8)
	R_B	8.63	8.72	5.80	7.67	6.30
R_F	7.39	7.65	6.06	7.54	6.22	
$C2/c$	SCOR	2.1433	2.0216	1.7629	1.7536	1.6051
	a_0 (Å)	16.861(2)	16.894(4)	16.8923(5)	16.900(2)	16.888(1)
	b_0 (Å)	11.9205(6)	11.951(2)	11.9329(4)	11.924(3)	11.922(2)
	c_0 (Å)	11.910(1)	11.943(3)	11.9432(4)	11.945(3)	11.943(2)
	β_0 (°)	134.97(1)	135.08(1)	134.079(1)	135.084(5)	135.071(4)
	V_0 (Å ³)	1693.5(3)	1702.9(7)	1699.9(1)	1700.1(6)	1698.2(4)
	$a_0 / \sqrt{2}$ (Å)	11.922	11.946	11.945	11.950	11.942
	Ca1	106(14)	96(10)	108(10)	92(10)	86(14)

	Ca2	98(10)	94(8)	112(12)	92(10)	92(16)
	Ca3	97(6)	102(7)	110(9)	104(7)	93(9)
	Ca4	108(7)	85(5)	86(3)	101(3)	100(3)
	<Ca>	102(9)	94(8)	106(8)	97(8)	93(10)
	Al1	100(20)	58(14)	60(20)	60(20)	60(20)
	Al2	98(18)	56(16)	60(20)	72(18)	80(20)
	Al3	120(20)	80(14)	60(20)	60(20)	60(20)
	Al4	100(20)	86(18)	60(20)	60(20)	60(20)
	<Al>	104(20)	70(16)	60(20)	63(20)	65(20)
	R_B	7.54	7.30	6.73	6.25	5.33
	R_F	6.48	6.95	8.38	7.03	5.53
	SCOR	2.7890	3.0879	2.2768	2.3167	1.7755
	a_0 (Å)	11.927(2)	11.926(3)	11.9467(8)	11.941(2)	11.931(1)
	b_0 (Å)	11.910(2)	11.945(2)	11.931(1)	11.925(2)	11.9260(9)
	c_0 (Å)	11.920(1)	11.929(2)	11.9303(8)	11.928(2)	11.940(1)
	< a_0 (Å)>	11.919(2)	11.933(2)	11.9360(9)	11.931(2)	11.932(1)
	α_0 (°)	90.01(2)	89.96(1)	90.060(4)	89.98(1)	89.96(7)
	β_0 (°)	90.081(9)	90.04(1)	90.007(5)	90.11(1)	90.12(5)
	γ_0 (°)	89.98(2)	89.89(1)	89.922(4)	89.94(1)	90.03(6)
	V_0 (Å ³)	1693.4(4)	1699.3(5)	1700.5(2)	1698.5(4)	1698.9(3)
	$\beta_0 - 90$ (°)	0.081(9)	-0.11(1) [#]	-0.078(4) [#]	0.11(1)	0.12(5)
	Ca1	120(50)	100(50)	110(20)	110(20)	100(20)
	Ca2	100(30)	100(30)	110(30)	80(20)	100(20)
	Ca3	90(20)	90(30)	90(20)	90(20)	80(10)
	Ca4	110(50)	100(40)	110(20)	100(20)	90(20)
	Ca5	110(20)	100(40)	110(30)	100(20)	100(20)
	Ca6	100(30)	90(30)	90(20)	100(20)	90(20)
	<Ca>	105(33)	97(37)	103(23)	97(20)	93(18)
	Al1	80(100)	80(100)	80(100)	60(60)	40(60)
	Al2	80(80)	60(80)	80(80)	80(80)	60(80)
	Al3	80(100)	60(80)	80(80)	80(80)	80(60)
	Al4	80(80)	60(100)	80(80)	80(80)	60(60)
	Al5	80(80)	80(80)	80(60)	80(60)	80(60)
	Al6	80(120)	60(80)	80(60)	80(60)	60(80)
	Al7	80(100)	60(120)	60(60)	40(80)	60(60)
	Al8	80(80)	60(120)	80(60)	60(60)	80(80)
	<Al>	80(92)	65(95)	78(72)	70(70)	65(68)
	R_B	11.2	12.0	7.94	9.01	7.65
	R_F	9.71	11.6	10.3	9.35	8.21
$\bar{1}$						

Calculated $4 \times c_0/a_0$ parameters and distortion α angles for the rhombohedral $R\bar{3}c$ and $R\bar{3}$ space groups, and specific geometry-mathematical transformations of the crystallographic axes within $la\bar{3}d$, $Fddd$ and $C2/c$ space groups, are also presented; * – the numbers in parentheses are the estimated standard deviations multiplied with SCOR and refer to the last significant number; ¹ – Tančić et al. (2020); [#] – $\gamma_0 - 90$ [°]; for the triclinic $\bar{1}$ space group, instead of 100 parameters for the background description, it was used 6 Chebyshev's polynomial parameters, and there were omitted 2 θ region from 4° to 10°

APPENDIX 2

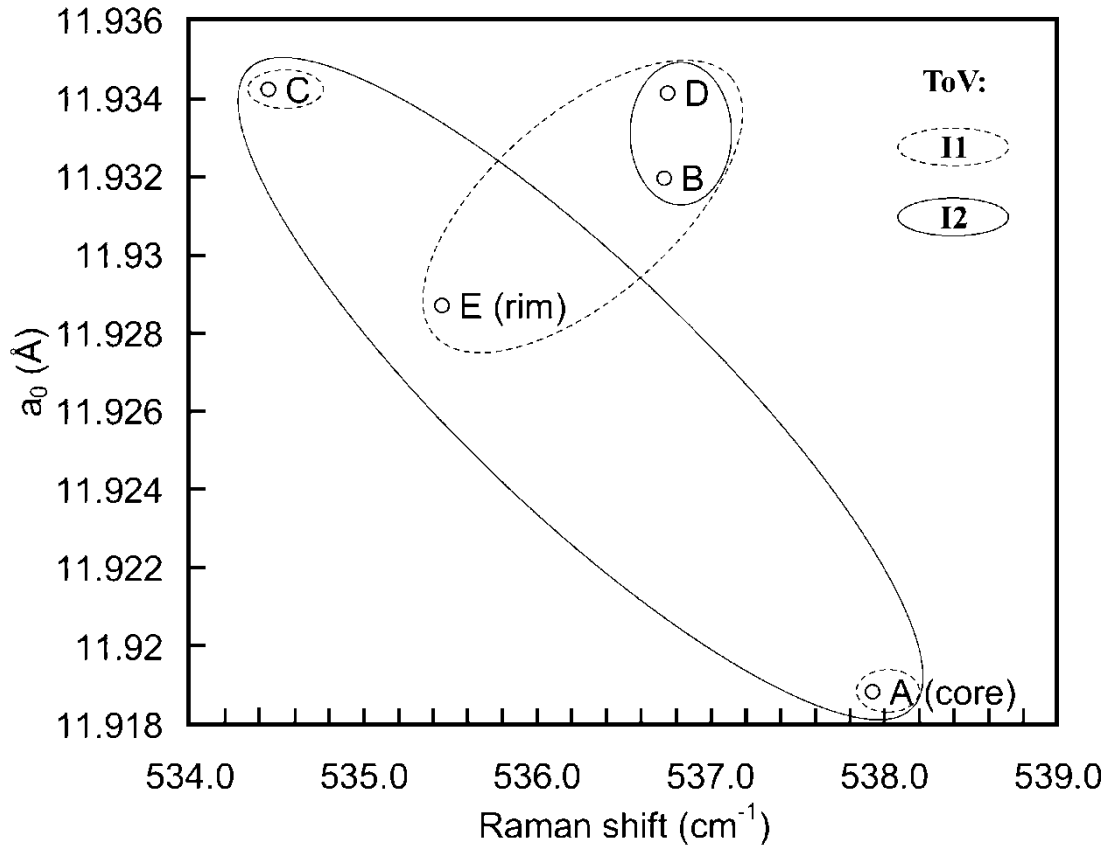
Applied constraints for the Ca and Al *sof's* (in at.%)
and for the <Si-O>, <Y-O>, <X'-O> and <X''-O> distances (in Å)

	A (core) ¹	B	C	D	E (rim)
Ca	98	98	98	98	98
Al	64	62	59	58	58
<Si-O>	1.644(5)	1.644(5)	1.644(5)	1.644(5)	1.644(5)
<Y-O>	1.960(5)	1.962(5)	1.965(5)	1.966(5)	1.966(5)
<X'-O>	2.336(5)	2.336(5)	2.337(5)	2.338(5)	2.338(5)
<X''-O>	2.494(5)	2.494(5)	2.494(5)	2.494(5)	2.494(5)
<D-O>	2.108(5)	2.109(5)	2.110(5)	2.110(5)	2.110(5)

¹ – Tančić et al. (2020); <D-O> = {<Si-O> + <Y-O> + <X'-O> + <X''-O>}/4

APPENDIX 3

Variations of the cubic a_0 axis ($Ia\bar{3}d$ space group) by Raman shift of band



VIITwo types of variations (ToV) are clearly visible: (I) proportional for zones B, D and E, and declination from it at zones A and C, and (II) inversely proportional for zones A, C and E, and declination from it at zones B and D

APPENDIX 4

Profile and structural parameters, unit cell dimensions and reliability factors for the $R\bar{3}c$ space group with constraints

Zone	A (core) ¹	B	C	D	E (rim)	
SCOR	2.4445	2.5216	2.2050	1.9755	1.7076	
Overall scale f. ($\times 10^{-4}$)	60(3)	75(4)	70(3)	130(20)	41(1)	
Eta (p-v) or m(p-vii)	0.32(9)	0.32(9)	0.08(7)	0.28(8)	0.13(8)	
Overall temp. factor	0.3(1)	0.4(1)	0.50(8)	0.8(1)	0.51(8)	
Halfwidth parameters	U	0.020(7)	0.06(1)	-0.009(5)	0.03(1)	0.012(5)
	V	0.004(9)	-0.03(1)	0.031(8)	0.03(1)	-0.018(8)
	W	0.010(2)	0.026(4)	0.014(2)	0.013(4)	0.031(3)
Preferred orientation	0.60(5)	0.36(5)	0.43(4)	-0.68(7)	0.25(3)	
Asymmetry parameters	As 1	0.03(6)	0.18(4)	0.07(5)	0.08(6)	0.04(4)
	As 2	0.048(8)	0.066(7)	0.053(7)	0.044(7)	0.038(6)
X parameter	0.004(2)	0.002(2)	0.009(1)	0.003(2)	0.008(2)	
Zero-point	0.011(8)	0.032(8)	-0.001(7)	0.02(1)	-0.043(6)	
a_0 (Å)	16.852(2)	16.867(2)	16.883(1)	16.873(2)	16.8611(9)	
c_0 (Å)	10.326(1)	10.341(2)	10.3304(8)	10.338(2)	10.3399(7)	
V_0 (Å ³)	2539.6(4)	2547.7(5)	2549.9(3)	2548.9(6)	2545.8(3)	
N-P+C	6420	6420	6420	6420	6397	
R_p	15.2	14.5	12.8	14.0	13.1	
R_{wp}	20.4	19.6	17.3	19.0	17.7	
R_{exp}	11.48	11.42	11.63	14.09	14.04	
Conventional R_p	23.7	23.0	20.2	22.8	25.1	
Conventional R_{wp}	26.9	26.0	22.9	25.7	26.9	
Conventional R_{exp}	15.14	15.13	15.38	19.12	21.35	
Chi2	3.17	2.95	2.22	1.81	1.59	
Chi2 (Bragg contr.)	3.65	3.30	2.49	2.03	1.67	
DW-stat.	0.8168	0.8498	1.1015	1.2918	1.3967	
DW-exp.	1.9642	1.9642	1.9642	1.9642	1.9715	
GoF	1.8	1.7	1.5	1.3	1.2	
R_B	10.1	9.84	7.46	8.07	6.92	
R_F	8.17	8.15	6.38	7.41	6.94	

* – the numbers in parentheses are the estimated standard deviations multiplied with SCOR and refer to the last significant number; ¹ – Tančić et al. (2020)

APPENDIX 5

Atom parameters (in Å) and calculated Ca and Al *sof*'s (at.%) for the $R\bar{3}c$ space group [without (a) and with (b) constraints]

Zone	A (core) ¹		B		C		D		E (rim)		
	a	b	a	b	a	b	a	b	a	b	
Ca1	x	0.625(1)	0.6239(6)	0.625(2)	0.6248(6)	0.628(1)	0.6258(5)	0.623(1)	0.6237(4)	0.620(1)	0.6239(4)
	y	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
	z	0.250	0.250	0.250	0.250	0.250	0.250	0.250	0.250	0.250	0.250
Ca2	x	0.129(1)	0.1274(6)	0.122(2)	0.1228(6)	0.127(1)	0.1236(6)	0.124(2)	0.1213(5)	0.127(2)	0.1235(5)
	y	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
	z	0.250	0.250	0.250	0.250	0.250	0.250	0.250	0.250	0.250	0.250
Mn2	x	0.129(1)	n. c.	n. c.	n. c.	0.127(1)	n. c.	n. c.	n. c.	0.127(2)	n. c.
	y	0.000	n. c.	n. c.	n. c.	0.000	n. c.	n. c.	n. c.	0.000	n. c.
	z	0.250	n. c.	n. c.	n. c.	0.250	n. c.	n. c.	n. c.	0.250	n. c.
Al1	x	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
	y	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
	z	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
Fe1	x	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
	y	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
	z	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
Al2	x	0.500	0.500	0.500	0.500	0.500	0.500	0.500	0.500	0.500	0.500
	y	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
	z	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
Fe2	x	0.500	0.500	0.500	0.500	0.500	0.500	0.500	0.500	0.500	0.500
	y	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
	z	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000
Si	x	0.839(1)	0.8338(7)	0.837(2)	0.8328(7)	0.834(2)	0.8334(6)	0.829(2)	0.8316(6)	0.833(2)	0.8325(5)
	y	0.041(2)	0.0395(7)	0.046(2)	0.0427(7)	0.041(1)	0.0427(6)	0.045(2)	0.0437(5)	0.043(2)	0.0418(5)
	z	0.417(2)	0.4174(9)	0.416(3)	0.418(1)	0.418(2)	0.4183(8)	0.417(2)	0.4185(8)	0.416(2)	0.4180(7)
O1	x	0.204(3)	0.2051(8)	0.210(5)	0.2096(8)	0.205(2)	0.2087(7)	0.215(3)	0.2091(6)	0.211(4)	0.2077(5)
	y	0.412(3)	0.4069(7)	0.405(5)	0.4087(7)	0.414(2)	0.4088(6)	0.417(3)	0.4092(6)	0.410(4)	0.4078(5)
	z	0.495(3)	0.493(1)	0.484(5)	0.490(1)	0.491(3)	0.4920(9)	0.490(2)	0.4898(8)	0.495(3)	0.4923(7)
O2	x	0.721(3)	0.7278(6)	0.729(4)	0.7268(7)	0.724(3)	0.7291(6)	0.721(3)	0.7274(5)	0.719(3)	0.7276(5)
	y	0.967(3)	0.9642(7)	0.965(4)	0.9633(7)	0.965(2)	0.9643(6)	0.960(3)	0.9643(5)	0.958(3)	0.9646(5)
	z	0.382(3)	0.379(1)	0.391(4)	0.380(1)	0.386(3)	0.3815(9)	0.370(2)	0.3773(8)	0.372(4)	0.3771(7)
O3	x	0.818(4)	0.8230(8)	0.821(5)	0.8226(8)	0.818(3)	0.8227(7)	0.826(4)	0.8236(6)	0.825(5)	0.8239(5)
	y	0.052(3)	0.0624(7)	0.061(5)	0.0610(7)	0.049(2)	0.0611(7)	0.067(4)	0.0610(6)	0.065(4)	0.0612(5)
	z	0.561(3)	0.5678(8)	0.580(5)	0.5730(9)	0.570(4)	0.5700(7)	0.565(2)	0.5726(7)	0.566(4)	0.5716(6)
O4	x	0.896(3)	0.9087(7)	0.905(4)	0.9077(7)	0.894(2)	0.9071(6)	0.907(3)	0.9056(6)	0.899(3)	0.9064(5)
	y	0.893(4)	0.9015(8)	0.898(5)	0.8998(6)	0.882(2)	0.8985(5)	0.895(3)	0.8977(5)	0.895(3)	0.8996(4)
	z	0.892(3)	0.8906(9)	0.891(5)	0.894(1)	0.895(3)	0.8945(9)	0.894(2)	0.8959(8)	0.890(4)	0.8938(7)
Ca1	100(4)	98	94(6)	98	100(4)	98	102(2)	98	98(4)	98	
Ca2	80(20)	98	96(4)	98	78(12)	98	96(2)	98	96(14)	98	
<Ca>	90(12)	98	95(5)	98	89(8)	98	99(2)	98	97(9)	98	
Al1	78(12)	53(1)	54(18)	60(6)	54(12)	55(2)	66(6)	61(2)	42(12)	59(5)	
Al2	84(4)	74(2)	68(6)	62(6)	70(4)	64(4)	60(2)	54(2)	62(4)	56(6)	
<Al>	81(8)	64(2)	61(12)	61(6)	62(8)	60(3)	63(4)	58(2)	52(8)	58(6)	

* – the numbers in parentheses are the estimated standard deviations multiplied with SCOR and refer to the last significant number;

¹ – Tančić et al. (2020); n. c. – not calculated

APPENDIX 6

Selected distances (in Å) for the $R\bar{3}c$ space group [without (a) and with (b) constraints]

Zone		A (core) ¹		B		C		D		E (rim)	
		a	b	a	b	a	b	a	b	a	b
Si–O(1)	1 ^s	1.55(3)	1.645(6)	1.69(3)	1.659(6)	1.56(2)	1.636(6)	1.59(2)	1.638(6)	1.62(3)	1.643(6)
Si–O(2)	1	1.77(2)	1.641(5)	1.67(2)	1.657(5)	1.69(2)	1.633(5)	1.74(3)	1.647(5)	1.79(3)	1.643(5)
Si–O(3)	1	1.56(2)	1.632(6)	1.76(3)	1.658(6)	1.61(2)	1.625(5)	1.57(2)	1.637(5)	1.62(3)	1.643(6)
Si–O(4)	1	1.43(4)	1.629(8)	1.60(4)	1.657(8)	1.35(3)	1.630(8)	1.68(4)	1.653(8)	1.57(5)	1.643(8)
<Si–O>		1.58(3)	1.637(6)	1.68(3)	1.658(6)	1.55(2)	1.631(6)	1.64(3)	1.644(6)	1.65(4)	1.643(6)
O(1)–O(2)	1	2.67(3)	2.700(5)	2.84(3)	2.783(5)	2.67(2)	2.743(5)	2.85(3)	2.751(5)	2.87(3)	2.730(5)
O(1)–O(3)	1	2.65(2)	2.699(5)	2.96(3)	2.813(6)	2.78(2)	2.753(5)	2.70(2)	2.800(5)	2.67(3)	2.763(6)
O(1)–O(4)	1	2.39(5)	2.554(8)	2.61(5)	2.547(7)	2.24(3)	2.520(7)	2.37(5)	2.521(7)	2.48(5)	2.546(7)
O(2)–O(3)	1	2.41(3)	2.544(5)	2.52(3)	2.574(6)	2.43(2)	2.525(5)	2.69(3)	2.593(5)	2.69(4)	2.586(6)
O(2)–O(4)	1	2.70(4)	2.761(7)	2.69(4)	2.757(7)	2.58(3)	2.713(7)	2.82(4)	2.722(7)	2.74(5)	2.735(7)
O(3)–O(4)	1	2.56(4)	2.764(7)	2.82(5)	2.754(7)	2.48(3)	2.714(7)	2.67(4)	2.701(7)	2.69(6)	2.726(7)
<O–O> _{tet}		2.56(4)	2.670(6)	2.74(4)	2.705(6)	2.53(2)	2.661(6)	2.68(4)	2.681(6)	2.69(4)	2.681(6)
Y(1)–O(4)	6	2.10(2)	1.961(3)	2.01(2)	1.963(3)	2.18(1)	1.974(3)	2.01(2)	1.981(3)	2.08(2)	1.972(3)
O(4)–O(4)	6	3.08(3)	2.776(6)	2.88(3)	2.819(6)	3.27(2)	2.851(6)	2.91(3)	2.881(6)	3.01(4)	2.838(6)
O(4)–O(4)	6	2.85(3)	2.770(6)	2.80(4)	2.734(6)	2.88(3)	2.731(6)	2.76(3)	2.720(6)	2.86(4)	2.740(6)
<O–O>		2.96(3)	2.773(6)	2.84(4)	2.776(6)	3.08(2)	2.791(6)	2.84(3)	2.800(6)	2.94(4)	2.789(6)
Y(2)–O(1)	2	2.02(2)	1.969(4)	1.88(2)	1.956(4)	2.01(2)	1.977(4)	2.04(2)	1.962(4)	2.02(2)	1.972(4)
Y(2)–O(2)	2	2.02(3)	1.968(6)	2.01(3)	1.962(6)	2.01(3)	1.979(6)	1.89(4)	1.969(6)	1.88(4)	1.971(6)
Y(2)–O(3)	2	2.11(2)	1.963(3)	1.91(2)	1.956(4)	2.11(2)	1.971(3)	1.94(2)	1.964(3)	1.95(3)	1.967(4)
<Y(2)–O>		2.05(2)	1.967(4)	1.93(2)	1.958(5)	2.04(2)	1.976(4)	1.96(3)	1.965(4)	1.95(3)	1.970(5)
<Y–O>		2.08(2)	1.964(4)	1.97(2)	1.960(4)	2.11(2)	1.975(4)	1.98(2)	1.973(4)	2.02(2)	1.971(4)
O(1)–O(2)	2	2.90(4)	2.833(8)	2.68(5)	2.782(8)	2.90(4)	2.822(8)	2.84(5)	2.822(8)	2.76(5)	2.829(8)
O(1)–O(2)	2	2.82(3)	2.733(7)	2.82(4)	2.759(7)	2.78(3)	2.773(7)	2.72(3)	2.736(7)	2.75(4)	2.747(7)
O(1)–O(3)	2	2.86(3)	2.713(5)	2.74(3)	2.783(5)	2.95(2)	2.780(5)	2.83(3)	2.796(5)	2.76(4)	2.775(5)
O(1)–O(3)	2	2.99(3)	2.846(5)	2.62(3)	2.749(6)	2.89(3)	2.804(5)	2.79(2)	2.757(5)	2.85(3)	2.796(6)
O(2)–O(3)	2	3.01(4)	2.755(7)	2.74(5)	2.757(7)	2.97(4)	2.753(7)	2.72(5)	2.769(7)	2.74(6)	2.768(7)
O(2)–O(3)	2	2.84(3)	2.804(7)	2.80(4)	2.783(7)	2.86(3)	2.833(7)	2.69(4)	2.793(7)	2.67(6)	2.802(7)
<O–O>		2.90(3)	2.781(6)	2.73(4)	2.769(7)	2.89(3)	2.794(6)	2.76(4)	2.779(6)	2.76(5)	2.786(7)
<O–O> _{oct}		2.93(3)	2.777(6)	2.78(4)	2.772(6)	2.98(2)	2.792(6)	2.80(4)	2.790(6)	2.85(4)	2.788(6)
X(1)–O(1)	2	2.59(2)	2.503(5)	2.46(3)	2.495(5)	2.67(2)	2.513(5)	2.51(3)	2.492(5)	2.41(3)	2.484(5)
X(1)–O(2)	2	2.40(3)	2.500(6)	2.56(3)	2.496(6)	2.43(3)	2.510(6)	2.41(3)	2.492(6)	2.45(3)	2.487(6)
X(1)–O(2)	2	2.25(2)	2.328(4)	2.23(2)	2.312(4)	2.24(2)	2.318(4)	2.37(1)	2.340(4)	2.34(3)	2.343(4)
X(1)–O(3)	2	2.24(2)	2.332(4)	2.30(3)	2.319(4)	2.23(2)	2.321(4)	2.39(3)	2.336(4)	2.38(4)	2.339(4)
<X(1)–O>		2.37(2)	2.416(5)	2.39(3)	2.406(5)	2.39(2)	2.416(4)	2.42(2)	2.415(5)	2.40(3)	2.413(5)
O(1)–O(1)	1	2.94(3)	2.843(6)	2.93(4)	2.935(6)	3.04(3)	2.904(6)	3.12(3)	2.942(6)	2.89(4)	2.876(6)
O(1)–O(2)	2	4.12(4)	4.169(8)	4.08(5)	4.105(8)	4.19(4)	4.158(8)	4.01(5)	4.119(8)	3.99(5)	4.129(8)
O(1)–O(2)	2	2.90(3)	2.833(5)	2.68(3)	2.782(5)	2.90(2)	2.822(5)	2.84(3)	2.822(5)	2.76(3)	2.829(5)
O(1)–O(2)	2	3.52(3)	3.474(6)	3.40(3)	3.466(6)	3.50(3)	3.436(6)	3.56(2)	3.503(6)	3.49(4)	3.487(6)
O(1)–O(3)	2	2.90(5)	2.996(8)	2.90(6)	2.914(8)	2.84(4)	2.930(8)	2.96(5)	2.920(8)	2.95(7)	2.933(8)
O(2)–O(2)	1	2.99(2)	2.856(6)	3.09(3)	2.904(6)	3.00(2)	2.911(6)	2.76(2)	2.831(6)	2.80(4)	2.824(6)
O(2)–O(2)	2	2.67(5)	2.936(9)	2.89(5)	2.926(9)	2.77(4)	2.960(9)	2.92(5)	2.933(9)	2.89(5)	2.930(9)
O(2)–O(3)	2	3.33(4)	3.432(8)	3.51(5)	3.472(8)	3.45(4)	3.470(8)	3.43(5)	3.464(8)	3.44(6)	3.458(8)
O(2)–O(3)	2	2.84(3)	2.804(5)	2.80(3)	2.783(8)	2.86(3)	2.833(5)	2.69(3)	2.793(5)	2.67(4)	2.802(5)
O(2)–O(3)	2	2.41(3)	2.544(5)	2.52(3)	2.574(7)	2.43(2)	2.525(5)	2.69(3)	2.593(5)	2.69(4)	2.586(7)
<O–O>		3.07(4)	3.115(7)	3.09(4)	3.105(7)	3.11(3)	3.116(7)	3.12(4)	3.115(7)	3.08(5)	3.112(7)
X(2)–O(1)	2	2.29(2)	2.334(4)	2.34(3)	2.317(5)	2.27(2)	2.324(5)	2.19(2)	2.330(5)	2.28(3)	2.340(5)
X(2)–O(3)	2	2.48(2)	2.495(5)	2.46(3)	2.492(5)	2.40(2)	2.506(5)	2.57(2)	2.499(5)	2.52(3)	2.484(5)
X(2)–O(4)	2	2.45(2)	2.337(5)	2.31(2)	2.317(5)	2.40(2)	2.326(5)	2.30(2)	2.329(5)	2.40(2)	2.338(5)
X(2)–O(4)	2	2.53(2)	2.499(5)	2.47(2)	2.489(5)	2.65(2)	2.514(5)	2.54(2)	2.499(5)	2.49(2)	2.487(5)
<X(2)–O>		2.44(2)	2.416(5)	2.40(2)	2.404(5)	2.43(2)	2.418(5)	2.40(2)	2.414(5)	2.42(2)	2.412(5)
<X–O>		2.40(2)	2.416(5)	2.40(2)	2.405(5)	2.41(2)	2.417(5)	2.41(2)	2.414(5)	2.41(2)	2.412(5)
O(1)–O(1)	1	4.09(3)	4.062(6)	3.86(4)	3.919(6)	4.06(3)	3.963(6)	3.77(3)	3.933(6)	3.92(4)	3.989(6)
O(1)–O(3)	2	2.93(3)	2.846(5)	2.62(3)	2.749(6)	2.89(3)	2.804(5)	2.79(2)	2.757(5)	2.85(3)	2.796(6)
O(1)–O(3)	2	2.90(3)	2.996(6)	2.90(4)	2.914(6)	2.84(3)	2.930(6)	2.96(4)	2.920(6)	2.95(5)	2.933(6)
O(1)–O(4)	2	2.39(4)	2.554(7)	2.61(5)	2.547(7)	2.24(3)	2.520(7)	2.37(4)	2.521(7)	2.48(5)	2.546(7)
O(1)–O(4)	2	3.29(3)	3.451(7)	3.42(4)	3.445(7)	3.29(3)	3.461(7)	3.38(4)	3.438(7)	3.36(4)	3.451(7)
O(3)–O(3)	1	4.18(2)	4.180(5)	3.94(3)	4.071(6)	3.99(3)	4.126(5)	4.30(3)	4.078(5)	4.25(4)	4.099(6)
O(3)–O(4)	2	2.83(5)	2.881(8)	2.88(5)	2.873(8)	2.90(4)	2.883(8)	2.90(5)	2.845(8)	2.78(7)	2.835(8)
O(3)–O(4)	2	3.47(3)	3.474(6)	3.47(3)	3.445(6)	3.32(3)	3.444(6)	3.47(3)	3.438(5)	3.57(5)	3.450(6)
O(4)–O(4)	2	3.08(3)	2.776(4)	2.88(4)	2.819(4)	3.27(2)	2.851(4)	2.91(4)	2.881(4)	3.01(4)	2.838(4)
O(4)–O(4)	2	2.94(2)	2.911(6)	2.92(3)	2.983(6)	3.01(2)	2.996(6)	2.99(2)	3.026(6)	2.90(3)	2.980(6)
<O–O>		3.11(3)	3.112(6)	3.07(4)	3.086(6)	3.09(3)	3.104(6)	3.09(3)	3.092(6)	3.11(4)	3.097(6)
<O–O> _{dod}		3.09(4)	3.114(6)	3.08(4)	3.096(6)	3.10(3)	3.110(6)	3.10(4)	3.104(6)	3.10(4)	3.104(6)
<D–O>		2.12(2)	2.108(5)	2.11(2)	2.107(5)	2.12(2)	2.110(5)	2.11(2)	2.111(5)	2.12(2)	2.110(5)

* – the numbers in parentheses are the estimated standard deviations multiplied with SCOR and refer to the last significant number; ^s – frequency of occurrences; ¹ – Tančić et al. (2020); <D–O> = {<Si–O> + <Y–O> + 2 × <X–O>}/4

APPENDIX 7

Selected angles (in °) for the $R\bar{3}c$ space group [without (a) and with (b) constraints]

Zone		A (core) ¹		B		C		D		E (rim)	
		a	b	a	b	a	b	a	b	a	b
O(1)–Si–O(2)	1 ^s	107(2)	110.5(4) ¹	115(2)	114.2(4)	111(2)	114.1(4)	117(2)	113.8(4)	115(2)	112.4(4)
O(1)–Si–O(3)	1	117(2)	110.9(4)	118(2)	116.0(5)	122(2)	115.2(4)	117(2)	117.5(4)	112(2)	114.5(5)
O(1)–Si–O(4)	1	107(3)	102.5(5)	104(3)	100.4(5)	100(2)	101.0(5)	93(3)	100.0(5)	102(4)	101.6(5)
O(2)–Si–O(3)	1	92(2)	102.0(4)	95(2)	101.9(4)	95(2)	101.6(4)	108(2)	104.3(4)	104(3)	103.9(4)
O(2)–Si–O(4)	1	114(3)	115.2(5)	111(3)	112.6(5)	116(3)	112.5(5)	111(3)	111.1(5)	109(3)	112.7(3)
O(3)–Si–O(4)	1	118(3)	115.9(5)	114(3)	112.3(5)	114(3)	113.0(5)	110(3)	110.3(5)	115(4)	112.2(5)
<O–Si–O>		109(2)	109.5(4)	110(2)	109.6(5)	110(2)	109.6(4)	109(2)	109.5(4)	110(3)	109.6(4)
O(4)–Y(1)–O(4)	6	94(2)	90.1(3)	92(2)	91.8(3)	97(1)	92.4(3)	93(2)	93.3(3)	93(2)	92.0(3)
O(4)–Y(1)–O(4)	6	86(2)	89.9(3)	88(2)	88.2(3)	83(1)	87.6(3)	87(2)	86.7(3)	87(2)	88.0(3)
<O–Y(1)–O>		90(2)	90.0(3)	90(2)	90.0(3)	90(1)	90.0(3)	90(2)	90.0(3)	90(2)	90.0(3)
O(1)–Y(2)–O(2)	2	92(2)	92.0(4)	87(2)	90.5(4)	92(2)	91.0(4)	92(2)	91.4(2)	90(3)	91.7(4)
O(1)–Y(2)–O(3)	2	88(2)	88.0(3)	93(2)	89.5(3)	88(2)	89.0(3)	88(2)	88.6(2)	90(2)	88.3(3)
O(1)–Y(2)–O(3)	2	88(1)	87.3(3)	92(2)	90.7(3)	91(1)	89.5(3)	91(2)	88.2(1)	88(2)	89.6(3)
O(1)–Y(2)–O(3)	2	92(1)	92.7(3)	88(2)	89.3(3)	89(1)	90.5(3)	89(1)	91.8(1)	92(2)	90.4(3)
O(2)–Y(2)–O(3)	2	93(2)	89.0(4)	89(2)	89.5(4)	92(2)	88.4(4)	91(2)	89.5(4)	91(3)	89.3(4)
O(2)–Y(2)–O(3)	2	87(2)	91.0(3)	91(2)	90.5(4)	88(2)	91.6(3)	89(2)	90.5(3)	89(3)	90.7(4)
<O–Y(2)–O>		90(2)	90.0(3)	90(2)	90.0(4)	90(2)	90.0(3)	90(2)	90.0(2)	90(2)	90.0(4)
<O–Y–O>		90(2)	90.0(3)	90(2)	90.0(4)	90(2)	90.0(3)	90(2)	90.0(2)	90(2)	90.0(4)
O(1)–X(1)–O(1)	1	69(1)	69.2(2)	73(2)	72.0(3)	69(1)	70.6(2)	77(1)	72.4(3)	74(2)	70.8(3)
O(1)–X(1)–O(2)	2	111(2)	112.8(3)	109(2)	110.7(3)	110(2)	111.8(3)	109(2)	111.5(3)	110(2)	112.3(3)
O(1)–X(1)–O(2)	2	73(1)	71.7(2)	70(1)	70.6(2)	71.7(9)	71.4(2)	71(1)	71.4(2)	71(1)	71.7(2)
O(1)–X(1)–O(2)	2	93(1)	91.9(3)	93(2)	92.2(3)	90(1)	90.6(3)	93(1)	92.9(3)	94(2)	92.5(3)
O(1)–X(1)–O(3)	2	73(1)	76.5(3)	75(2)	74.4(3)	70(1)	74.5(3)	74(2)	74.4(3)	76(2)	74.9(3)
O(2)–X(1)–O(2)	1	74(1)	69.7(2)	74(1)	71.2(2)	76(1)	70.9(2)	70(1)	69.2(2)	70(2)	69.2(2)
O(2)–X(1)–O(2)	2	70(2)	74.8(3)	74(2)	74.9(3)	72(2)	75.5(3)	75(2)	74.7(3)	74(2)	74.6(3)
O(2)–X(1)–O(3)	2	92(2)	90.4(3)	92(2)	92.2(3)	96(2)	91.7(3)	91(2)	91.7(3)	91(3)	91.5(3)
O(2)–X(1)–O(3)	2	75(1)	70.8(2)	70(1)	70.5(3)	76(1)	71.7(2)	68(1)	70.6(2)	67(2)	70.9(2)
O(2)–X(1)–O(3)	2	65(1)	66.2(2)	68(1)	67.5(2)	65.8(9)	66.0(2)	69(1)	67.4(2)	69(2)	67.1(2)
<O–X(1)–O>		80(1)	80.5(3)	80(2)	80.5(3)	81(1)	80.4(3)	80(1)	80.6(3)	80(2)	80.6(3)
O(1)–X(2)–O(1)	1	127(2)	121.0(3)	112(2)	115.5(3)	126(1)	117.0(3)	119(2)	115.1(3)	119(2)	117.0(3)
O(1)–X(2)–O(3)	2	78(1)	72.1(2)	66(1)	69.6(2)	76(1)	70.9(2)	71(1)	69.5(2)	73(1)	70.8(2)
O(1)–X(2)–O(3)	2	75(1)	76.6(2)	74(2)	74.5(3)	75(1)	74.6(2)	76(2)	74.3(2)	76(2)	74.8(3)
O(1)–X(2)–O(4)	2	61(1)	66.3(3)	68(2)	66.7(3)	57(1)	65.6(3)	64(2)	65.5(3)	64(2)	66.0(3)
O(1)–X(2)–O(4)	2	86(1)	91.1(3)	91(2)	91.5(3)	84(1)	91.3(3)	90(2)	90.7(3)	90(2)	91.2(3)
O(3)–X(2)–O(3)	1	113(1)	113.8(2)	107(2)	109.6(3)	112(1)	110.8(2)	114(1)	109.4(2)	115(2)	111.2(3)
O(3)–X(2)–O(4)	2	69(2)	70.5(3)	71(2)	70.5(3)	70(1)	70.1(3)	69(2)	69.4(3)	67(2)	69.5(3)
O(3)–X(2)–O(4)	2	90(1)	91.9(2)	94(2)	91.4(2)	88(1)	90.8(2)	90(1)	90.8(2)	93(2)	91.3(2)
O(4)–X(2)–O(4)	2	76(1)	70.0(2)	74(2)	71.7(2)	80.7(9)	72.0(2)	73(1)	73.2(2)	76(2)	72.0(2)
O(4)–X(2)–O(4)	2	72(1)	73.9(2)	75(1)	76.6(2)	73.1(9)	76.4(2)	76(1)	77.5(2)	73(1)	76.2(2)
<O–X(2)–O>		81(1)	81.1(2)	80(2)	80.6(3)	80(1)	80.6(2)	81(2)	80.4(2)	81(2)	80.7(3)
<O–X–O>		80(1)	80.8(2)	80(2)	80.6(3)	80(1)	80.5(2)	80(2)	80.5(2)	80(2)	80.6(3)

* – the numbers in parentheses are the estimated standard deviations multiplied with SCOR and refer to the last significant number; ^s – frequency of occurrences; ¹ – Tančić et al. (2020)

APPENDIX 8

Number (N°) of calculated Bragg's positions (Y_{calc}) and differences between starting and ending corresponding Y_{calc} 's (ΔY_{calc} , in $^{\circ}$)

<i>hkl</i>	<i>10 4 0</i> (-)		<i>10 4 2</i> (-)		<i>8 8 0</i> (+)		<i>8 8 4</i> (+)		<i>12 2 0</i> (-)		<i>12 2 2</i> (+)		<i>8 8 8</i> (+)	
Zone	N°	ΔY_{calc}	N°	ΔY_{calc}	N°	ΔY_{calc}	N°	ΔY_{calc}	N°	ΔY_{calc}	N°	ΔY_{calc}	N°	ΔY_{calc}
A ¹	6	0.057	4	0.043	2	0.046	4	0.071	2	0.018	7	0.074	2	0.116
B		0.118		0.088		0.094		0.143		0.036		0.152		0.234
C		0.080		0.060		0.065		0.098		0.024		0.104		0.161
D		0.058		0.043		0.046		0.071		0.018		0.075		0.116
E		0.145		0.109		0.116		0.177		0.044		0.187		0.289

¹ – Tanić et al. (2020); all reflections are marked with *hkl*'s belonging to the $Ia\bar{3}d$ space group; the symbols “+” and “-” denote the presence and the absence of doublets, respectively

APPENDIX 9

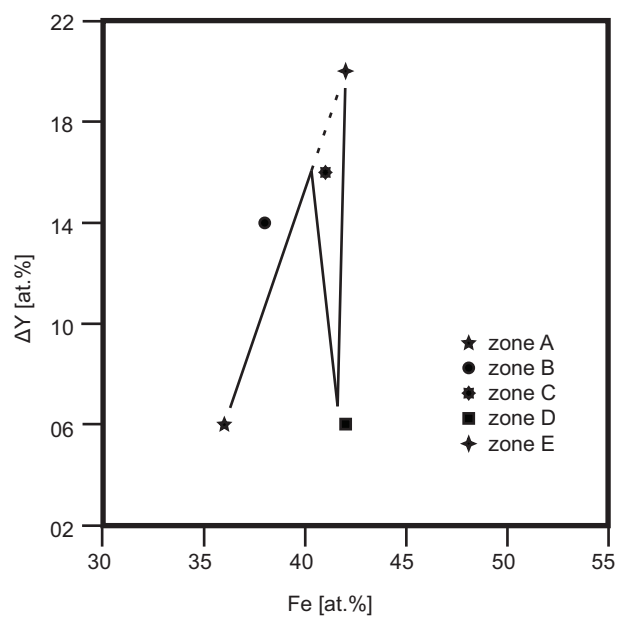
Differences between Al *sof*'s (in at. %), and average <Y–O> and <O–O> distances (in Å) in octahedral sites and calculated degree of ordering (*m*) and intersite distribution coefficient (K_{1-2}) for the $R\bar{3}c$ and $Fddd$ space groups [without (a) and with (b) constraints]

Zone		A (core) ¹		B		C		D		E (rim)	
		a	b	a	b	a	b	a	b	a	b
Δ Al	$R\bar{3}c$	6(8)	21(2)	14(12)	2(6)	16(8)	9(3)	6(4)	7(2)	20(8)	3(6)
	$Fddd$	4(8)	4(10)	0(10)	4(12)	2(8)	6(8)	4(5)	4(4)	4(8)	8(1)
Δ <Y–O>	$R\bar{3}c$	0.05(2)	0.006(4)	0.08(2)	0.005(4)	0.14(2)	0.002(4)	0.05(2)	0.016(4)	0.13(2)	0.002(4)
	$Fddd$	0.02(3)	0.000(4)	0.02(3)	0.001(4)	0.04(3)	0.000(4)	0.02(3)	0.005(4)	0.03(4)	0.001(4)
Δ <O–O>	$R\bar{3}c$	0.06(3)	0.008(6)	0.11(4)	0.007(6)	0.19(2)	0.003(3)	0.08(4)	0.021(6)	0.18(4)	0.003(6)
	$Fddd$	0.04(4)	0.001(6)	0.02(4)	0.001(6)	0.06(4)	0.001(6)	0.02(4)	0.007(6)	0.05(6)	0.000(6)
<i>m</i>	$R\bar{3}c$	0.037	0.165	0.115	0.016	0.129	0.076	0.048	0.043	0.192	0.026
	$Fddd$	0.024	0.031	0	0.032	0.015	0.051	0.031	0.034	0.032	0.069
K_{1-2}	$R\bar{3}c$	1.481	2.524	1.810	1.088	1.988	1.454	1.294	1.333	2.253	1.131
	$Fddd$	1.312	1.190	1	1.185	1.095	1.282	1.190	1.179	1.185	1.391

* – the numbers in parentheses are the estimated standard deviations multiplied with SCOR and refer to the last significant number; ¹ – Tančić et al. (2020)

APPENDIX 10

Variations of the Fe^{3+} and Al^{3+} over the octahedral sites in the $R\bar{3}c$ space group by ΔAl



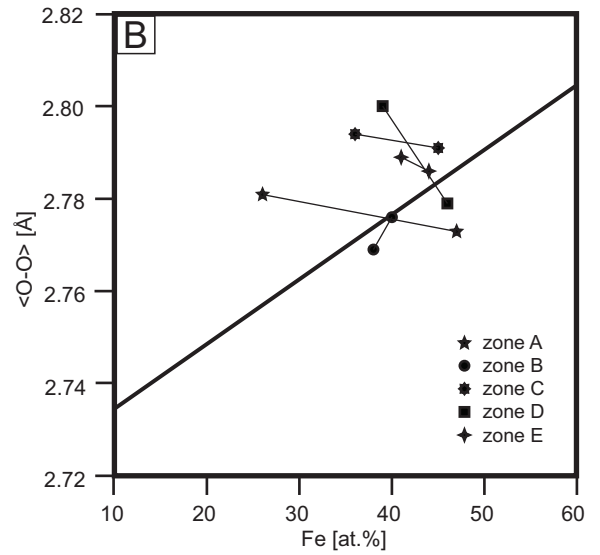
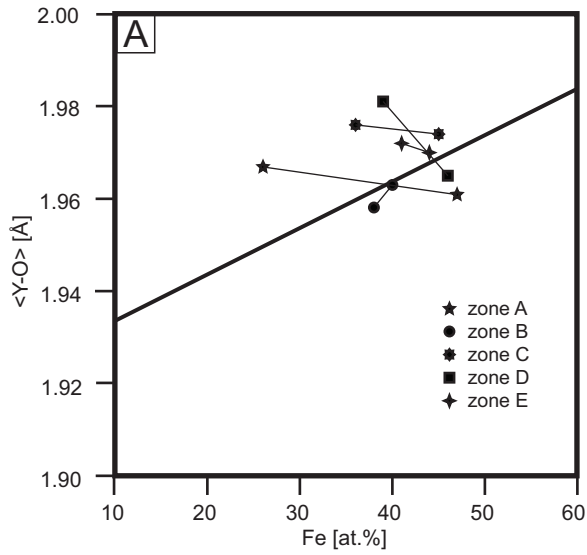
APPENDIX 11

Determined Si–Y1, Si–X2 and Y1–X2 distances (in Å)

	A (core)	B	C	D	E (rim)
Si–Y1	3.24(1)	3.32(1)	3.31(2)	3.44(2)	3.35(2)
Si–X2	3.38(1)	3.43(2)	3.42(1)	3.49(2)	3.44(2)
Y1–X2	3.377(6)	3.308(7)	3.361(4)	3.327(8)	3.358(5)

APPENDIX 12

Variations of the Fe^{3+} and Al^{3+} over the octahedral sites in the $R\bar{3}c$ space group, by average:
 A – $\langle Y-O \rangle$; B – $\langle O-O \rangle$ distances with constraints



Linear Grs-Adr joins (Novak and Gibbs, 1971) were also plotted for comparison

APPENDIX 13

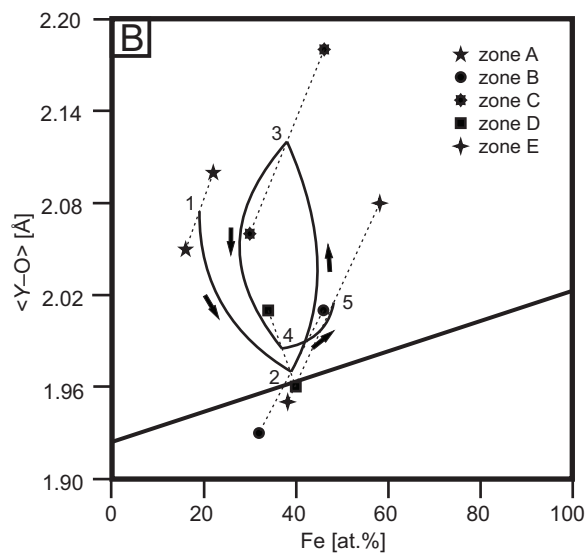
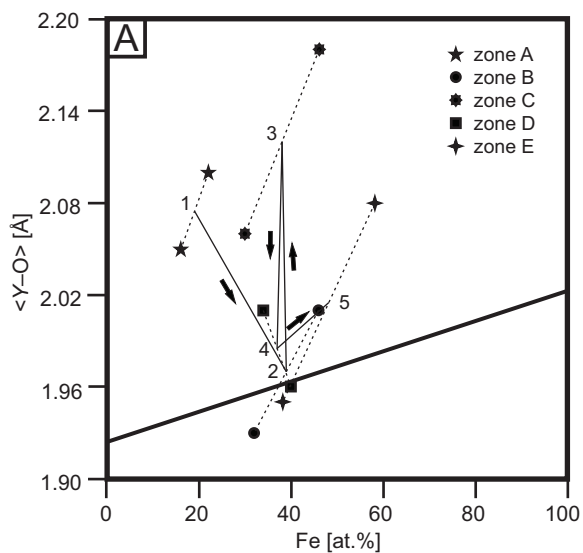
Calculated polyhedral volumes, quadratic elongations and angle variances in the tetrahedron and two octahedrons for the $R\bar{3}c$ space group [without (a) and with (b) constraints]

	Zone	A (core) ¹		B		C		D		E (rim)	
		a	b	a	b	a	b	a	b	a	b
Tetrahedron	polyhedral volume	1.94156	2.22042	2.36897	2.30182	1.83061	2.19356	2.22672	2.24588	2.26404	2.24952
	quadratic elongation	1.02945	1.00892	1.02100	1.01101	1.03708	1.01028	1.02401	1.00103	1.01163	1.00722
	angle variance	92.57748	36.87321	75.59731	44.99269	106.96640	42.07073	79.89321	40.74125	28.39711	29.40202
Octahedron-1	polyhedral volume	12.19868	10.04237	10.78950	10.07710	13.44318	10.22631	10.72094	10.31607	11.88520	10.21280
	quadratic elongation	1.00632	1.00000	1.00073	1.00096	1.01843	1.00189	1.00268	1.00344	1.00259	1.00126
	angle variance	21.30664	0.01247	2.55438	3.37979	58.95992	6.54798	9.23038	11.80922	8.93208	4.41588
Octahedron-2	polyhedral volume	11.48921	10.12718	9.59907	10.00429	11.34777	10.27427	9.92271	10.11234	9.82322	10.19424
	quadratic elongation	1.00292	1.00131	1.00341	1.00011	1.00232	1.00041	1.00263	1.00041	1.00226	1.00037
	angle variance	7.32937	4.60396	6.14373	0.36546	3.94984	1.45068	2.53165	1.48096	1.90723	1.32142

¹ – Tančić et al. (2020)

APPENDIX 14

**Different presentation models of the Fe³⁺ and Al³⁺ variations over the octahedral sites
by average <Y-O> distances: A – (pseudo) cyclical of the oscillatory nature;
B – (pseudo) rotational or spiral**



Linear Grs-Adr joins (Novak and Gibbs, 1971) were also plotted for comparison