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Polymorphism of K_2ZrF_6

Lubomír Smrčok^{1, †}, Armel Le Bail², Miroslav Boca^{*, 1}, Aydar Rakhmatullin³

¹ *Institute of Inorganic Chemistry, Department of molten systems, Dubravská cesta 9, Bratislava; 845 36, Slovakia, miroslav.boca@savba.sk*

² *Institut des Molécules et des Matériaux du Mans, CNRS UMR 6283, Université du Maine, Avenue Olivier Messiaen, 72085 Le Mans Cedex 9, France, Armel.le_Bail@univ-lemans.fr*

³ *Conditions Extremes et Matériaux: Haute Temperature et Irradiation, 1D avenue de la Recherche Scientifique CS 90055, 45071 Orleans cedex 2, France, aydar.rakhmatullin@cnsr-orleans.fr*

* corresponding author: Miroslav Boča, miroslav.boca@savba.sk

† in memoriam: Lubomír Smrčok

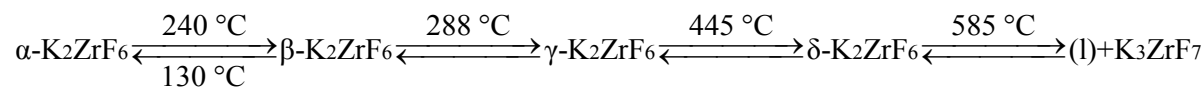
KEYWORDS: di-potassium zirconium fluoride, polymorphism, synchrotron powder diffraction, ¹⁹F MAS NMR

ABSTRACT: Heating of the K_2ZrF_6 room temperature monoclinic form-I reveals a complex polymorphism investigated by synchrotron powder temperature-dependent diffraction. The first thermal event at 240 °C corresponds to the quasi-simultaneous apparition of four different phases (noted II to V) of which only three could be indexed and two completely characterized structurally. Form-II (structure unrefined) is very probably hexagonal ($a = 6.4473(1)$ Å, $c = 3.8606(1)$ Å at 264 °C, $Z = 1$). Form-III is cubic (space group $Fm\bar{3}m$, $a = 9.0804(1)$ Å at 264 °C, $Z = 4$), related to K_3ZrF_7 . Form-IV remains unindexed, presenting broad diffraction peaks. Form-V is present as a weak impurity at this stage, increasing in proportion after the second thermal event at 288 °C where a new form-VI is built. Form-V is tetragonal (space group $P4/nmm$, $a = 6.3281(1)$ Å, $c = 9.3671(2)$ Å at 295 °C, $Z = 2$) and form-VI is orthorhombic (space group $Cmcm$, $a = 6.40840(5)$ Å, $b = 20.7697(2)$ Å, $c = 19.9888(2)$ Å, at 295 °C, $Z = 16$). At 317 °C, form-VI transforms into form-V. At 449 °C, a different orthorhombic form-VII is disclosed (space group $Pnma$, $a = 10.3084(1)$ Å,

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3 $b = 6.0291(1) \text{ \AA}$, $c = 23.8023(2) \text{ \AA}$, at $495 \text{ }^\circ\text{C}$, $Z = 8$), coexisting with the cubic form-III. On
4 cooling, form VII of which an approach of the crystal structure is provided, returns to form V at
5 $411 \text{ }^\circ\text{C}$ which looks stable down to $139 \text{ }^\circ\text{C}$ giving form I back. Isolated ZrF_6 octahedra are present
6 in forms-III, -V, -VI, -VII. Form-VI contains also dimeric entities Zr_2F_{12} built from a ZrF_7
7 monocapped trigonal prism sharing a corner with a ZrF_6 octahedron. Form-VII structure is related
8 to form-V by quadrupling the tetragonal a parameter, the reversible $\text{V} \leftrightarrow \text{VII}$ transition is very
9 probably topotactic. The coexistence of several phases at each temperature above the first
10 transition at $240 \text{ }^\circ\text{C}$ was observed. Peak multi-splitting is observed at many temperatures that
11 suggests the occurrence of coexisting micro-phases. The parallel existence of several phases was
12 observed also by ^{19}F MAS NMR spectroscopy measured from $250 \text{ }^\circ\text{C} \rightarrow 410 \text{ }^\circ\text{C} \rightarrow \text{RT}$.
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22 INTRODUCTION

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24 In the concept of so-called “Generation IV” Molten Salts Nuclear Reactors (MSR) fission power
25 should be produced in a circulating molten salt fuel mixture. Currently, these types of reactors are
26 considered to be fuelled with uranium or thorium fluorides dissolved in a mixture of molten
27 fluorides, with Na/K and Zr fluorides being a primary option ¹. The compounds belonging to the
28 KF-ZrF₄ system have several potential uses: i) as solvent media in a reactor core or as solvent for
29 pyrochemical reprocessing of actinides and lanthanides produced in nuclear reactions; ii) as
30 moderators; iii) as cooling media of the reactor core, iv) as high temperature transport media
31 providing transport of heat to hydrogen production facilities employed for the Hydrogen Economy
32 Concept ². In all of the above applications the compatibility of the molten salt with reactor
33 construction materials is necessary to avoid chemical attack by a corrosive liquid. Considering the
34 working regime of a MSR it is evident that reliable reference data on the dynamics of phase
35 transitions in zirconium fluorides are essential for design of a cooling unit. However, the only
36 available phase diagram of KF-ZrF₄ system was reported in 1961 ³ and although it has brought
37 initial information, some results are seriously limited by the use of low resolution laboratory
38 powder diffraction. None of the high temperature phases was the subject of a crystal structure
39 refinement by the Rietveld method. Among the compounds belonging to the KF-ZrF₄ system it is
40 K₂ZrF₆ which seems to be, mainly for the economic reasons, the most promising candidate for
41 future application. For its solid-solid phase transformations the following scheme was suggested
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For the region above 585 °C a series of phase transformations resulting in the co-existence of solid and liquid phases was suggested, but the region has not yet been explored in details. At room temperature, the crystal structure of K_2ZrF_6 is monoclinic, $C2/c$ ^{4,5}. The structure of $\beta\text{-K}_2\text{ZrF}_6$ is not reliably known in spite of several suggestions^{6,7}. Phase transitions in the temperature range of ~240 to 245 °C were independently confirmed only by NMR⁷. Recently, some more detailed analysis of phase transformation of K_2ZrF_6 was reported in the region up to 330 °C^{8,9} using XRD, DTA/DSC, Raman spectroscopy and statistical methods PCA and MCR, as well as phase diagram of the system $(\text{LiF-NaF-KF})_{\text{eut}}\text{-K}_2\text{ZrF}_6$ ¹⁰. Detailed room temperature analysis by XPS and NMR spectroscopy was provided, as well^{11,12}.

The very poor knowledge of the K_2ZrF_6 thermal behaviour regarding to its potential industrial and economic importance incited us to re-examine the problem by synchrotron temperature-dependent diffraction experiments. Our results led to the observation of seven K_2ZrF_6 phases instead of four, requiring to change the above Greek notation by using forms-I to VII. Correspondences with the polymorphism of K_2HfF_6 ¹³, for which structure models were proposed (but no Rietveld or single crystal structure refinement) for six different phases, are discussed.

EXPERIMENTAL SECTION

Preparation of K_2ZrF_6 : For the preparation of the title compound the following chemicals were used: ZrO_2 (Chiron, 98.5 %), KHF_2 (Lachema, 99%), KF (Sigma-Aldrich, 99 %), ZrF_4 (prepared at the Institute of Chemistry KSC RAS, Apatite, Russia; min. 99.5%; no impurities could be detected by powder XRD), HF (38–40 wt % in H_2O ; Lachema). KF was dried in vacuum at 130 °C for 24 h; other chemicals were used without further purification. The handling of KF and ZrF_4 was done in a glove box under dry argon atmosphere (Messer Tatragas, 99.9990 %) with moisture content below 10 ppm.

Zirconium oxide (0.01 mol, 1.232 g) was dissolved in hydrofluoric acid (10 wt % in H_2O ; 30 mL) by continuous stirring and heating (80–90 °C) in a platinum dish. Potassium hydrogen fluoride (0.02 mol, 1.562 g) was added and the reaction mixture was stirred for ca. one hour. The resulting clear solution was filtered through a folded paper filter into a plastic beaker and left to crystallise at ambient temperature. After two days, the oblong colourless crystals were removed

and dried in a desiccator at ambient temperature (1.147 g, 40 %). The studied compound can be prepared by the described procedure using zirconium fluoride (0.01 mol, 1.672 g) instead of zirconium oxide and/or potassium fluoride (0.02 mol, 1.162 g) instead of potassium hydrogen fluoride. It is crucial to use the right concentration of hydrofluoric acid. The use of highly concentrated solution is resulting into formation of solvated product $K_2ZrF_6 \cdot HF$ ¹⁴, while from pure water solution K_3ZrF_7 crystallises¹⁵. Two K_2ZrF_6 samples were made, noted samples **1** and **2** below, which were used for mainly two different synchrotron experiments (both fraction might differ in grain size).

DSC-TGA Analysis: Thermogravimetric experiments were carried out on a thermoanalyzer TA instrument DT-Q600 under argon at a heating rate of $5\text{ }^\circ\text{C min}^{-1}$ from room temperature up to $500\text{ }^\circ\text{C}$ and back, allowing to characterize a strong hysteresis (Figure 1). This DSC shows mainly four endothermic effect on heating (240 , 288 , 317 and $445\text{ }^\circ\text{C}$) and two exothermic effects on cooling (411 and $139\text{ }^\circ\text{C}$). If one does not consider the $317\text{ }^\circ\text{C}$ peak, the heating seems to be compatible with the previous model³ of $\alpha \rightarrow \beta \rightarrow \gamma \rightarrow \delta$ transitions given in the introduction, but not with any reversibility (also compare with⁹).

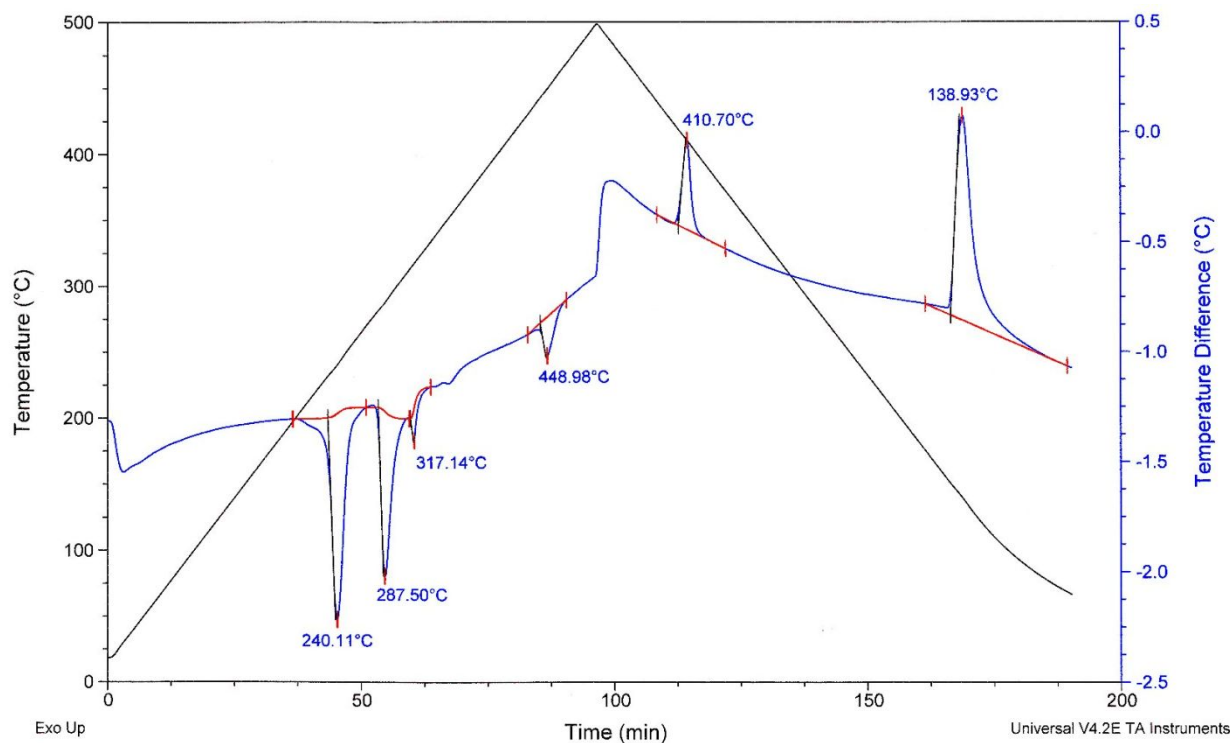


Figure 1. DSC thermal analysis of K_2ZrF_6 by DSC under argon at a heating rate of $5\text{ }^\circ\text{C min}^{-1}$ from room temperature up to $500\text{ }^\circ\text{C}$ and back.

¹⁹F MAS NMR: The High temperature (HT) MAS NMR experiments were acquired on a Bruker Avance III HD NMR spectrometer operating at 17.6 T using the previously described diode laser heated system ⁹. In addition, to enhance resolution at room temperature a probe with 1.3 mm diameter rotors spinning at a frequency of 50 kHz was used. The parameters were: pulse width 0.95 μs ($\pi/2$), relaxation delay 300 s ($6T_1$), and 4 scans. The ¹⁹F chemical shift was referenced to CFC1₃. The DMFIT program was applied to simulate the spectra and fit the peaks ¹⁶.

Powder X-ray Diffraction (PXRD): Data were recorded using beamline I11 at the Diamond Light Source, Didcot, UK. Samples were mounted in a sapphire capillary, in dry box under argon atmosphere, that was closed by glue on both sides for measurements in transmission geometry. Two main experiments were made, using the Mythen detector configuration on sample **1** and the multianalyzing crystal-detectors (MACs) on sample **2**, both operating at wavelength close to 0.827 Å. The beamline set-up and characteristics are described in the literature ¹⁷⁻¹⁹. The diffraction data were collected in order to obtain significant patterns in the temperature range 20–500 °C.

Data interpretations were sometimes facilitated by the previous results about K₂HfF₆ in 1983 by Saalfeld and Guse ¹³. From their GUINIER-powder diagrams (high-temperature chamber), they indexed a tetragonal phase (190-250 °C, $a = 6.29(8)$ Å, $c = 9.27(4)$ Å at 210 °C, suggested space group $P4/mmm$) and proposed it to be related to distorted variants of the cubic elpasolite ²⁰. KNaAlF₆ having similar cell parameters ($a \sim 6.5$ Å, $c \sim 9.5$ Å for (NH₄)₃MF₆, M = In, Sc) described by Bode and Voss ²¹. Saalfeld and Guse ¹³ identified a cubic K₂HfF₆ phase (210-380 °C, $a = 9.01(6)$ Å at 240 °C, space group $Fm\bar{3}m$, proposing a structure related to elpasolite or to K₂PtCl₆) ²² and sometimes a hexagonal phase (170-220 °C, $a = 6.40(5)$, $c = 3.81(8)$ at 200 °C, no systematic extinction) which they proposed to be similar to β_1 -K₂UF₆ and β_1 -K₂ThF₆, with the space group $P\bar{6}2m$, that were investigated by Zachariasen ²³. Saalfeld and Guse ¹³ noted that the γ -K₂ZrF₆ powder pattern from Novoselova *et al.* ³ was analogous to that of their hexagonal K₂HfF₆ phase, unfortunately they never confirmed their structure models by any Rietveld refinement. These K₂HfF₆ hexagonal, cubic and tetragonal three phases were recognized (using search-match identification process by means of the EVA software ²⁴ combined with the ICDD-PDF-4+ powder patterns database) ²⁵ to occur simultaneously in the case of K₂ZrF₆, with similar cell parameters as those of the Hf-based compounds, they were noted phases II, III and V, respectively. A fourth phase noted IV is also present at this stage, all of these phases corresponding to the " β -K₂ZrF₆" in the Novoselova *et al.* notation. ³.

A contour plot of a synchrotron thermo-diffractogram of K_2ZrF_6 covering the range 185-300 °C is shown in Figure 2, with phase numbering. This figure shows that the 288 °C thermal event corresponds to the complete collapse of the hexagonal phase-II (phase IV having not totally disappeared yet at 295 °C), a strong decrease in the proportion of the cubic phase-III, a strong increase in the proportion of the tetragonal phase-V and the occurrence of a new phase VI for which no similar powder pattern could be found in the K_2HfF_6 previous study.

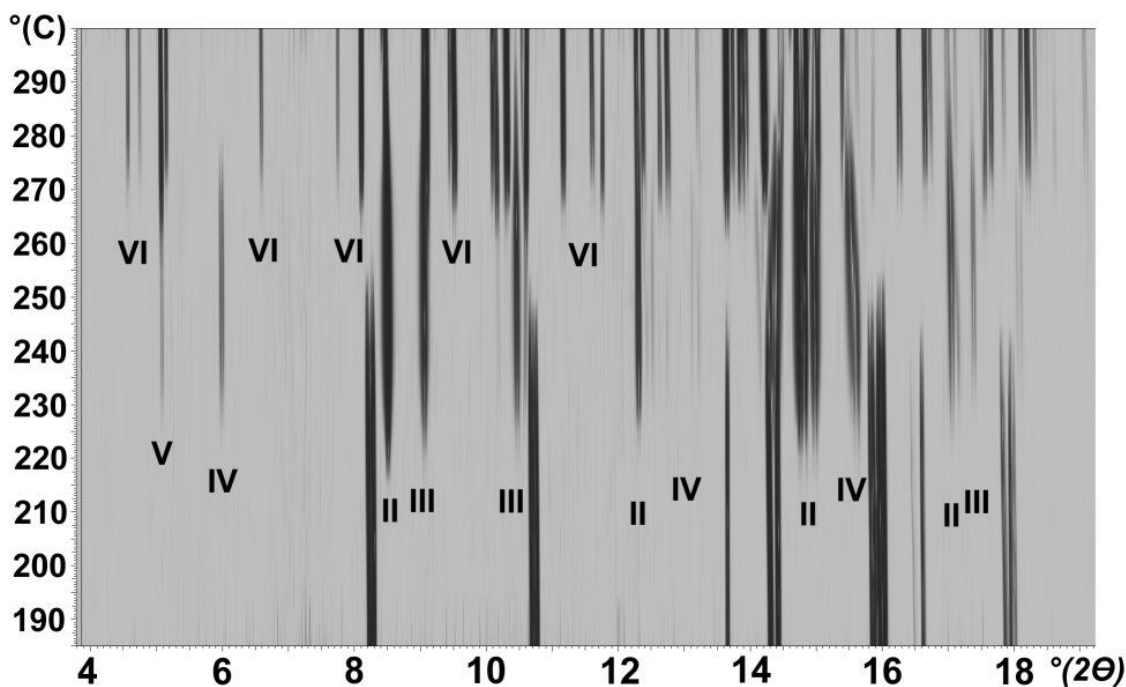


Figure 2. Contour plot from the synchrotron thermo-diffractometry on K_2ZrF_6 sample 1 in the range 185-300 °C, on heating.

The 317 °C DSC peak in Figure 1 is interpreted as the transition from form-VI to the tetragonal form-V which is the main phase up to 449 °C corresponding to the transition between the tetragonal form-V to a new phase VII accompanied by the reappearing of more of the cubic phase-III. On cooling, the 411 °C peak of the DSC corresponds to the reverse VII→V transition, with a strong hysteresis. The tetragonal phase-V is then stable on cooling down to 139 °C, where K_2ZrF_6 returns back to phase-I, monoclinic.

The new phases VI and VII were indexed by using the *McMaille*²⁶ software and were found to be both orthorhombic. Phase IV could never be indexed. It is to be noted that its diffraction peaks are quite broader than for phases II (showing the narrowest peaks) and III. Hypotheses that the

phase IV peaks could be satellites of either a commensurate or an incommensurate supercell of either phases II or III found no success, so it cannot be excluded that what we call form-IV is in fact a mixture of 2 phases. Then the *ab initio* structure determination of phases V, VI and VII were undertaken with success by using the direct space Monte Carlo software *ESPOIR*²⁷ (but the space group *P4/mmm* and the model proposed¹³ for the tetragonal K_2HfF_6 phase finally could not be applied to K_2ZrF_6 form-V). Rietveld²⁸ refinement as well as Le Bail fits²⁹ were realized by using the *FULLPROF*³⁰ software. The cubic structure of K_2ZrF_6 form-III was better Rietveld-refined at 495 °C when mixed with form-VII, using the K_3ZrF_7 cubic structure³¹ as a starting model. Attempts to refine the hexagonal structure of form-II using the various proposed models for K_2HfF_6 were never satisfying due to the presence of phases III, IV (the latter unindexed) and V. Crystallographic data for the K_2ZrF_6 forms I to VII are summarized in Table 1.

All the solid-state DFT structures optimizations were done according the procedure described elsewhere³².

Table 1. Crystallographic data for the K_2ZrF_6 forms I to VII (IV excluded)

	I	II	III	V	VI	VII
crystal system	monoclinic	hexagonal	cubic	tetragonal	orthorhombic	orthorhombic
space group	C2/c	?	Fm-3m	P4/nmm	Cmcm	Pnma
temperature (C)	22*	264	495	200**	295 / 54	495
(K)	295	537	768	473	568 / 327	768
<i>a</i> (Å)	6.56646(2)	6.4473(1)	9.15641(3)	6.33605(2)	6.40840(5) / 6.37122(8)	10.30844(7)
<i>b</i> (Å)	11.44440(4)				20.7697(2) / 20.5603(3)	6.02909(3)
<i>c</i> (Å)	6.94202(2)	3.8606(1)		9.29458(5)	19.9888(2) / 19.6642(3)	23.8017(2)
β (deg)	90.5668(3)					
<i>V</i> (Å ³)	521.661(3)	138.975(2)	767.675(5)	373.136(3)	2660.52(4) / 2575.90(6)	1479.29(2)
<i>Z</i>	4	1	4	2	16	8
<i>V/Z</i>	130.41	138.97	191.92	186.57	166.28/160.99	184.91
Wavelength (Å)	0.82563	0.82695	0.82695	0.82695	0.82711	0.82695
2 θ range (deg)	7-100		3-49	4-59	3-50 / 3-60	3-49
refined coordinates	13		3	6	32	32
soft restraints	none		none	none	none	18 (ZrF ₆)
<i>R_p</i>	3.82		5.72	7.73	3.04 / 3.10	5.72
<i>R_{wp}</i>	4.94		9.69	10.5	4.65 / 4.45	9.69
<i>R_B</i>	1.99		4.14	5.10	7.63 / 6.03	13.4
<i>R_F</i>	0.96		5.67	7.97	7.85 / 5.25	22.3
other phases	ZrO ₂ , K ₃ ZrF ₅ O ?	III, IV, V	VII, ZrO ₂	III, ZrO ₂	IV, V / V	III, ZrO ₂

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3 *after cooling from 470 °C
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5 **on cooling direction from 500 °C
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8 **RESULTS AND DISCUSSION**

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10 **K₂ZrF₆ form-I.** In the whole range between the room temperature and the DSC peak at 240 °C,
11 form-I (previously noted α -K₂ZrF₆) is found to be monoclinic ^{4, 5}, however, peak shapes are so
12 distorted and even split (in two or three components) that the Rietveld refinements are always quite
13 bad. There was some controversy in the past about the symmetry: monoclinic or orthorhombic,
14 monoclinic at room temperature and then possibly orthorhombic at higher temperature. For sample
15 **1**, the 0k0 reflections are split clearly in two main peaks (Figure 3), suggesting the existence of at
16 least two microphases presenting different cell parameters. The splitting is visible at all
17 temperatures. Comparison of samples **1** and **2** at 200 °C show that sample **2** looks much better than
18 sample **1** but presents itself a small 0k0 splitting (see the 060 below) and is not free of peak shape
19 problems (see the 133 or 422 in Figure 3). The Rietveld refinement for sample **2** at 200 °C led to
20 $R_p = 13.0\%$ and $R_B = 11.5\%$, and it was not better than $R_p = 35.1\%$ and $R_B = 24.3\%$ for sample
21 **1**. Our explanation about these differences is that sample **1** was too highly crushed and that sample
22 **2** was not enough crushed (the bad fit for sample **2** is not only due to broad peaks like the 422 or
23 133, but also to some abnormally intense peaks typical of the presence of single crystals in the
24 powder). Similar problems on sample grain size response was observed also on DTA/DSC
25 experiments ⁹.
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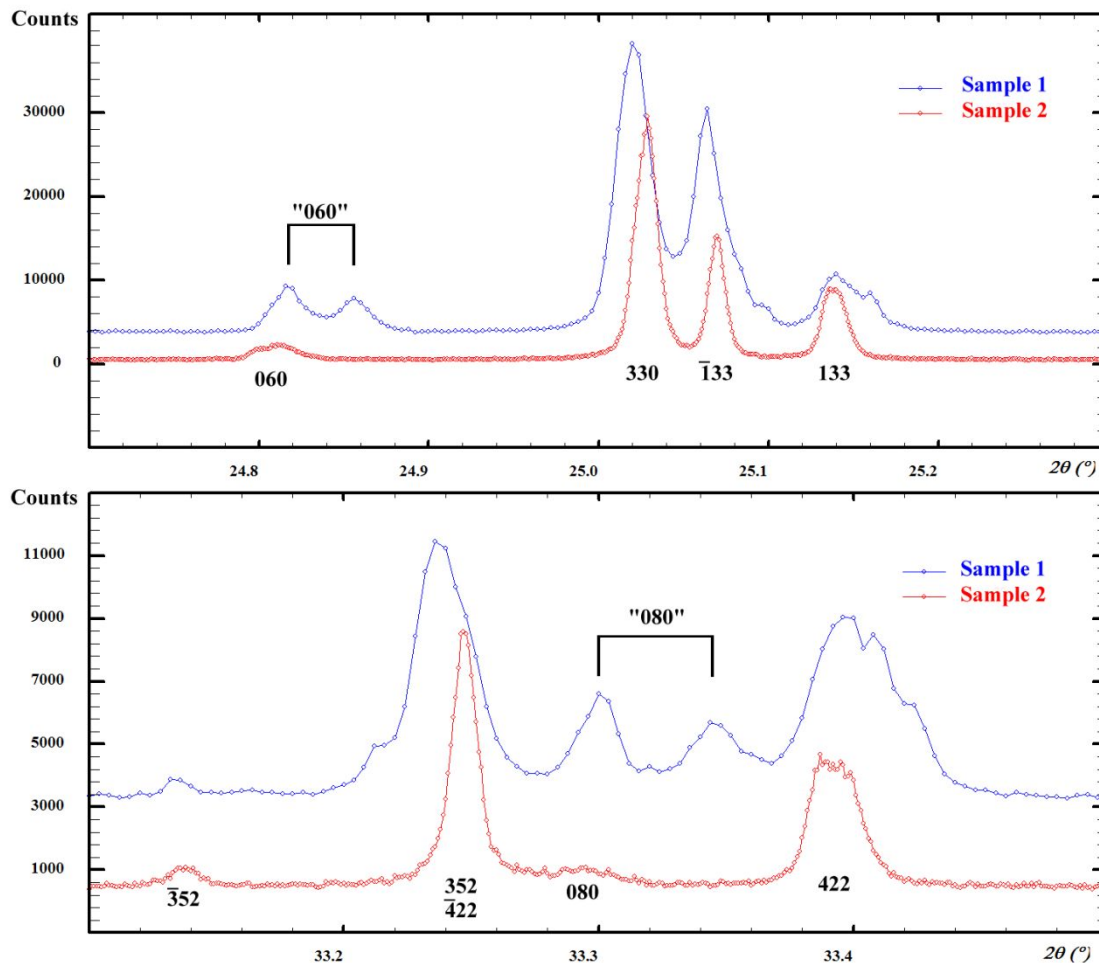


Figure 3. Parts of the synchrotron diffraction patterns of the two starting K_2ZrF_6 samples at 200 °C illustrating their differences and suggesting the coexistence of micro-phases with slightly different cell parameters.

Then, sample 2 was placed 12 h at 470 °C, then cooled freely in order to see if some high temperature phase could be quenched. The result was the form-I presenting broader peaks than for unheated sample 1 and 2, but which produced an excellent fit (Figure 4) by the Rietveld method in spite of some anisotropic broadening which was treated by separating the much narrower $h00$ and $hh0$ from the others. This sample contains monoclinic $P2_1/c$ ZrO_2 as impurity, and also a third phase never encountered before, having peaks at positions very close to those of the cubic form-III, but convincingly indexed in a rhomboedral cell with parameters $a = 6.34818(5)$, $c = 15.65071(13)$ Å, in too small proportion for an accurate structure refinement ($R_B \sim 10\%$ was obtained with the $R\bar{3}m$ space group suggesting the K_3ZrF_5O formula, to be confirmed), so it was treated by a Le Bail fit, whereas m - ZrO_2 was treated by using the atomic coordinates from the literature, fixed.

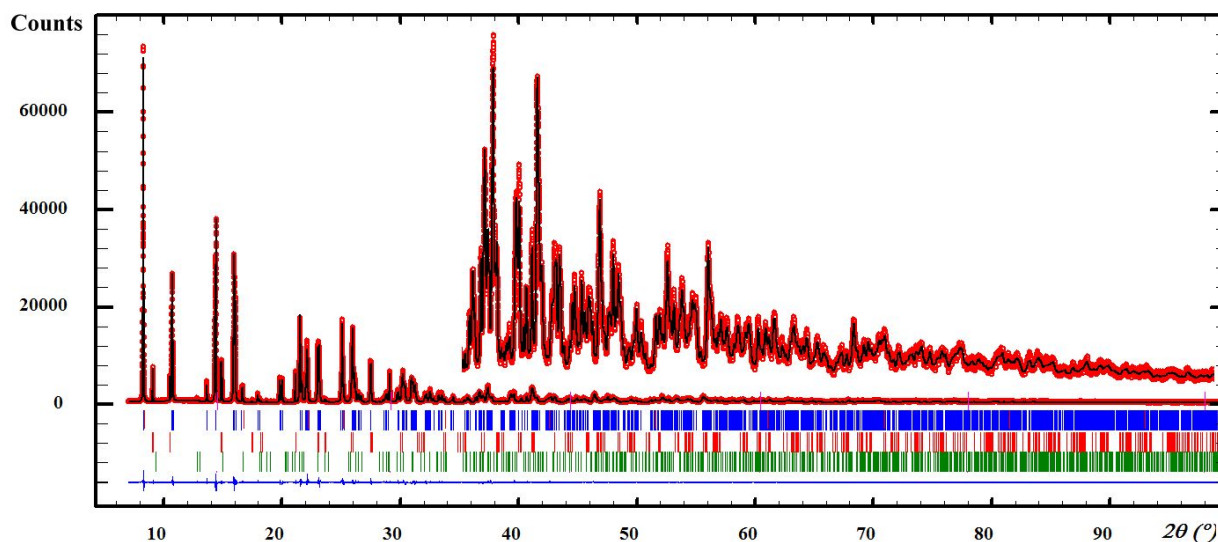


Figure 4. Refined diffraction pattern from synchrotron data for K_2ZrF_6 form-I at 22 °C after free cooling from 470 °C. Red dots represent the observed data, the black line represents the calculated ones. Bragg ticks are the peak positions (3 sets from top to bottom: reflections of form-I, new rhombohedral phase related the cubic K_3ZrF_7 , monoclinic ZrO_2) and the blue curve shows the difference between the observed and calculated patterns, $\lambda = 0.82563$ Å.

The Rietveld refinement from synchrotron data leads to similar results as from the two previous single crystal studies^{4,5} (none presenting anisotropic thermal parameters because of crystals being of low quality, which is compatible with our problems with sample **1** and **2** freshly prepared) and confirms the monoclinic cell (Table 2). Interatomic distances are in Table 3. The structure is built up from infinite chains of ZrF_8 square antiprisms sharing edges, extending along the c -axis (Figure 5). The tenfold coordination of the potassium is harder to define, but can be described as a strongly distorted cube having two adjacent faces capped.

Table 2. Crystallographic parameters for K_2ZrF_6 form-I at 22 °C (C2/c) after cooling from 470 °C. Second line, from Hoppe and Mehlhorn (1976)⁴ third line from Gerasimenko *et al.* (2006)⁵

atom	site	x	y	z	B_{iso} or U_{iso}
K	8f	0.01939(7)	0.34331(4)	0.00129(7)	1.725(9)
		0.0197(2)	0.3428(1)	0	0.84
		0.01844(5)	0.34287(3)	0.00238(3)	0.02044(6)
Zr	4e	0	0.04711(1) 0.04761(7)	1/4	0.167(4) 0.71

			0.04741(1)		0.01031(4)
F1	8f	0.28774(12)	-0.01238(7)	0.22121(13)	1.56(2)
		0.2889(4)	-0.014(3)	0.220(1)	0.91
		0.2895(1)	-0.0140(1)	0.2191(2)	0.0239(2)
F2	8f	0.19031(14)	0.19056(7)	0.27435(15)	2.17(2)
		0.1900(5)	0.1918(3)	0.274(1)	0.71
		0.1899(1)	0.1922(1)	0.2746(1)	0.0193(2)
F3	8f	-0.03405(15)	0.09824(6)	0.54599(11)	1.58(2)
		-0.0362(5)	0.0985(3)	0.539(1)	0.96
		-0.0336(1)	0.0986(1)	0.5450(1)	0.0193(2)

Table 3. Selected bond distances (angstroms) in K_2ZrF_6 form-I from synchrotron X-ray diffraction refinement

bond	length	bond	length	bond	length
Zr-F1 x 2	2.0201(9)	K-F2	2.6897(11)	K-F2	2.8048(11)
Zr-F2 x 2	2.0691(10)	K-F1	2.7236(10)	K-F3	2.8256(8)
Zr-F3 x 2	2.1503(8)	K-F2	2.7240(11)	K-F1	2.8274(10)
Zr-F3 x 2	2.1946(8)	K-F2	2.7418(12)	K-F3	3.0206(11)
average	2.1085	K-F1	2.7879(10)	K-F1	3.1272(10)
				average	2.8273

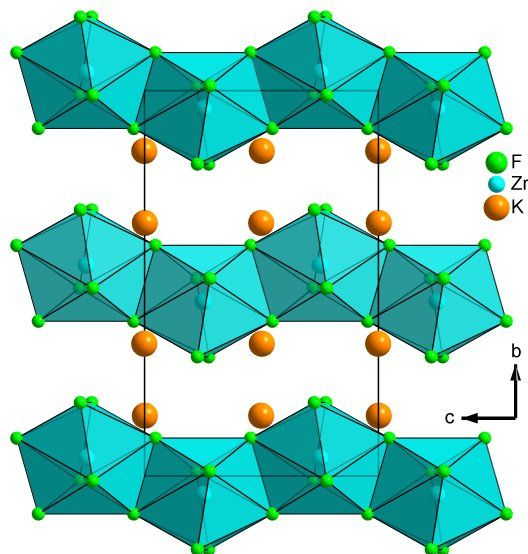
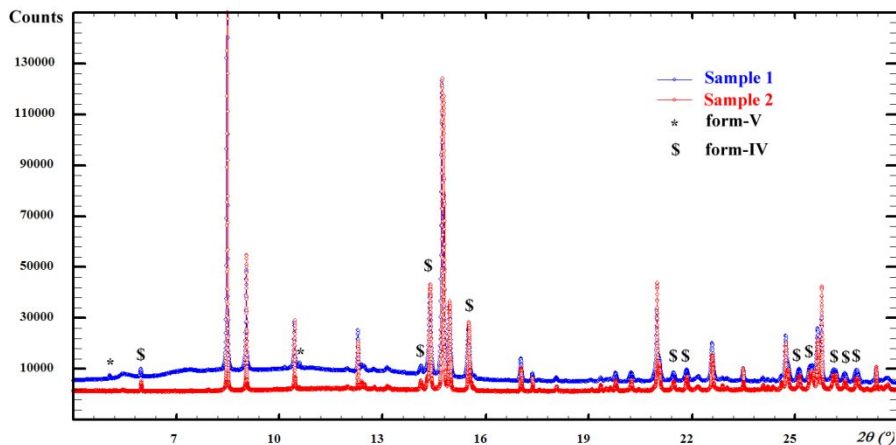


Figure 5. Projection of the unit cell of K_2ZrF_6 form-I along the a axis showing the infinite chains of ZrF_8 square antiprisms sharing edges.

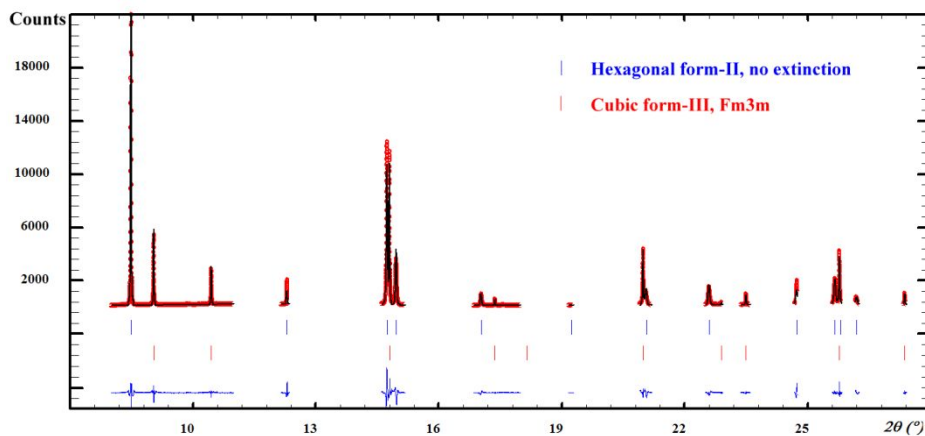
K_2ZrF_6 forms II, III and IV (= " β - K_2ZrF_6 "). In spite of their strong differences, the two " K_2ZrF_6 form-I" samples produce very similar results after the first DSC peak at 240°C. The

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3 comparison of a part of the two patterns at 265 °C is made in Figure 6. The diffraction peaks are
4 narrower for sample 2 but this is due more probably to a better instrumental resolution.
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21 **Figure 6.** Comparison of the two samples synchrotron patterns at 265 °C. The tetragonal form-V is more present in
22 the case of sample 1. Some of the main peaks corresponding to form-IV (unindexed) are indicated.
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26 Once a convincing interpretation by pattern matching followed by peak indexing was established
27 (see supplementary material) showing the simultaneous presence of up to four phases at this
28 temperature (265 °C), a Le Bail fit was realized, including forms II and III, using excluded zones
29 to discard forms IV (unindexed) and V, leading to the result shown in Figure 7. It can be noted
30 that no peak splitting is observed, and that the peaks of the cubic phase (form-III) are narrower
31 than those of the hexagonal form-II.
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51 **Figure 7.** Le Bail fit on selected zones of the sample 2 synchrotron pattern at 265 °C using only the dominant forms
52 II and III.
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3 The absence of extinction leads to sixteen possible space groups for the hexagonal (or trigonal)
4 form-II. It was proposed¹³ for the equivalent K_2HfF_6 form ($a = 6.40$, $c = 3.81$ Å, "*sometimes*
5 *additional lines occur*" in the 170-220 °C range), but not confirmed by any refinement, that it could
6 be isostructural with the β_1 - K_2MF_6 ($M = U, Th$) investigated by Zachariasen²³, with the space
7 group $P\bar{6}2m$. Zr would have an unusual ninefold coordination according to that model.
8 Unfortunately, no satisfying Rietveld fit could be obtained by following this approach. No more
9 success was obtained by trying other models selected among the A_2BX_6 known structures with
10 relatively similar cell parameters, including the K_2GeF_6 structure-type³³.

11
12 The cubic form-III could be best refined ($R_B = 4.14$ %) by applying the K_3ZrF_7 model³¹ to the
13 495 °C synchrotron pattern (refined together with form-VII). Crystallographic parameters are
14 gathered in Table S1 and interatomic distances in Table S2. The same model used in the attempts
15 to fit the 264 °C complex pattern could lead to a Rietveld Bragg reliability $R_B \sim 14$ % while nothing
16 better than $R_B \sim 25$ % was attained for form-II. It is believed that form-II belongs to the β_1 - K_2MF_6
17 ($M = U, Th$) structure type but that disorder exists and could not be characterized due to the mixture
18 of phases. Indeed, the K_3ZrF_7 compound presents a strong disorder of the F atom positions so that
19 it is difficult to estimate if all the zirconium polyhedra are ZrF_7 ones or if there is a distribution of
20 three possibilities including also ZrF_6 and ZrF_8 ones. This is not the same for K_2ZrF_6 form-III since
21 all polyhedra around the zirconium have to be octahedra. If only the F1 atoms are considered, they
22 build perfect ZrF_6 octahedra ($Zr-F = 1.977$ Å), but there is a non-negligible proportion of F atoms
23 at the F2 position corresponding to longer Zr-F distances (2.178 Å) compatible with larger
24 coordination which could not be compensated by smaller ones in this case, since the average
25 polyhedra is ZrF_6 that time. It is more probable that all polyhedra are ZrF_6 octahedra adopting five
26 possible positions: that corresponding to F1 or one of the four possibilities suggested by F2.
27 Moreover, there is a supplementary cause for a disorder of the F atoms in K_2ZrF_6 form-III which
28 is the partial occupancy of the potassium sites (fully occupied in K_3ZrF_7). In order to answer to the
29 question "has this phase really a K_2ZrF_6 formula or is it in fact K_3ZrF_7 ?" we fixed the stoichiometry
30 to the latter and the Rietveld R_B value increased from 4.14 to 9.87 %.

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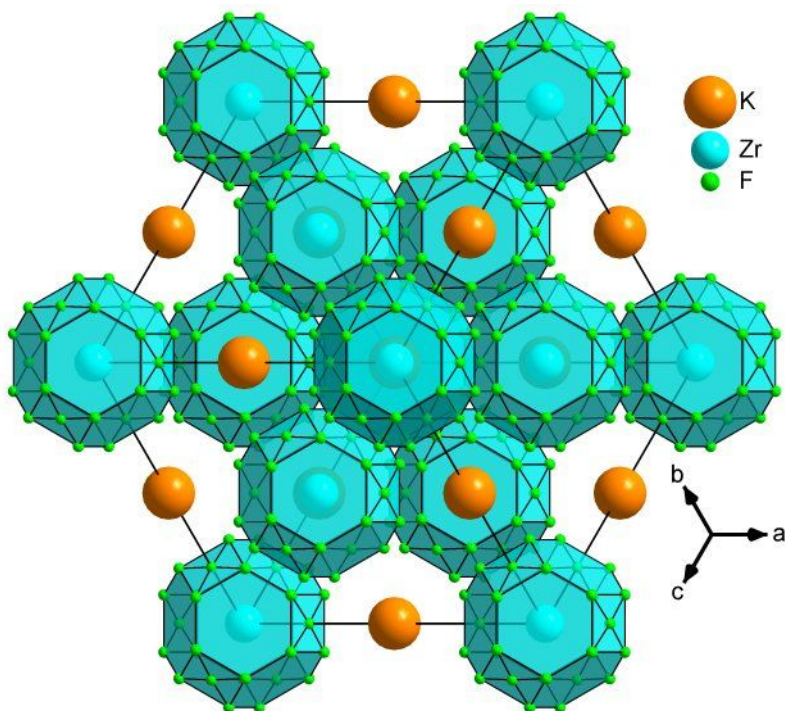


Figure 8. The structure of K_2ZrF_6 form-III projected along the $[111]$ direction, with all possible positions for the F atoms represented, delimiting a kind of isolated sphere around of the Zr atoms due to the various orientations of the ZrF_6 octahedra.

K_2ZrF_6 form-V. This tetragonal form is present in large temperature ranges on heating (240-449 °C) and on cooling (411-139 °C), however, it is never a unique phase and the diffraction peaks are showing strange shapes/splitting, especially at the higher temperatures (Figure 9). It is to be noted that the a cell parameter decreases when the temperature increases, whereas c increases in a normal thermal expansion behaviour.

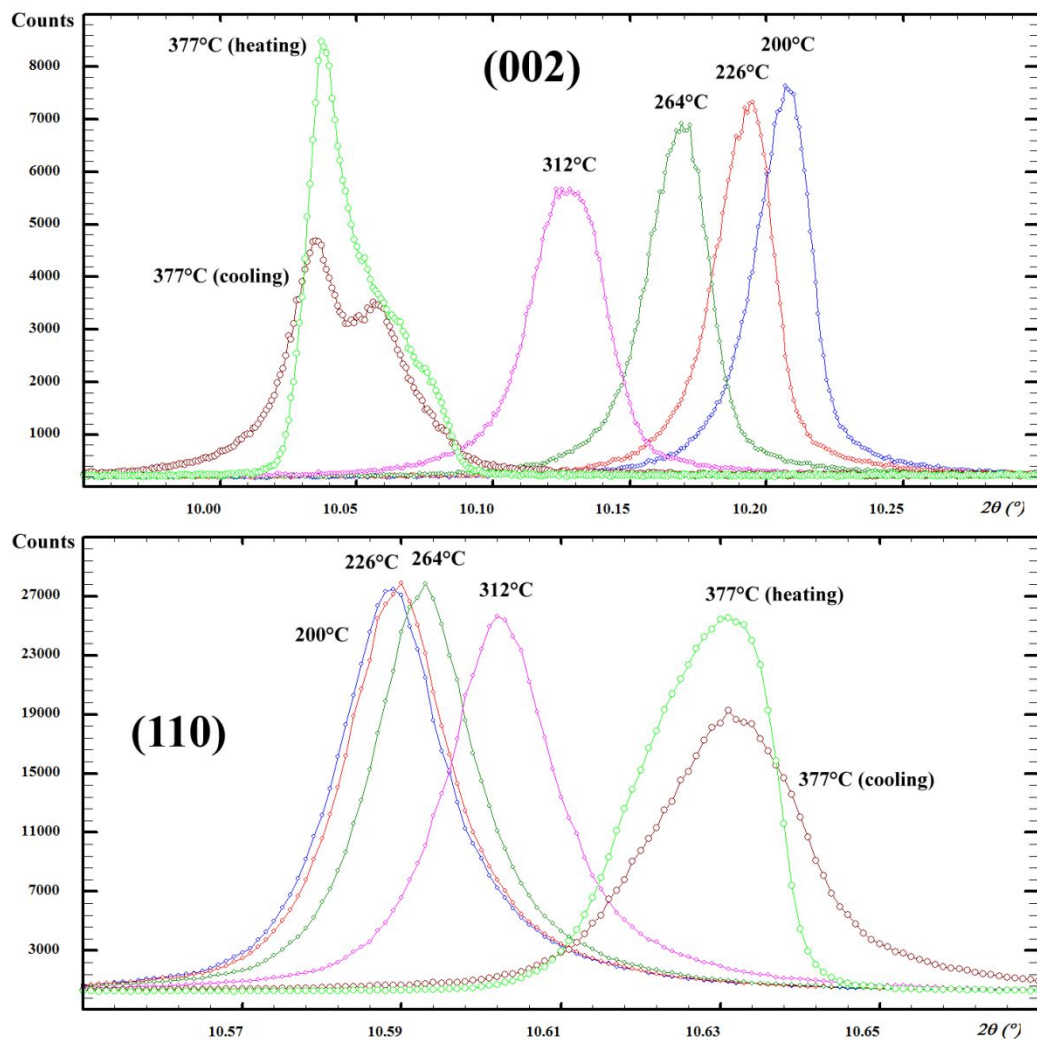


Figure 9. Splitting of the (002) reflection (K_2ZrF_6 form-V) at the higher temperatures, strange peak shapes, a cell parameter decreasing with the increasing temperature as seen from the (110) reflection positions. Where not specified, temperatures were on cooling from 495 °C.

The best pattern selected for the structure refinement is that of sample 2 at 200 °C on cooling. The space group $P4/mmm$ proposed (no refinement) for the tetragonal form of K_2HfF_6 could not fit for K_2ZrF_6 , the powder pattern showing clearly systematic extinctions leading to the $P4/nmm$ space group. After structure solution, the refinement could be managed with an estimation of the anisotropic thermal parameters. The anisotropic peak broadening was undertaken by a simple approach, separating the narrower reflections (hh0) from the others. The crystallographic parameters and interatomic distances are in Tables S3 and S4, respectively. The Rietveld plot is in Figure 10.

The crystal structure (Figure 11) of form-V is built up from ZrF_6 octahedra, and $K(1)F_6$ octahedra (as in KF) alternating in corner-sharing chains with the ZrF_6 ones (Figure 12). In these $K(1)F_6$ octahedra, the potassium atom is strongly off-centre. The second independent potassium atom $K(2)$ is in a 8+4 coordination very close to a cuboctahedron (the square faces are rectangularly distorted).

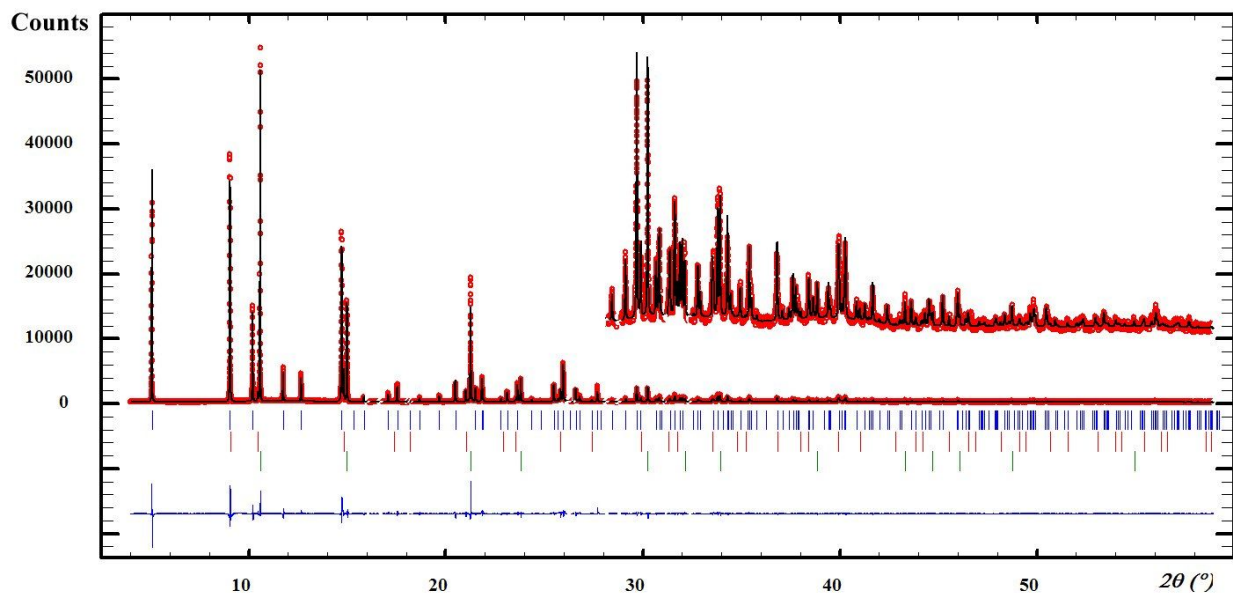


Figure 10. Refined diffraction pattern from synchrotron data for K_2ZrF_6 form-V at 200 °C on cooling. Red dots represent the observed data, the black line represents the calculated ones. Bragg ticks are the peak positions (3 sets from bottom to up: $hk0$ reflections of form-V, cubic K_2ZrF_6 form-III, other peaks of form-V) and the blue curve shows the difference between the observed and calculated patterns, $\lambda = 0.82695 \text{ \AA}$. Traces of $m\text{-}ZrO_2$ were managed by small excluded zones.

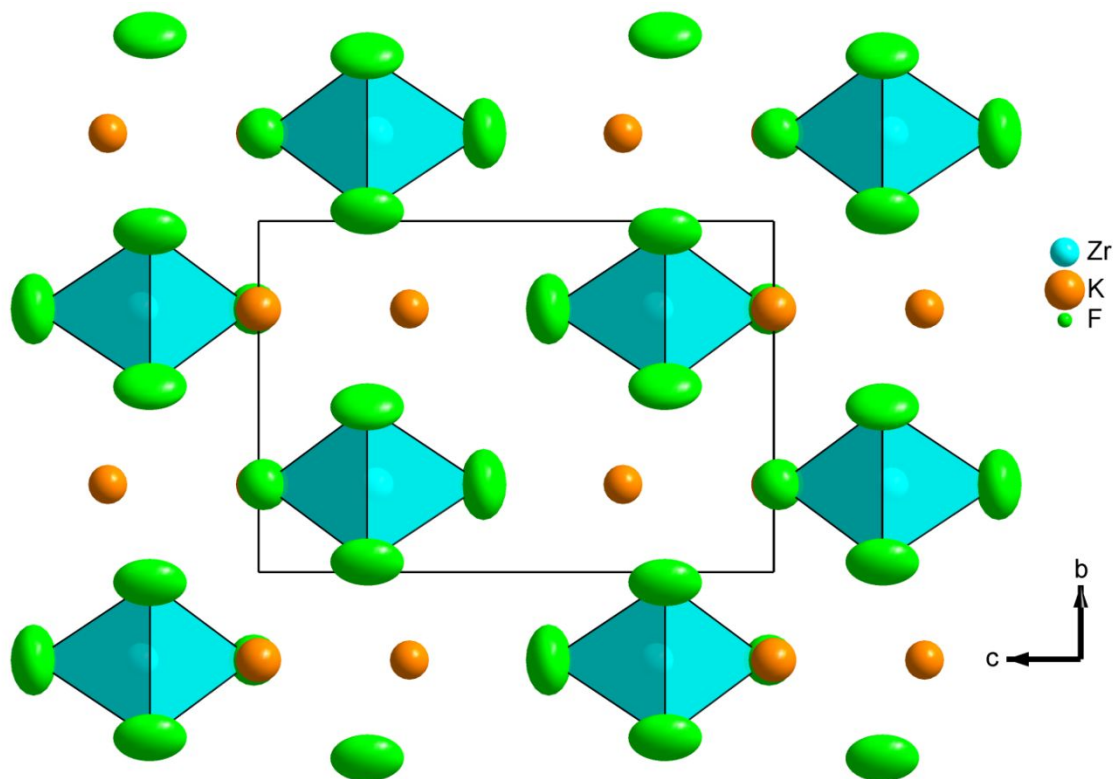


Figure 11. Unit cell projection of the K_2ZrF_6 form-V (at 200 °C on cooling) structure along the a axis. Ellipsoids are represented at 50% probability.

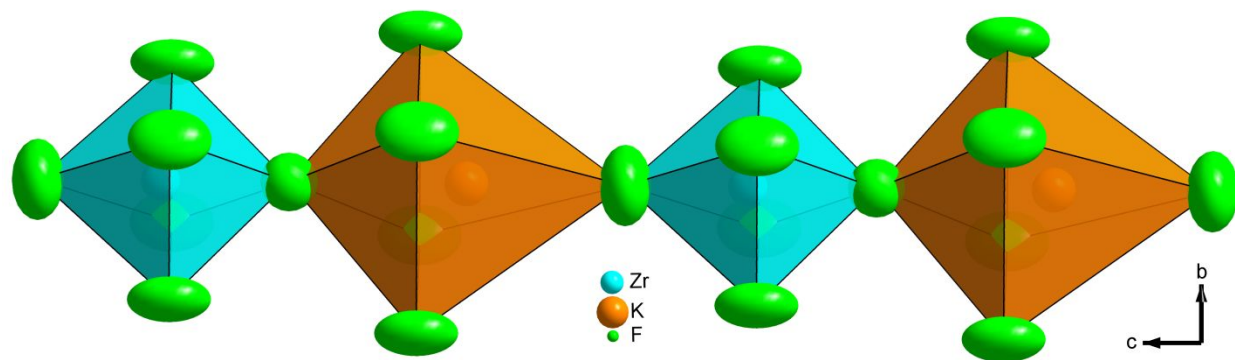
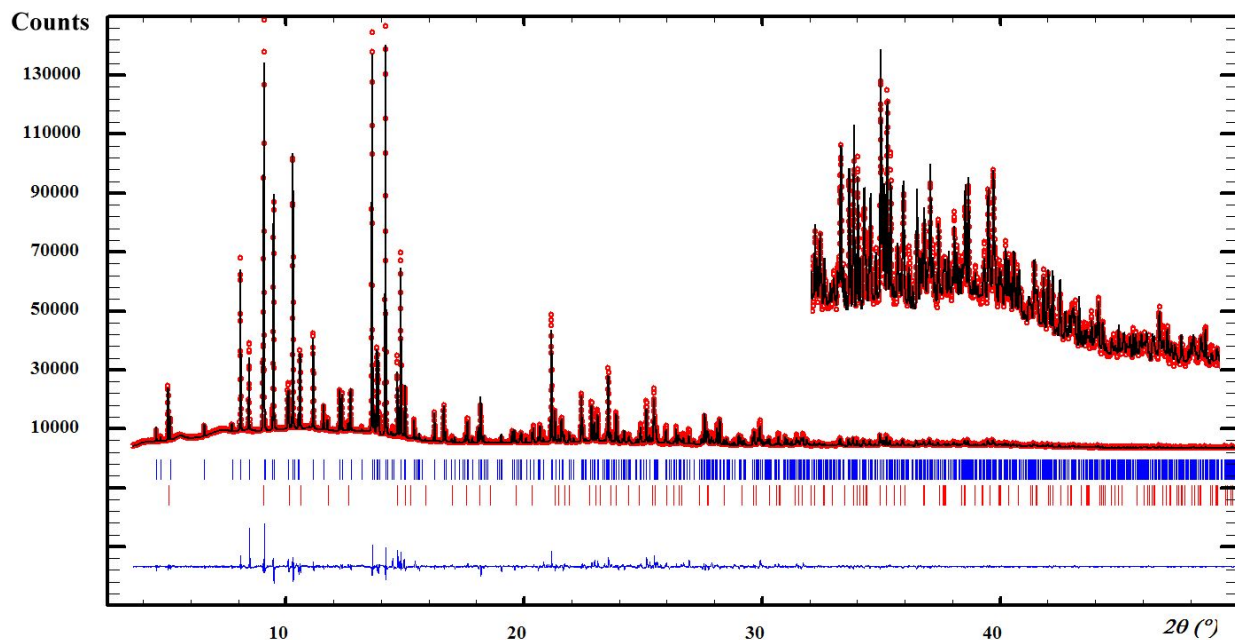


Figure 12. Alternating corner-sharing trans-chains of ZrF_6 and KF_6 octahedra along the c axis of the K_2ZrF_6 form-V structure. Ellipsoids are represented at 50% probability.

K_2ZrF_6 form-VI. This orthorhombic form (space group $Cmcm$) has a small domain of existence in temperature (288-317 °C) and is mixed with form-V and form-III. The unindexed form-IV continues also to exist but its proportion decreases with the temperature increasing and the peaks become even broader than previously. A relatively fast cooling was made in situ during the synchrotron measurements (from sample 1) from 300 to 54 °C, and the forms VI and V were also

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3 observed quenched at that temperature. The refined crystallographic parameters and interatomic
4 distances are in Tables S5 and S6, respectively, and the Rietveld plots are in Figure 13 (295 °C)
5 and 14 (54°C).
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31 **Figure 13.** Refined diffraction pattern from synchrotron data for K_2ZrF_6 form-VI at 295 °C. Red dots represent the
32 observed data, the black line represents the calculated ones. Bragg ticks are the peak positions (up: form-VI, bottom:
33 form V) and the blue curve shows the difference between the observed and calculated patterns, $\lambda = 0.82711 \text{ \AA}$.
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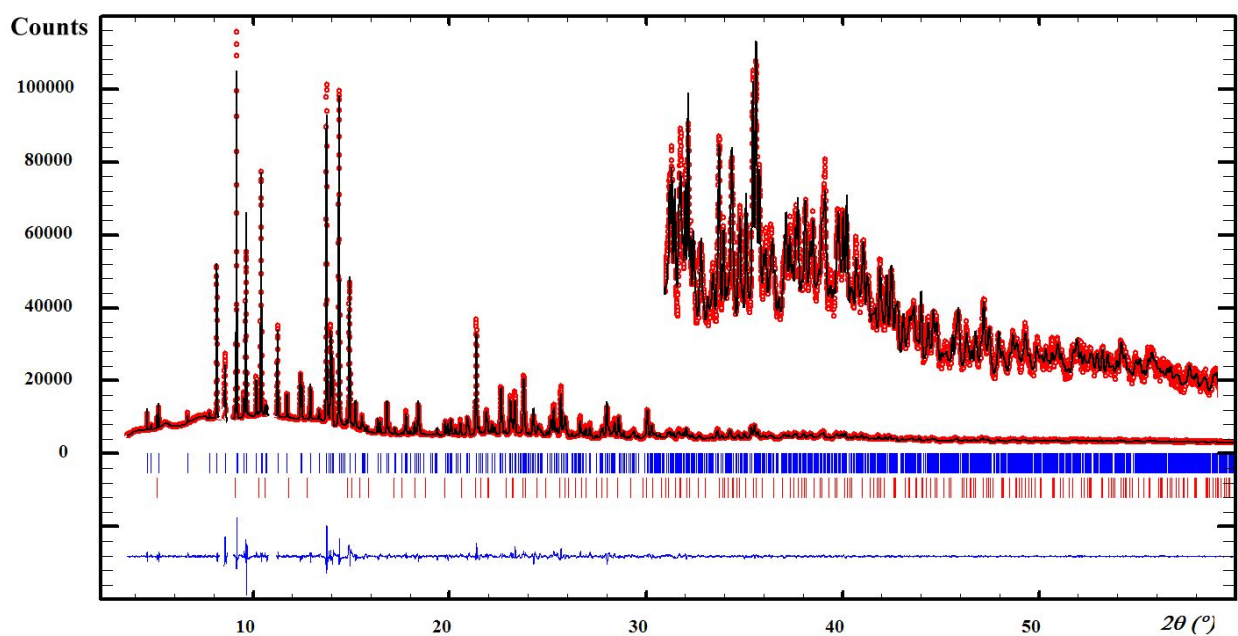


Figure 14. Refined diffraction pattern from synchrotron data for K_2ZrF_6 form-VI at 54 °C after natural cooling from 300 °C. Red dots represent the observed data, the black line represents the calculated ones. Bragg ticks are the peak positions (up: form-VI, bottom : form V) and the blue curve shows the difference between the observed and calculated patterns, $\lambda = 0.82711$ Å.

As for form-V, K_2ZrF_6 form-VI contains one isolated $Zr(1)F_6$ octahedra but there is also an uncommon dimeric entity formulated Zr_2F_{12} built from a $Zr(2)F_7$ monocapped trigonal prism sharing a corner with a $Zr(3)F_6$ octahedron (Figure 15). There are five different potassium atom sites, showing all a different polyhedral shape: $K(1)F_8$ is in a bicapped trigonal prism; $K(2)F_{11}$ is a tricapped cube; $K(3)F_7$ is a monocapped trigonal prism; $K(4)F_6$ is an octahedron; $K(5)F_8$ is a hexagonal-based bipyramid. There is a considerable decrease of the b and c axes lengths between 295 and 54 °C. The atom which had a quite abnormally big thermal factor at 295 °C was F(9), but it is F(7) at 54 °C. Both fluorine atoms belong to the $Zr(3)$ environment and show abnormally short Zr-F distances which could be released according to some positional disorders presenting differences at the two temperatures.

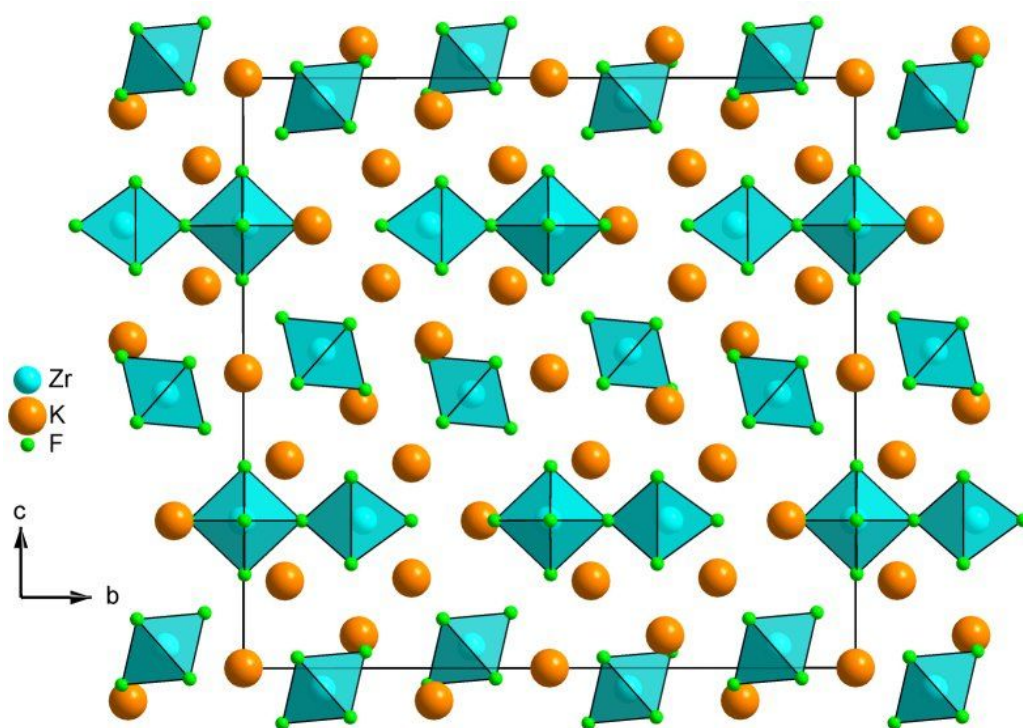


Figure 15. Unit cell projection of the K_2ZrF_6 form-VI (at 295 °C) structure along the short a axis showing the isolated ZrF_6 octahedra and the dimeric entities formulated Zr_2F_{12} built from a ZrF_7 monocapped trigonal prism sharing a corner with a ZrF_6 octahedron.

K₂ZrF₆ form-VII. The pattern selected for structure solution and refinement is at 495 °C. Some ZrO₂ is appearing due to the high reactivity of fluorides and the contact with the sapphire capillary. Two ZrO₂ forms are present, the monoclinic and the tetragonal. The cubic form-III is present in large quantity so that its structure could be refined simultaneously as noted previously. The form-VII refined crystallographic parameters and selected interatomic distances are in Tables S7 and S8, respectively, the Rietveld plot is in Figure 16.

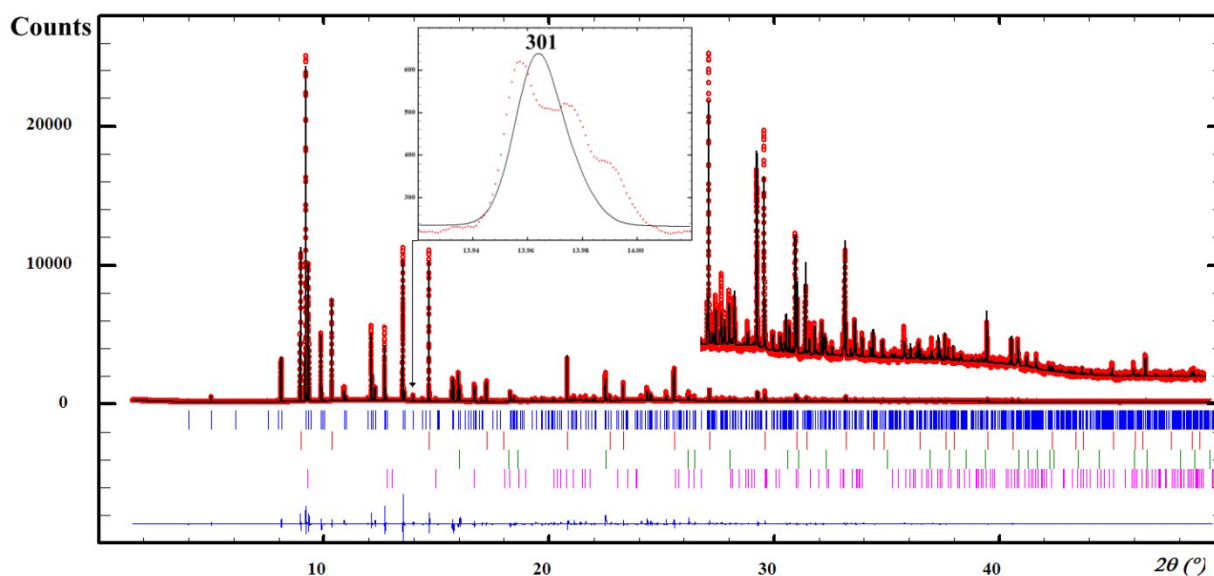


Figure 16. Refined diffraction pattern from synchrotron data for K₂ZrF₆ form-VII at 495 °C. Red dots represent the observed data, the black line represents the calculated ones. Bragg ticks are the peak positions (from up to down: orthorhombic form-VII, cubic form-III, tetragonal ZrO₂, monoclinic ZrO₂) and the blue curve shows the difference between the observed and calculated patterns, $\lambda = 0.82695 \text{ \AA}$. The splitting of some form-VII peaks is exemplified by the enlarged part showing the tripled 301 peak.

There are considerable differences between the DFT-optimized atomic coordinates and the Rietveld-refined ones and therefore on the interatomic distances (Tables S7 and S8). The indexing and the space group assignment were straightforward from the first 20 peaks at small diffraction angles, and the Le Bail fit allowed to extract intensities that led very directly to the full structural model by the direct space method. However, when the final Rietveld fit is examined in details, many peaks are badly fitted due to splitting (see especially the 301 peak split in three components in the Figure 16). There is no splitting at all for form-III peaks. Even if the ZrF₆ octahedra were

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3 recognized at the solution stage, soft restraints had to be used on the Zr-F distances during the
4 refinement, in spite of which some distances are too short in Table S8. We have obviously to
5 consider this structure as an imperfect approach, and even if the DFT optimization is capable to
6 provide more realistic interatomic distances (to a point that we have chosen this time to show the
7 drawing of the optimized crystal structure), using the fixed DFT coordinates does not improve the
8 Rietveld fit, indeed, it is worst. Anyway, this phenomenon of peak splitting was also seen for form-
9 V especially at the vicinity of the reversible V \leftrightarrow VII transition, and also for some of the form-I
10 various preparations. From the comparison of the crystal structures of forms VII and V (Figure
11 17), the V \leftrightarrow VII transition is more probably topotactic, given the cell parameters obvious relations
12 $a_{\text{VII}} \sim c_{\text{V}}$, $b_{\text{VII}} \sim b_{\text{V}}$, $c_{\text{VII}} \sim 4 a_{\text{V}}$ and the observation that relatively small translations and rotations
13 of the ZrF₆ octahedra and translations of the K atoms allow to reproduce one crystal structure from
14 the other.
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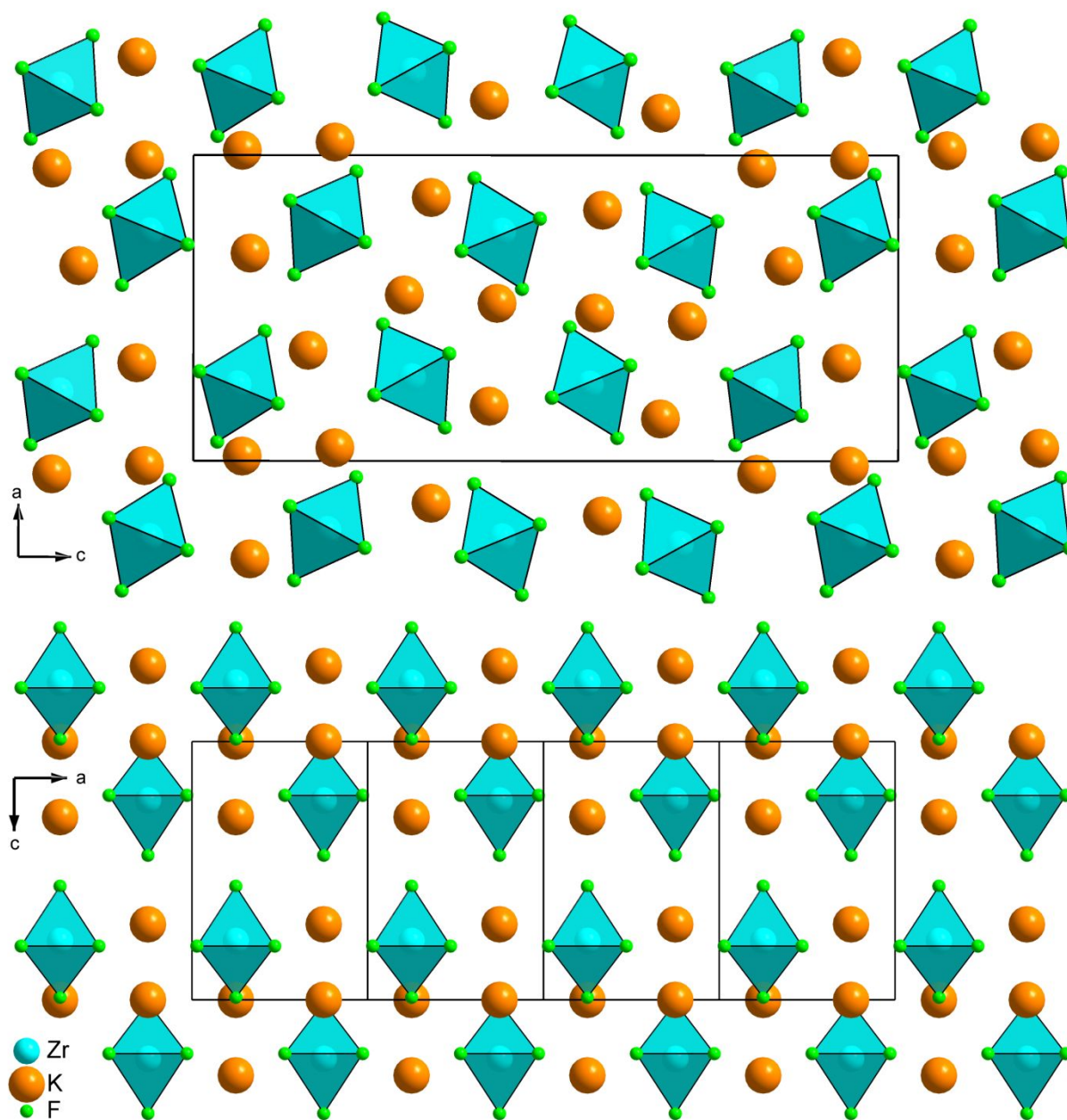


Figure 17. K_2ZrF_6 form-VII (up) DFT-optimized model compared to form-V (down) extended on four times the a_V cell parameter. The comparison shows the rotations and translations of ZrF_6 octahedra as well as the translations of the K atoms occurring during the topotactic transition.

However, such a non-reconstructive transition suggests that a strong disorder may affect both phases since either $4 a_V$ or $4 b_V$ can become c_{VII} , this could explain the quasi impossibility to fit the form-V powder patterns at high temperature, so that what we call "micro-phases" could correspond to different local arrangements at the interface of domains having the main structure.

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3 It is hard to give the polyhedra shapes around of the potassium atoms, if the limit in the K-F
4 maximum distance is fixed at 3.5 Å, the polyhedra shapes may be slightly different in the DFT-
5 optimized and Rietveld-refined models.
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8 Based on the XRD results the scheme a scheme of phase transformations of K_2ZrF_6 is suggested
9 on Figure 18. In order to support the suggested models high temperature ^{19}F MAS NMR
10 spectroscopy was applied. As was mentioned in our previous work ⁹, spectrum at 250 °C contains
11 two resonances at -16.3 ppm and at -33.7 ppm with ratio 3:1, respectively. It should be noted that
12 the signals have very different values of the chemical shift anisotropy, 28 and 10 ppm, respectively.
13 An increase of the temperature from 250 °C to 290 °C leads to a decrease of the line width (full
14 width at half-maximum, FWHM) for low field peak from 740 to 460 Hz and increase for high field
15 signal from 440 Hz to 640 Hz. In addition, a small shift of around 1 ppm in low field was measured.
16 At 310 °C two new peaks appear at -15.9 and -19.1 ppm. In the range of the temperature 310-340
17 °C we observe 4 signals. We note that at 340 °C two initially observed peaks almost disappeared.
18 At 360 °C the integral intensity of the resonance at -15.5 ppm reaches maximum and then it starts
19 to decreases at 380 °C until the signal disappears completely at 410 °C. At 410 °C only one peak
20 was observed. NMR data are summarized in Table 3.
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31 Usually, at room temperature (or at lower temperatures) the high-resolution solid state NMR
32 spectrum should contain the number of lines corresponding to the number of crystallographically
33 non-equivalent atoms depending on the local chemical environment, in other words different
34 nucleus in a molecule resonates at different frequencies. On other hand, the NMR spectra acquired
35 in molten state consist in a single Lorentzian line, characteristic of a rapid exchange at the
36 timescale of NMR between the different atomic configurations around the observed nucleus ³⁴. In
37 some cases the measured chemical shift might thus be the average of the individual chemical shifts
38 of the different species even at room temperature or temperatures below melting as was discussed
39 many times for K_2TaF_7 ³⁵ and refs. therein. Some other examples of the dynamics of fluorine atoms
40 can be Ba_5AlF_{13} ³⁶, Rb_3TaOF_6 ³⁷, or $(NH_4)_2ZrF_6$ ³⁸.
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48 In the present case, we observed a number of fluorine resonances much smaller than the number
49 of non-equivalent fluorines. We explain this discrepancy by the dynamic behavior of fluorine
50 atoms, which consists of a fast hopping of the fluorine on each position around one polyhedron. We
51 think that in the measured spectra of K_2ZrF_6 we observe a dynamic behavior of fluorine atoms,
52 each signal corresponds to a different polyhedron and the chemical shift is the barycenter of the
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3 individual resonances. Similar behavior was observed also in the case of $K_{18}Ta_5Zr_5F_{63}$ compound
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7 In the range 250 °C to 290 °C we detected two signals for K_2ZrF_6 . The value of the chemical shift
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9 at -33.7 ppm is slightly different from others, and it is very close to the signal of K_3ZrF_7 (-34/-37
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11 ppm at RT^{9, 12}), and we assigned this signal to the phase-III related to K_3ZrF_7 . The second signal
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13 can be assigned to β - K_2ZrF_6 . At 410 °C NMR spectrum is characterized by one line at -18.2 ppm
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15 and it can be attributed to the phase-VII with ZrF_6 environments (Figure 19). The phase-VI consists
16
17 of two different polyhedrons: ZrF_6 and Zr_2F_{12} . The signal at -15.4 ppm was assigned to dimeric
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19 fragment Zr_2F_{12} and signal around -18.4 ppm to the island like ZrF_6 fragment. The phase transition
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21 between 390 °C and 410 °C is accompanied not by the appearance of a new signal, but by the
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23 disappearance of the signal at -15.3 ppm corresponding to the dimer Zr_2F_{12} and preserving the
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25 signal at -18.3 ppm. Based on this observations it could be estimated that β -phase that was for the
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27 first time mentioned by³ and lastly discussed by⁹ probably consist of structural motives based on
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29 bridging of zirconium atoms by fluorine atoms; Zr-F-Zr.

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31 The spectral changes were not reversible upon cooling to RT. A cooling up to 170 °C does not
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33 lead to changes in the spectrum, only one line was observed, which just shifted from -18.2 ppm
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35 (410 °C) to -20 ppm (170 °C). Then two new small peaks were observed. In order to reach the
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37 maximum resolution, a very fast spinning rate of 50 kHz was used after cooling the sample. As in
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39 case the heating-cooling cycle RT- 250 °C-RT⁹, in addition to the expected signals for the α -phase
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41 (I=97.3 %), two small signals at -34.6 ppm (I=1.6 %) and -36.7 ppm (I=0.1 %) were detected. This
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43 spectral evolution during the cooling process is in a good agreement with the obtained XRD data.
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Table 3. ^{19}F NMR data obtained from the simulation of the ^{19}F NMR spectra at 17.6 T and MAS 7 kHz during the heating of the sample from RT up to 410 °C

T, °C	Zr_2F_{12}		ZrF_6		$\beta\text{-K}_2\text{ZrF}_6$		phase-III (K_3ZrF_7)		
	δ_{iso} , ppm (± 0.1 ppm)	Integral, % ($\pm 1\%$)	δ_{iso} , ppm (± 0.1 ppm)	Integral, % ($\pm 1\%$)	δ_{iso} , ppm (± 0.1 ppm)	Integral, % ($\pm 1\%$)	δ_{iso} , ppm (± 0.1 ppm)	Integral, % ($\pm 1\%$)	
410	-	-	-18.2	100	-	-	-	-	phase-VII
390	-15.3	16.8	-18.3	83.2	-	-	-	-	
380	-15.4	39	-18.4	61	-	-	-	-	
380	-15.4	52.9	-18.4	47.1	-	-	-	-	phase-VI
360	-15.5	75	-18.8	25	-	-	-	-	
340	-15.6	75	-18.8	23	-13.3	1	-32.9	1	phase-VI;
320	-15.8	70	-19	24	-14	4	-33.1	2	β -phase;
310	-15.9	7	-19.1	30	-14.7	51	-33.2	12	phase-III
290	-	-	-	-	-15.1	80	-33.2	20	phase-II +
		-		-					Phase-IV =
									β -phase;
250	-		-		-16.3	77	-33.7	23	phase-III

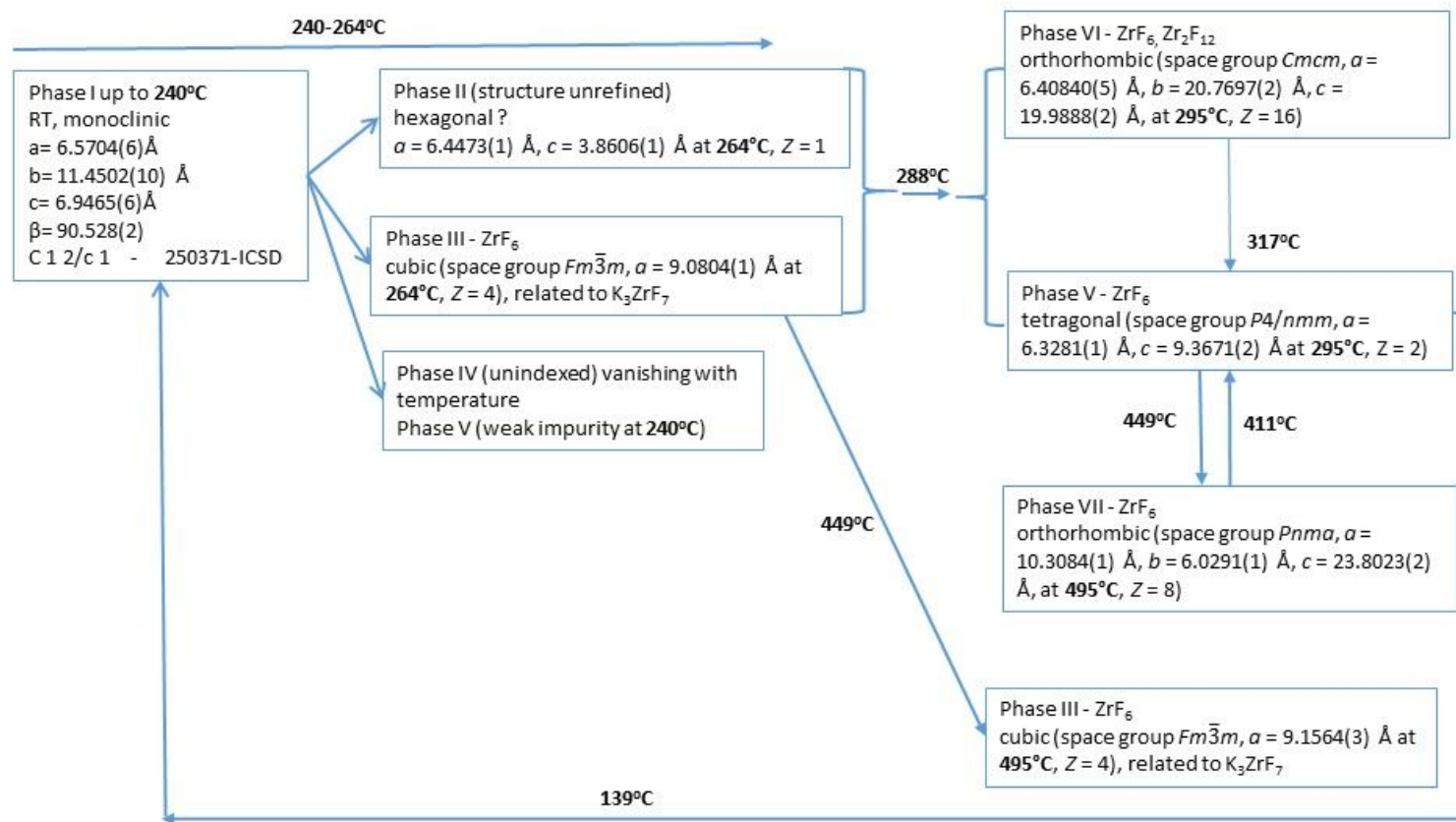


Figure 18. Scheme of phase transformations of K_2ZrF_6 .

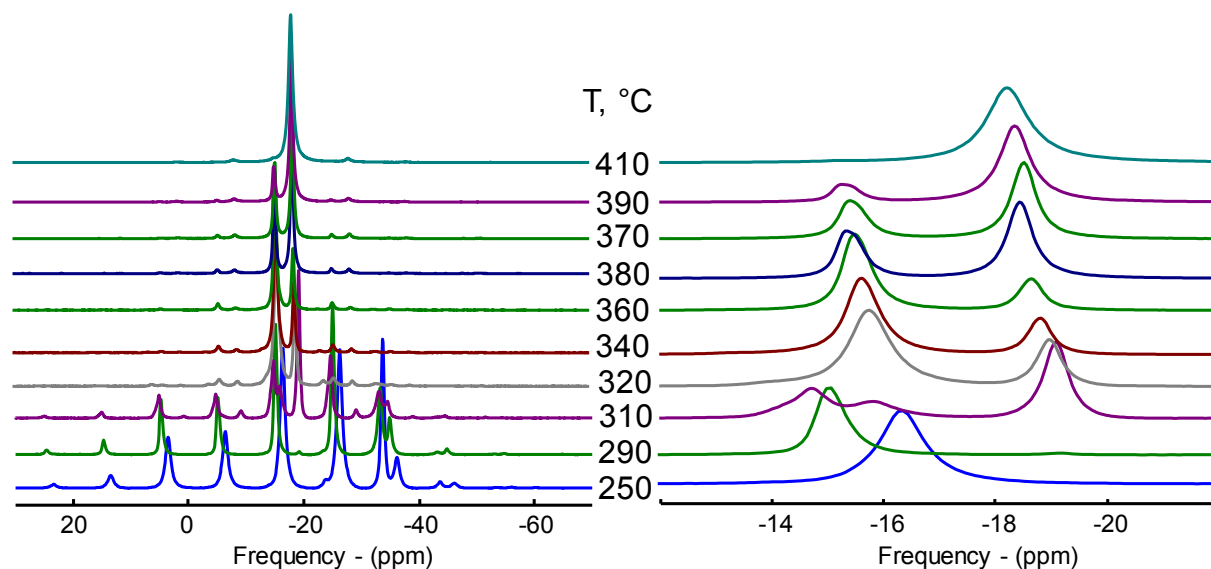


Figure 19. In the left ^{19}F NMR spectra of K_2ZrF_6 acquired at 17.6 T and MAS 7 kHz as a function of temperature. In the right extended view of the region from 12 ppm to 22 ppm.

CONCLUSIONS

From this study, not everything is explained yet (forms II and IV, more details on hypothetical micro-phases) but a considerable step is made in the knowledge of the K_2ZrF_6 complex polymorphism that reveals to be seriously different from the old expectations. An incredible diversity of polyhedral shapes is adopted by the zirconium and the potassium atoms, with coordinations from 6 to 7 and 6 to 12, respectively. The " $\beta\text{-K}_2\text{ZrF}_6$ " phase characterized previously by hyperfine quadrupolar interaction of ^{181}Ta at Zr sites ⁴⁰, or by RAMAN ⁴¹ and NMR ⁷ before any crystallography study, is a mixture of up to four phases, indeed, two of which being still not completely understood (forms-II and IV). Three new crystal structures types were established *ab initio* from powder diffraction data for forms-V, VI and VII. Rietveld refinements were concluding for the ubiquitous cubic form-III related to K_3ZrF_7 suggesting a possible conduction by K^+ ions as a consequence of 2/3 sites occupied. Strange enough is the fact that the forms VI and VII at high temperature present both a lower symmetry (orthorhombic with $Z = 16$ and 8, respectively) than the tetragonal form-V ($Z = 2$) existing also at much lower temperatures. A scheme of phase transformations of K_2ZrF_6 was suggested. It could be noted that the phase transformation of K_2ZrF_6 (here all Zr atoms are eight coordinated at RT forming ZrF_8) is related with the breaking of infinite

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3 chains of bridged zirconia atoms through F atoms followed by the formation of phase with dimeric
4 fragments finished with island like fragment of ZrF_6 .

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6 A re-examination of the polymorphism of K_2HfF_6 looks desirable, given the high degree of
7 isomorphism of Zr- and Hf-based fluorides. The fact is that no K_2HfF_6 powder patterns
8 corresponding to our K_2ZrF_6 complex forms VI and VII (nor IV) was observed, though the forms
9 I, II, III and V were listed¹³ together with some others which we could not disclose. In case of
10 future use of the K_2ZrF_6 formulation for the “Generation IV” Molten Salts Nuclear Reactors, these
11 results will have to be taken into consideration, especially the strong variation of the volume
12 occupied by the molecular unit in the various solid states when cooling from the molten state. The
13 nice dream of a phase transition from a crystal structure to another, both perfectly ordered, is not
14 fulfilled in the K_2ZrF_6 case, for reasons remaining to be explained.
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24 **ASSOCIATED CONTENT**

25 Supplementary material contain crystallographic parameters and selected bond distances or
26 K_2ZrF_6 forms III, V, VI and VII phases tables of indexed powder patterns of phases I-VII, full
27 dataset for K_2ZrF_6 at 265°C and a CIF file for every synchrotron refinement including the Fullprof
28 .pcr and .dat files for Rietveld or Le Bail fits. This material is available free of charge via the
29 internet at <http://pubs.acs.org>.
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36 **CORRESPONDING AUTHOR**

37 *Institute of Inorganic Chemistry, Department of molten systems, Dubravská cesta 9, Bratislava;
38 845 36, Slovakia. E-mail: miroslav.boca@savba.sk.
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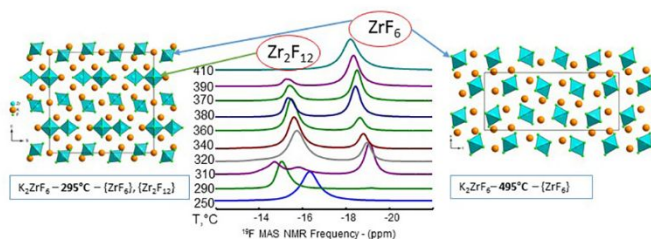
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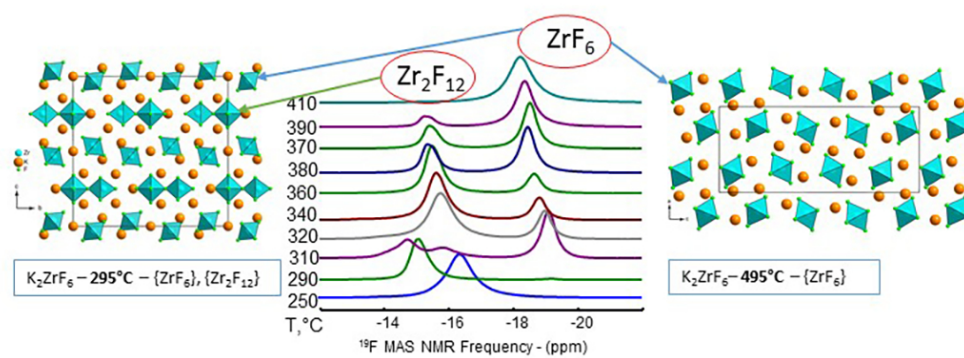
Polymorphism of K_2ZrF_6

Lubomír Smrčo[†], Armel Le Bail, Miroslav Boca, Aydar Rakhmatullin*



SYNOPSIS

Phase transformations of K_2ZrF_6 were analysed *in situ* using synchrotron powder diffraction and high temperature ^{19}F MAS NMR. Exceptional multiple solid-solid transformation was described.



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