Strontium Phosphate Composite Designed to Red-Emission at Different Temperatures

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Abstract: Eu³⁺-doped Sr₁₀(PO₄)₆(OH)₂–Sr₆(PO₄)₂ (SrHAp-TSP) composites were obtained via the microwave-stimulated hydrothermal method and post-heat-treated from 750 to 950 °C. Concentration of the Eu³⁺ ions was set to be 0.5, 1, 2, 3, 5 mol% in a ratio of the strontium ions molar content. The structural and morphological properties were investigated by X-ray powder diffraction (XRPD), scanning electron microscopy (SEM) and fourier transform infrared spectroscopy (FT-IR) techniques. The average particle size of the studied materials annealed at 750, 850 and 950 °C were counted about 100, 131 and 173 nm, respectively. The luminescence properties depending on the dopant ion concentration, heat-treatment temperature, excitation wavelength and temperature were investigated. In the emission spectra, a broad peak corresponding to the 4f⁵d⁷ → 4F (⁷S₂) emission of Eu³⁺ ions as well as narrow 4f-4f transitions typical for Eu³⁺ ions can be observed. The luminescence intensity of the 1 mol% Eu³⁺:Sr₁₀(PO₄)₆(OH)₂–Sr₆(PO₄)₂ was measured depending on the ambient temperature in the range of 80–550 K. The CIE 1931 (International Commission on Illumination) chromaticity diagram was determined from emission spectra measured in 80, 300 and 550 K. The reduction mechanism of the Eu³⁺ to the Eu²⁺ was explained by the charge compensation mechanism based on the Kröger–Vink notation. The decay times were measured and the Judd–Ofelt (J–O) theory was applied to analyze the observed structural and spectroscopic features.

Keywords: luminescence; strontium hydroxyapatite; strontium phosphate; europium (III) ions; biomaterial; hydrothermal synthesis; Kröger–Vink notation

1. Introduction

Apatite (Ap) is a large family of compounds commonly occurring in nature. Calcium hydroxyapatite (Ca₁₀(PO₄)₆(OH)₂ – CaHAp) is ideal representative of this family of minerals, which can be easily doped with different kinds of ions [1,2]. Apatite is applied in plenty industrial fields as catalysts, laser hosts, luminescent materials, gas sensors, ionic conductors and adsorbents [3,4]. Moreover, calcium phosphates are also used in implantology due to high similarity to the inorganic part of hard tissues in bones, high biocompatibility, osteoconductivity, nontoxicity, non-immunogenicity and bioactivity [5–7]. Apatite doped with lanthanide ions are extensively investigated as bio-labels that can be used in in vitro and in vivo imaging [8–10].

Strontium is a trace element present in human bones and tooth impacting on the strength, healing, microarchitecture, and bone formation. Besides, strontium (II) ions were employed in osteoporosis therapy in the form of orally administrated strontium ranelate salt [11]. Moreover, strontium apatites seem to be good candidates as host lattices for luminescent dopants due to their excellent thermal and chemical stability as well as excellent luminescence properties. As an example, the Sr₆(PO₄)₂Cl: Eu²⁺ material was used as the blue component in fluorescent lamp [12].
Designing of phosphor materials activated by lanthanide ions has enabled their applications in such technologies as solid-state lighting, flat panel displays, light-emitting diodes (LEDs), biological diagnostic, and biomedical areas due to their excellent properties [13]. Eu$^{3+}$ ions are the most frequently used activator applied as red-emitting phosphor e.g., in commercially available phosphors Y$_2$O$_5$: Eu$^{2+}$ and Y$_2$O$_3$: Eu$^{3+}$ [14,15]. Moreover, europium can occur in Eu$^{2+}$ ($F$) and Eu$^{3+}$ ($F$) oxidation states of suitable stability. Both show many different features as emitting centers. The Eu$^{3+}$ ion is sensitive to oxidation under some conditions; however, it possesses good stability in the solid-state materials. It is of particular interest because it exhibits strong and tunable emission from blue to red [16]. The emission spectrum consists of the broad band related to the allowed $4f^5d^1 \rightarrow 4f^6 (5S_{7/2})$ transition. Its luminescence properties strongly depend on its environment in the host lattice. The Eu$^{3+}$ ion is also characterized by unique optical properties with a relatively strong red-orange emission. Its luminescence results from forbidden, intra-configurational f-f transitions roughly independent on crystal field influence. However, the intensity of particular transitions are highly sensitive to Eu$^{3+}$ ions surrounding, so additionally, Eu$^{3+}$ ions are very useful as a luminescence probe [17–19].

Luminescence materials co-doped with Eu$^{3+}$ and Eu$^{2+}$ is tough to obtain because the reduction of oxidation state in solid states requires a thermal treatment in reducing atmosphere (H$_2$, H$_2$/N$_2$, CO). However, it is interesting that Eu$^{2+}$-doped materials can be obtained in a nonreducing atmosphere at elevated temperatures in certain compounds due to reduction of Eu$^{3+}$ ions to Eu$^{2+}$ ions by the crystallographic site [20]. This abnormal reduction phenomenon was observed in crystal structures with BO$_4$, SiO$_4$, AlO$_4$, or PO$_4$ tetrahedral groups.

In the present work, for the first time to the best of our knowledge [7,8,21–24], synthesis of strontium phosphate composite (Sr$_{(10)}$(PO$_4$)$_4$(OH)$_2$–Sr$_{(2)}$(PO$_4$)$_3$) co-doped with Eu$^{3+}$ and Eu$^{2+}$ ions prepared in the air at high temperature and the structural, morphological as well as spectroscopic properties have been studied.

2. Materials and Methods

2.1. Synthesis of Composite

Strontium phosphate (Sr$_{(10)}$(PO$_4$)$_4$(OH)$_2$–Sr$_{(2)}$(PO$_4$)$_3$) composite doped with Eu$^{3+}$ ions was synthesised by microwave stimulated hydrothermal method and then post-treated at elevated temperature [7,8]. The Sr(NO$_3$)$_2$ (99.0% min Alfa Aesar, Karlsruhe, Germany), (NH$_4$)$_2$HPO$_4$ (≥ 99.0% Fluka, Bucharest, Romania), Eu$_2$O$_3$ (99.99% Alfa Aesar, Karlsruhe, Germany) and NH$_3$H$_2$O (99% Avantor, Gliwice, Poland) for pH modification were used as substrates. The Eu$^{3+}$ ions concentration was set on 0.5, 1, 2, 3 and 5 mol% in proportion of strontium ions molar content. Firstly, the stoichiometric Eu$_2$O$_3$ mass was digested in the HNO$_3$ (ultrapure Avantor, Gliwice, Poland) to receive Eu(NO$_3$)$_3$·xH$_2$O. Europium nitrate hydrate was re-crystallized three times in order to eliminate the HNO$_3$ excess. Then, the stoichiometric Sr(NO$_3$)$_2$ mass was dissolved in water and the Eu(NO$_3$)$_3$·xH$_2$O also was added in. Thereafter, 0.5353 g (4.05 mmol) of (NH$_3$)$_2$HPO$_4$ was put to the previous mixture and the pH value was modified to 10 by means of ammonia. The suspension was transferred into Teflon vessel (Ertiec, Wroclaw, Poland) and placed in the microwave hydrothermal reactor (MV 02-02, Ertiec, Wroclaw, Poland). The mixture was heat treated at 280 °C for 90 min under 60 atm autogenous pressure. The obtained product was rinsed with de-ionized water (Hydrolab, Straszyn, Poland) several times and dried at 70 °C for 24 h. The thermal treatment from 750 to 950 °C for 3 h was the last step during materials preparation.

2.2. Characterisation

Powder diffraction patterns were received by using an X’Pert Pro diffractometer (PANalytical, Almelo, The Netherlands) equipped with Ni-filtered Cu Kα radiation ($\lambda = 0.154$ nm, $V = 40$ kV, $I = 30$ mA). The structural refinement was done by using a Maud program version 2.93 (University of Trento – Italy, Department of Industrial Engineering, Trento, Italy) [25,26] based on the apatite hexagonal and trigonal tristontium diphosphate crystal structures with better approximation and indexing of the Crystallographic Information File (CIF). The quality of structural refinement was
supervised by R-values ($R_w$, $R_{wp}$, $R_{all}$, $R_p$, and $q$), which were followed to get a structural refinement with better quality and reliability.

A FEI Nova NanoSEM 230 scanning electron microscope (Hillsboro, OR, USA) equipped with energy dispersive spectrometer spectrometer (EDS; EDAX Genesis XM4; Hillsboro, OR, USA) and operating at an acceleration voltage in the range 3.0–15.0 kV and spot 4.0–4.5 was used to determine the surface morphology and the elements mapping. The EDS analysis was done to confirm the contents of elements in the obtained materials.

A Nicolet iS50 FT-IR (Thermo Scientific, Waltham, MA, USA) spectrometer equipped with an Automated Beamsplitter exchange system (iS50 ABX containing DLaTGS KBr detector) (Thermo Scientific, Waltham, MA, USA) and HeNe laser as an IR radiation source were used to measure IR spectra. Polycrystalline mid-IR spectra were collected in the 4000–400 cm$^{-1}$ range in KBr pellets at the temperature of 295 K and spectral resolution of 4 cm$^{-1}$.

The excitation spectra were recorded using an FLS980 Fluorescence Spectrometer (Edinburgh Instruments, Kirkton Campus, UK) equipped with 450 W Xenon lamp. The excitation of 300 mm focal length monochromator was in Czerny–Turner configuration. The excitation arm was supplied with holographic grating of 1800 lines/mm grating blazed at 250 nm. The excitation spectra were corrected to the excitation source intensity. The emission spectra were measured with a PMA-12 photonic multichannel analyser (Hamamatsu, Hamamatsu City, Japan) equipped with back-thinned charge-coupled device (BT-CCD) line (Hamamatsu, Hamamatsu City, Japan). As an excitation source a pulsed 266 nm line of Nd: YAG laser (3rd harmonic) and a pulsed 395 nm line from Ti: Sapphire tunable laser pumped by 532 nm line from Nd:YAG laser was used (LOTIS TII, Minsk, Belarus). The detection setup is calibrated and has flat response in whole working range (350–1100 nm). The measurements were carried out in 300 K and 80 K as well as in the temperature range 80–550 K using Linkam THMS600 cooling/heating stage (Linkam, Epsom, UK).

The luminescence kinetics were measured by using Jobin-Yvon THR1000 monochromator (HORIBA Jobin-Yvon, Palaiseau, France) equipped with Hamamatsu R928 photomultiplier (Hamamatsu, Hamamatsu City, Japan) and 1200 lines/mm grating blazed at 500 nm. As an excitation source a pulsed 395 nm line from Ti: Sapphire tunable laser (LOTIS TII, Minsk, Belarus) pumped by 532 nm line from Nd: YAG laser was used. The decay profiles were collected using a LeCroy WaveSurfer 400 MHz oscilloscope (Teledyne LeCroy, Chestnut Ridge, NY, USA). The luminescence kinetics were monitored at 612 nm line which is the maximum of the most intense electric dipole transition ($^2$D$_{0} \rightarrow ^2F_{2}$) and the effective emission lifetimes were calculated by the following equation:

$$\tau_{m} = \frac{\int_{0}^{\infty} t I(t) dt}{\int_{0}^{\infty} I(t) dt} = \frac{\int_{0}^{t_{\text{max}}} t I(t) dt}{\int_{0}^{t_{\text{max}}} I(t) dt}$$

(1)

where $I(t)$ is the luminescence intensity at time $t$ corrected for the background, and the integrals are calculated over the range of $0 < t < t_{\text{max}}$, where $t_{\text{max}} >> \tau_{m}$.

A blue luminescence decay curve was excited by a femtosecond laser system Libra Series (Coherent, Wilsonville, USA; 1 mJ, 89 fs, 800 nm) coupled to a Light Conversion OPerA-Solo Optical Parametric Amplifier (Coherent, Wilsonville, USA) and a C5680 streak camera (Hamamatsu, Hamamatsu City, Japan) was used as a detector. The decay curve fitting was performed via Origin 2019 software (OriginLab, Northampton, MA, USA) by using following single exponential decay formula:

$$I(t) = I_0 + A \exp(-t/\tau)$$

(2)

where $I(t)$ and $I_0$ are the luminescence intensity at time $t$ and $t >> \tau$, $A$ is a constant, and $\tau$ is the decay time for an exponential component.

3. Results and Discussion

3.1. Structural and Morphological Analysis

The formation of the strontium phosphate composite ($\text{Sr}_{10}$(PO$_4$)$_3$(OH)$_2$–Sr$_x$(PO$_4$)$_2$) doped with $x$ mol% Eu$^{3+}$ ions crystalline powders (where $x = 0.5, 1, 2, 3, 5$ mol%) was followed by the powder XRD
measurements both as a function of annealing temperature and of concentration of dopant ions (see Figure 1). The crystallinity was detected for all prepared materials sintered at different temperatures (750 – 950 °C per 3 h). The presence of both phases in the final product was confirmed by the correlation of the diffraction patterns with the reference standard of the hexagonal strontium hydroxyapatite ascribed to the P6_3/m space group form Inorganic Crystal Structure Database (ICSD-2866) [27] and the trigonal tristrontium diphosphate ascribed to the R3̅-m space group (ICSD-150869) [28]. All the materials heat-treated at 750–950 °C have shown detectable crystallinity in the form of sharp, narrow diffraction peaks. All registered diffraction peaks were assigned to the hydroxyapatite and tristrontium diphosphate. No extra diffraction peaks were observed. The ratio of strontium phases is different and are depended on the heat-treating temperature as well as on europium(III) ions concentration (see Table 1). With an increase of annealing temperature, the content of the Sr_3(PO_4)_2 crystal phase in composite increases due to the dehydration process of hydroxyapatite occurring at elevated temperature. It is not possible to detect any trend according to content of particular crystal phases when the concentration of the Eu^{3+} in composites increases. The Sr_3(PO_4)_2 crystal phase amount in composite increases from 37% for 750 °C temperature to 52% for 950 °C in thermally treated materials. This phenomenon is related to dehydration process occurring in hydroxyapatite matrix under elevated temperature.

![X-ray powder diffraction patterns](image-url)

**Figure 1.** X-ray powder diffraction patterns of the Sr_{10}(PO_4)_6(OH)_2–Sr_3(PO_4)_2 composite doped with 3 mol% Eu^{3+} ions as a function of annealing temperature (up) and annealed at 750 °C composites as a function of dopant ions concentration (bottom).
The structural refinement has been performed to get the unit cell parameters and percentage composition of the composite. The hexagonal and trigonal phases formation as well as the effective incorporation of Eu\(^{3+}\) ions have been confirmed. A good agreement between the recorded XRD pattern and the theoretical fit has been found. It indicates the success of the Rietveld refinement method. More details regarding Rietveld refinement are displayed in Table 1. As can be seen, it is possible to observe a shrinkage of both cells volume and a parameters of the SrHAp as well as the TSP materials with an increase of the Eu\(^{3+}\) ions concentration, which is caused by smaller ionic radii of dopant (Sr\(^{2+}\) (CN9) - 1.31 Å, Eu\(^{3+}\) (CN9) - 1.12 Å and Sr\(^{2+}\) (CN7) - 1.21 Å, Eu\(^{3+}\) (CN7) - 1.01 Å) as well as tristrontium diphosphate (Sr\(^{2+}\) (CN6) - 1.18 Å, Eu\(^{3+}\) (CN6) - 0.937 Å and Sr\(^{2+}\) (CN10) - 1.36 Å, Eu\(^{3+}\) (CN9) - 1.12 Å) [29]. Moreover, the a and V cells parameters of both composite materials marginally decreased after thermal treatment, which is an expected and well-known consequence caused by increase of particle size. An increase of annealing temperature caused an increase of Sr\(^{(PO_4)}\) crystal phase content in composite.

**Table 1.** Unit cell parameters (a and c), crystal cell volume (V), the percentage of particular phases calculated based on the Rietveld methods, and refined factors (R\(\omega\)) presented for the Sr\(^{(PO_4)}\)(OH)\(_2\)--Sr\(^{(PO_4)}\): powder composites as a function of dopant concentration and sintering temperature.

| Sample                  | Cell parameters | Phase | SRHAp (%) | TSP (%) | R\(\omega\) |
|-------------------------|-----------------|-------|-----------|---------|-------------|
|                         | Belovite Sr\(^{(PO_4)}\)(OH)\(_2\)|       |           |         |             |
|                         | Tristrontium Bis(phosphate(V)|       |           |         |             |
|                         | Sr\(^{(PO_4)}\): |       |           |         |             |
|                         | a (Å) | c (Å) | V (Å\(^3\)) | a (Å) | c (Å) | V (Å\(^3\)) |       |           |         |             |
| s. c.                   | 9.745 (1) | 7.265 (1) | 597.49 (15) | 5.3975 (2) | 19.818 (3) | 500.01 (8) | –       | –         | –        | –         |
| 0.5% mol Eu\(^{3+}\); 750°C | 9.769 (1) | 7.282 (1) | 601.84 (07) | 5.3912 (5) | 19.794 (9) | 498.23 (6) | 78.16 | 21.84 | 1.38 |
| 1% mol Eu\(^{3+}\); 750°C | 9.767 (3) | 7.282 (1) | 601.59 (43) | 5.3913 (1) | 19.798 (1) | 498.35 (5) | 82.79 | 17.21 | 1.30 |
| 2% mol Eu\(^{3+}\); 750°C | 9.769 (2) | 7.278 (5) | 601.51 (01) | 5.3921 (2) | 19.802 (6) | 498.60 (4) | 80.36 | 19.64 | 1.20 |
| 3% mol Eu\(^{3+}\); 750°C | 9.758 (9) | 7.270 (1) | 599.49 (65) | 5.3892 (6) | 19.795 (8) | 497.89 (1) | 62.65 | 37.35 | 1.28 |
| 5% mol Eu\(^{3+}\); 750°C | 9.760 (5) | 7.270 (1) | 599.74 (22) | 5.3886 (5) | 19.796 (7) | 497.80 (6) | 80.55 | 19.45 | 1.55 |
| 3% mol Eu\(^{3+}\); 850°C | 9.748 (5) | 7.262 (7) | 597.61 (01) | 5.3904 (1) | 19.798 (7) | 498.18 (9) | 50.81 | 49.19 | 2.12 |
| 3% mol Eu\(^{3+}\); 950°C | 9.735 (1) | 7.251 (1) | 595.11 (44) | 5.3904 (5) | 19.796 (4) | 498.13 (8) | 47.69 | 52.31 | 1.46 |

The visualization of the ideal hexagonal strontium apatite unit cell and trigonal tristrontium diphosphate unit cells as well as the coordination polyhedral of Sr\(^{2+}\) cations in hydroxyapatite lattice are current: four at the Sr(1) site with Cs point symmetry situated on the ternary axes and six at the Sr(2) site with Cs symmetry [30,31]. Moreover, in apatite structure, cis and trans symmetry of Cs crystallographic site correlated with arrangements of Eu\(^{3+}\) ions along the channel is well-known [8]. In the Sr\(^{(PO_4)}\): matrix, there are two nonequivalent crystallographic sites of Sr\(^{2+}\) ions: three at the Sr(1) site and six at the Sr(2) site. The atoms at the Sr(1) crystallographic site are placed on the threefold axis coordinated by 6 oxygen atoms belonging to the PO\(^{4-}\) group and site symmetry Cs\(_v\). The ions at the Sr(2) site are coordinated by ten oxygen atoms having Cs symmetry [32]. Nevertheless, a charge and an ionic radii incompatibility between Sr\(^{2+}\) and Eu\(^{3+}\) ions are present, leading to some structural defects correlated with the charge balance necessity.
The morphology of the Sr$_{10}$(PO$_4$)$_6$(OH)$_2$-Sr$_3$(PO$_4$)$_2$ composite doped with 3 mol% Eu$^{3+}$ ions annealed at 750, 850 and 950 °C was studied using a Scanning Electron Microscope. The particles are aggregated, their shape is asymmetrical and elongated in one direction (see Figure 3). The morphology of particles was not changed with an increase of the annealing temperature. However, the particle size was changed with an increase of the annealing temperature. The particle size distributions were counted and displayed as histograms. The size distributions are relatively wide, which is caused by different grain growth speed of each phase. The average size of the Sr$_{10}$(PO$_4$)$_6$(OH)$_2$-Sr$_3$(PO$_4$)$_2$ composite doped with 3 mol% Eu$^{3+}$ annealed at 750, 850 and 950 °C are about 100 (± 3.4), 131 (± 2.6) and 173 nm (± 4.6 nm), respectively. The energy-dispersive X-ray spectroscopy was used to confirm the content of elements in obtained materials.

The element maps were recorded by the SEM-EDS technique to analyze the elements distribution in the obtained material (Figure 4). The results confirmed the stated stoichiometry of dopants and regular distribution of elements.
Figure 3. The particle size distribution (left) based on SEM images (middle) as well as EDS spectra (right) of the Sr$_{10}$(PO$_4$)$_6$(OH)$_2$–Sr$_3$(PO$_4$)$_2$ composite doped with 3 mol% Eu$^{3+}$ ions annealed at 750 (a), 850 (b) and 950 °C (c).
Figure 4. SEM image and EDS elemental maps of the 3 mol% Eu$^{3+}$:Sr$_{60}$(PO$_4$)$_6$(OH)$_2$–Sr$_3$(PO$_4$)$_2$: composite annealed at 750 °C.

The FT-IR spectra were measured to identify the existence of orthophosphate and hydroxyl groups in obtained materials (Figure 5). The infrared spectra consist of ordinary PO$_4^{3-}$ vibration bands: the $\nu_2$ bending at 448 cm$^{-1}$; 564 cm$^{-1}$, the $\nu_4$ vibration at 596 cm$^{-1}$; the $\nu_1$ symmetric stretching at 950 cm$^{-1}$; the $\nu_3$ antisymmetric stretching at 1033 cm$^{-1}$ and 1081 cm$^{-1}$. Moreover, the vibrational transitions observed at 1632 cm$^{-1}$ 3432 cm$^{-1}$ (v OH) confirm the presence of a hydroxyl group in the obtained materials [33–35].
Figure 5. FT-IR spectra of the annealed 3 mol% Eu³⁺: Sr₁₀(PO₄)₆(OH)₂–Sr₃(PO₄)₂ composite annealed at different temperatures.

3.2. Emission and Excitation Spectra

The synthesised materials' excitation-emission spectra were measured at room temperature and observed at 615 nm emission wavelength of the maximum of the most intense ¹D₀ → ⁷F₁ electric dipole transition. All recorded spectra were alike to each other, thus only the representative excitation spectra of 1 mol% Eu³⁺: Sr₁₀(PO₄)₆(OH)₂–Sr₃(PO₄)₂ annealed at 750 °C was presented in Figure 6. The weak intraconfigurational 4f-4f transitions with sharp lines are characteristic for the Eu³⁺ ions as well as the intense ligand-to-metal charge transfer (CT) O²⁻ → Eu³⁺ with broad band in the UV range are typical for the Eu³⁺ ions incorporated into oxide matrix. The narrow lines observed at 339.5 nm (29 445 cm⁻¹) were attributed to the ⁷F₀ → ⁵H₄, ⁵L₅, ⁵G₆, at 383.5 nm (26 076 cm⁻¹) to the ⁷F₀ → ⁵D₄, ⁵S₅, at 415.8 nm (24 050 cm⁻¹) to the ⁷F₀ → ⁵D₃, at 465.3 nm (21 492 cm⁻¹) to the ⁷F₀ → ⁵D₂, at 533.3 nm (18 751 cm⁻¹) to the ⁷F₀ → ⁵D₁ and at 572 nm (17 483 cm⁻¹) to the ⁷F₀ → ⁵D₀ transition. It is well known that the 4f-sub-shells of lanthanide ions (RE³⁺) are well protected by the 5s and 5p sub-shells and the energy levels of the 4f-electrons are weakly influenced by the RE³⁺ ion vicinity. These transitions are forbidden by Laporte parity rule that is why the f-f transition intensities are weak. However, quite the opposite, the charge-transfer (CT) transition is allowed and strongly affected by electron-lattice coupling. Moreover, its energy level is depended on an ion local symmetry [36–38]. The CT maximum is located at 253 nm (39 526 cm⁻¹).
The emission spectra of \( x\text{Eu}^{3+}: \text{Sr}_{10} \left( \text{PO}_4 \right)_{6} \left( \text{OH} \right)_{2} - \text{Sr}_3 \left( \text{PO}_4 \right)_{2} \) (where \( x = 0.5 - 5 \text{ mol\%} \)) materials annealed at the temperature 750, 850 and 950 °C were recorded upon pulsed 266 nm and 395 nm excitation at 300 K as well as at 80 K (Figure 7) and as a function of \( \text{Eu}^{3+} \) ions concentration excited by 266 nm at 300 K and 80 K (Figure 8). The spectra excited by 395 nm line were normalized to the \( ^{5}D_0 \rightarrow ^{7}F_1 \) transition, which can be treated as an internal reference. In the emission spectra measured upon excitation by 266 nm, a broad peak at about 421 nm was observed, which corresponds to the \( 4f^6 5d^1 \rightarrow 4f^7 \left( ^8S_{7/2} \right) \) emission of \( \text{Eu}^{2+} \) ions as well as narrow \( 4f-4f \) transitions were present, which are typical for \( \text{Eu}^{3+} \) ions. As can be seen, an increase of the annealing temperature results in an increase of the \( \text{Eu}^{2+} \) ions emission intensity and a decrease of \( \text{Eu}^{3+} \) emission intensity (see Figure 7 upside). The \( \text{Eu}^{2+} \) emission measured at 80 K is less intense compared to the \( \text{Eu}^{3+} \) emission measured at 300 K. A change in the annealing temperature and measurement temperature results in changes of the \( \text{Eu}^{2+} \) to \( \text{Eu}^{3+} \) relative intensity ratio. Moreover, the intensity of the \( \text{Eu}^{2+} \) ions emission depends on the measurement temperature and it is more intense in higher temperatures (see Figure 7). This was caused by an increase of \( \text{Eu}^{2+} \) fraction in the composite. The detailed description of reduction mechanism is provided in Section 3.3 and 3.4.
Figure 7. Emission spectra of the Sr_{10}(PO_{4})_{6}(OH)_{2}-Sr_{3}(PO_{4})_{2} composite doped with 3 mol\% Eu^{3+} ions as a function of post-heat treatment temperature excited by 266 nm (above) and by 395 nm line (below) measured at 300 K (left) and 80 K (right).

The Eu^{3+} emission spectra consist of characteristic electron transitions occurring in the 4f shells appearing in the red region of the electromagnetic radiation. The \(^{5}D_{0} \rightarrow {^{7}}F_{0}\) transition is located at 576.9 nm (17 334 cm\(^{-1}\)), the \(^{5}D_{0} \rightarrow {^{7}}F_{1}\) transition at 585.9 nm (17 068 cm\(^{-1}\)), the \(^{5}D_{0} \rightarrow {^{7}}F_{2}\) transition at 616.5 nm (16 221 cm\(^{-1}\)), the \(^{5}D_{0} \rightarrow {^{7}}F_{3}\) transition at 655.2 nm (15 262 cm\(^{-1}\)), the \(^{5}D_{0} \rightarrow {^{7}}F_{4}\) transition at 700.5 nm (14 275 cm\(^{-1}\)), the \(^{5}D_{0} \rightarrow {^{7}}F_{5}\) transition at 742.7 nm (13 464 cm\(^{-1}\)) and the \(^{5}D_{0} \rightarrow {^{7}}F_{6}\) transition at 805.3 nm (12 418 cm\(^{-1}\)). The \(^{5}D_{0} \rightarrow {^{7}}F_{0}\) transition is split into three components located at 571.6 nm (17 495 cm\(^{-1}\)), 576.9 nm (17 334 cm\(^{-1}\)) and 579.9 nm (17 244 cm\(^{-1}\)), indicating that Eu^{3+} ions are located at least into three different crystallographic sites in this composite [39–42]. A difference between the intensity ratio of components is visible.

The hypersensitive \(^{5}D_{0} \rightarrow {^{7}}F_{2}\) emission transition is the most intense in all cases. The intensity of this transition is very sensitive to variations in the local environment of Eu^{3+} ions in the host. On the other hand, the intensity of the \(^{5}D_{0} \rightarrow {^{7}}F_{1}\) magnetic dipole transition remains almost independent on the crystal field. When Eu^{3+} ions occupy a centrosymmetric site, the only permissive transition is magnetic one. In the opposite, the electric dipole transition is cardinal. The asymmetry ratio of the relevant emission intensities (R) is defined followingly [43]:

\[
R = \frac{\int {^{5}}D_{0} \rightarrow {^{7}}F_{2}}{\int {^{5}}D_{0} \rightarrow {^{7}}F_{1}}
\]  

The integrated intensities ratio of these transitions is used to evaluate the asymmetry of the Eu^{3+} ions coordination polyhedron. The higher the relationship between these transitions is the less centrosymmetric the local environment of Eu^{3+} ions becomes. The annealing temperatures and Eu^{3+} ions concentration impacts on the R value is gathered in Table 2. Any relationship between an increase of annealing temperature and dopant concentration is not observed. The crystallography sites in phosphate lattice are randomly occupied by Eu^{3+} ions what is related to the Eu^{3+} ion...
concentration and annealing temperature. Moreover, the observed emission is related to different crystallographic sites. That is why the R value behaves incalculably in such cases.

The Stark’s components are better visible in the case of emission spectra measured at 80 K (see Figure 7). The emission spectra upon excitation by 395 nm line were also measured to be compared. As can be seen, some differences between these two excitation lines are well visible as different shapes, ratios, and intensity of transitions. The 3 mol% Eu³⁺: Sr₁₀(PO₄)₆(OH)₂–Sr₃(PO₄)₂ material stands out by the abnormal high intensity of the ⁵D₀ → ⁷F₀ transition upon excitation by the 395 nm line.

The emission spectra of the Eu³⁺-doped Sr₁₀(PO₄)₆(OH)₂–Sr₃(PO₄)₂ composite depend on the Eu³⁺ ions concentration measured upon 266 nm excitation at 300 and 80 K is presented in the Figure 8. As can be observed, an increase of the europium ions concentration in material led in a decrease of the photoluminescence (PL) intensity of Eu²⁺ ions. This insight can be caused by the Eu²⁺ → Eu³⁺ energy transfer, which prompts in the quenching of the Eu³⁺ emission. In materials doped with low Eu³⁺ ions concentration, the Eu²⁺ ions are weakly quenched due to the statistically enormous distance between these two ions. An increase of dopant concentration caused an increase of the Eu²⁺ → Eu³⁺ energy transfer efficiency as a result of a bigger concentration of ions in the host [7, 44]. This could also be caused by the different emission intensity in two compounds due to the change of Sr₁₀(PO₄)₆(OH)₂ to Sr₃(PO₄)₂ proportion with an increase of Eu ions doping.

On the other hand, the decrease of Eu²⁺ PL intensity may be clarified by the fact that the amount of V‴Sr vacancies also increase with an increase of the Eu³⁺ concentration and an excess of the Eu³⁺ ions would occupy the V‴Sr position. Consequently, the double negative vacancy (V‴Sr) would be converted into a positive charge hereupon, it cannot move negative charge to Eu³⁺ and the reduction was inhibited. To have electroneutrality, the composite may absorb oxygen from air.

![Figure 8: Emission spectra of the Sr₁₀(PO₄)₆(OH)₂–Sr₃(PO₄)₂ composite doped with Eu³⁺ ions as a function of the Eu³⁺ ions concentration annealed at 750 °C, excited by 266 nm line, measured at 300 K (left) and 80 K (right).](image)

The Ω₂ and Ω₄ intensity parameters was calculated based on the Judd–Ofelt theory and the results are listed in Table 2. The Ω₂ parameter value changes lightly with an increase of Eu³⁺ ion concentration and with an increase of annealing temperature, but without any well visible tendency. It indicates the changes in the europium coordination polyhedra and could be related to changes of the Eu³⁺–O²⁻ bond covalency. The Ω₄ value brings information about the electron density variations around Eu³⁺ cations. However, the Eu³⁺ ions symmetry changes cannot be directly interpreted by the Ω₂ value but can add some information about the electron density deviations in the O²⁻ anions surrounding, what influences on the CT band position.
Table 2. Decay rates of radiative (A_rad), nonradiative (A_nonrad) and total (A_tot) processes of the $^3D_0 \rightarrow ^7F_i$ transitions, luminescence lifetimes (T), intensity parameters ($\Omega_i$, $\Omega_o$), quantum efficiency ($\eta$) and asymmetry ratio (R) of the Sr$_{10}$(PO$_4$)$_6$(OH)$_2$-Sr$_3$(PO$_4$)$_2$: composites.

| Sample | A_rad ($s^{-1}$) | A_nonrad ($s^{-1}$) | A_tot ($s^{-1}$) | T (ms) | $\Omega_i$ (10$^{-3}$cm$^2$) | $\Omega_o$ (10$^{-3}$cm$^2$) | $\eta$ (%) | R |
|--------|-----------------|---------------------|-----------------|--------|---------------------------|---------------------------|-----------|---|
| 750 °C | 162.56          | 86.11               | 253.67          | 3.94   | 3.2704                    | 3.2757                    | 66.05     | 2.321 |
| 850 °C | 154.13          | 73.81               | 227.94          | 4.39   | 2.9401                    | 2.9301                    | 67.62     | 2.087 |
| 950 °C | 164.66          | 150.97              | 315.63          | 3.17   | 3.3232                    | 2.9414                    | 52.17     | 2.399 |

\[
x \text{mol% Eu}^{3+}: \text{Sr}^{3+} \text{(PO}_4\text{)}_6\text{(OH)}_2 \rightarrow \text{Sr}^{3+} \text{(PO}_4\text{)}_2: \text{750 °C}
\]

3.3. Abnormal Reduction Mechanism of Eu$^{3+} \rightarrow$ Eu$^{2+}$

The abnormal reduction of the Eu$^{3+}$ to the Eu$^{2+}$ ion is known in the literature for solid-state materials made at elevated temperature in air [45–47]. Su et al. introduced four demands for these abnormal reductions of the Eu$^{3+}$ ions to the Eu$^{2+}$ ions in a solid-state host:

1. no oxidizing ions are present in compounds,
2. the Eu$^{3+}$ ions substitute the cations with lower valences in compounds,
3. similar radii of the substituted cation and Eu$^{2+}$ ion, and
4. the tetrahedral anion groups (BO$_4$, PO$_4$, AlO$_4$, SiO$_4$) are present in compound [47].

In the Sr$_{10}$(PO$_4$)$_6$(OH)$_2$-Sr$_3$(PO$_4$)$_2$: composite, the Sr$^{3+}$ and PO$_4$$^{3-}$ are not oxidizing ions, which satisfied condition (1). The Eu$^{3+}$ ions substituted Sr$^{3+}$ sites in this composite, which means that Eu$^{3+}$ ions substituted cations with lower valences. Moreover, the Sr$^{3+}$ ion has similar radii to Eu$^{2+}$ (in hydroxyapatite: Sr$^{2+}$ (CN9) - 1.31 Å, Eu$^{3+}$ (CN9) - 1.30 Å and Sr$^{2+}$ (CN7) - 1.21 Å, Eu$^{2+}$ (CN7) - 1.20 Å), in tristrontium diphosphate: Sr$^{2+}$ (CN9) - 1.18 Å, Eu$^{3+}$ (CN6) - 1.17 Å and Sr$^{2+}$ (CN10) - 1.36 Å, Eu$^{2+}$ (CN9) – 1.35 Å [29]) and meet conditions (2) and (3). Furthermore, the condition (4) is also achieved, because this composite is built by PO$_4$$^{3-}$ anions groups. Based on this conditions and emission spectra, it is expected that the Eu$^{3+}$ to Eu$^{2+}$ reduction occur in these composites. The Eu$^{3+}$ ions were partially reduced to the Eu$^{2+}$ and this reduction in composites obtained in air atmosphere at elevated temperature can be clarified by the charge compensation mechanism based on the Kröger–Vink notation (′ sign means charge neutrality, ″ means positive charge and ′ negative charge in particular crystallographic site. The vacancy is marked by V and interstitial atom by i). When the trivalent Eu$^{3+}$ ions substituted the Sr$^{3+}$ ions, two Eu$^{3+}$ ions would substitute for three Sr$^{3+}$ ions to preserve charge electroneutrality and a double negative vacancy in the Sr crystallographic site (V$_{Sr}$) is created. Consequently, two Eu$^{3+}$ positive defects and one V$_{Sr}$ negative vacancy would be designed by each substitution for every two Eu$^{3+}$ ions:

\[
\text{EuO}_3 + 3\text{Sr}^{3+} \rightarrow 2(\text{Eu}^{3+})^{\prime}\text{Sr} + \text{V}^{\prime}\text{Sr} + 3\text{CaO};
\]

Two relative negative charges are composed on the hydroxyl sides:

\[
\text{EuO}_3 + 2\text{Ca}^{2+}\text{Ca} + 2\text{OH}^{-}\text{OH} \rightarrow 2(\text{Eu}^{3+})^{\prime}\text{Ca} + 2\text{O}^{\prime}\text{OH} + 2\text{CaO} + \text{H}_2\text{O};
\]

In addition, charge balance can also be reached by the creation of interstitial oxygen O$^{	ext{i}}$ with double relative negative charge:

\[
\text{EuO}_3 + 2\text{Ca}^{2+}\text{Ca} \rightarrow 2(\text{Eu}^{3+})^{\prime}\text{Ca} + \text{O}^{\prime}\text{i} + 2\text{CaO};
\]

The vacancy V$_{Sr}$ could behave as a donor of electrons pending the two Eu$^{3+}$ defects turn into acceptors of electrons. Via thermal stimulation, the negative charges of the V$_{Sr}$ vacancy defects would be thereafter moved to the Eu$^{3+}$ positions and reduce them to their Eu$^{2+}$ form followingly:
The higher annealing temperature, the mechanism of the Eu$^{3+}$ to Eu$^{2+}$ reduction is more probable, which is well visible on the emission spectra.

3.4. Thermal Stability

The luminescence intensity of the 1 mol% Eu$^{3+}$: Sr$_{10}$(PO$_4$)$_6$(OH)$_2$–Sr$_3$(PO$_4$)$_2$ was measured depending on the measurement temperature in the range of 80–550 K (see Figure 9). It was observed that the PL intensity of Eu$^{3+}$ ion decreases with an increase of the measurement temperature but on the other hand the PL intensity of Eu$^{2+}$ ion increases. Mechanism of this behavior has been described as abnormal intense Eu$^{2+}$ emission in high temperature [16,48]. It could be related to the presence of two possible defect traps of $[2$(Eu$^{3+}$)$_n$ – O$_{i''}$] or $[2$(Eu$^{3+}$)$_n$ – V$_{Sr}^{-}$] in this material. The Eu$^{3+}$ ions could recombined with the electrons released from defects O$_{i''}$ or V$_{Sr}^{-}$; nevertheless, the electrons do not get into the Eu$^{2+}$ ion but loosely couple with Eu$^{3+}$ in the near environment (Eu$^{3+}$ + e$^-$) manifesting emission of Eu$^{2+}$. All the defects can be inert at low temperature what yield low luminescence of Eu$^{2+}$ ion when the temperature decrease. The CIE 1931 chromaticity diagram is presented in the Figure 10 determined from emission spectra measured in 80, 300 and 550 K of temperature.

**Figure 9.** The luminescence intensity dependence on the measurement temperature of 1 mol% Eu$^{3+}$: Sr$_{10}$(PO$_4$)$_6$(OH)$_2$–Sr$_3$(PO$_4$)$_2$ in the temperature range of 80–550 K. Inset: dependence of integrated intensity of the Eu$^{2+}$ and Eu$^{3+}$ emission on the measurement temperature.
3.5. Fluorescence Dynamics

The luminescence kinetics were analyzed for all synthesized materials to determine the comprehensive characteristics of the luminescence properties. The decay times at room temperature (RT) corresponding to the $^5D_0 \rightarrow ^7F_2$ transition were excited by pulse radiation of 395 nm and monitored at 615 nm (Figure 11). The decay curves are not single-exponential, what is compatible with the presence of nonequivalent crystallographic sites of Eu$^{3+}$ ions accordingly, the lifetimes values were calculated as the effective emission decay time by using Equation (1). The decay profiles and the lifetime values are alike to each other and only minor differences were observed. The decay times of Eu$^{3+}$ ions in Sr$_{10}$(PO$_4$)$_6$(OH)$_2$--Sr$_3$(PO$_4$)$_2$ are typical for this kind of ion and are longer than for strontium apatite [7,8], which is caused by the presence of Sr$_3$(PO$_4$)$_2$ phase.
The decay curve (blue component) of the 1 mol% Eu$^{3+}$: Sr$_{10}$(PO$_4$)$_6$(OH)$_2$–Sr$_3$(PO$_4$)$_2$ composite annealed at 750 °C was measured under excitation at 266 nm and monitored at 430 nm (Figure 12). The single exponential shape of the decay curve indicates to the Eu$^{2+}$ ion emission from one crystallographic site. The lifetime value was calculated based on Equation (2) and was determined to be 0.674 µs. The emission spectra of blue emission from the streak camera is presented in Figure 13.

Figure 12. Luminescence kinetics of the blue component of the 1 mol% Eu$^{3+}$: Sr$_{10}$(PO$_4$)$_6$(OH)$_2$–Sr$_3$(PO$_4$)$_2$ composite annealed at 750 °C. Excited by 266 nm and observed at 430 nm.
Figure 13. The emission spectrum image form streak camera of the 1 mol% Eu$^{3+}$: Sr$_{10}$(PO$_4$)$_6$(OH)$_2$-Sr$_3$(PO$_4$)$_2$ composite annealed at 750 °C.

4. Conclusions

A microwave-assisted hydrothermal method was used with success to produce Eu$^{3+}$-doped Sr$_{10}$(PO$_4$)$_6$(OH)$_2$-Sr$_3$(PO$_4$)$_2$: composites. The structural and luminescence properties were investigated in a function of dopant ions concentration and annealing temperature. The formation of the strontium phosphate composite was confirmed by the XRPD analysis. The unit cell parameters as well as the percentage content of crystal phases, were calculated by Rietveld refinement method. The obtained materials have had a tendency to agglomerate to minimize their excess surface energy and their shapes were irregular and elongated. There were no observed changes in morphology with an increase of the annealing temperature. There were observed the narrow 4f-4f transitions typical for Eu$^{3+}$ ions and a broadband at 421 nm related to the 4f$^5$5d$^1$ → 4f$^6$(S$^7/2$) transition from Eu$^{2+}$ ions. Moreover, it has been described as an abnormal intense emission of Eu$^{2+}$ ion due to recombination of the excited state of Eu$^{3+}$ ion with the electrons released from defects (O$^{2-}$ or V$_{Sr}^{-}$).

The decay curves of Eu$^{3+}$ ions are not single-exponential as a result of the nonequivalent crystallographic sites of Eu$^{3+}$ ions in the matrix. However, the lifetime value of Eu$^{2+}$ ions is single-exponential and was determined to be 0.674 µs.

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