Emission tunability and local environment in europium-doped OH⁻-free calcium aluminosilicate glasses for artificial lighting applications

Aline M. Farias a, Marcelo Sandrini a, José Renato M. Viana b, Mauro L. Baesso a, Antônio C. Bento a, Jurandir H. Rohling a, Yannick Guyot b, Dominique De Ligny c, Luiz Antônio O. Nunes d, Flávio G. Gandra e, Juraci A. Sampaio f, Sandro M. Lima g, Luis Humberto C. Andrade e, Antônio N. Medina a,*,

a Departamento de Física, Universidade Estadual de Maringá, Av Colombo, 5790, 87020-900, Maringá, PR, Brazil
b Laboratoire de Physico-Chimie des Matériaux Luminescents, Université de Lyon, Université Claude Bernard Lyon 1, Villeurbanne, UMR 5620 CNRS 69622, France
c Department of Materials Science and Engineering, University of Erlangen Nürnberg, Martens str. 5, 91058, Erlangen, Germany
d Instituto de Física de São Carlos, Universidade de São Paulo, Av. Trabalhador São-Carlense 400, 13566-590, São Carlos, SP, Brazil
e Instituto de Física Gleb Wataghin, Universidade Estadual de Campinas, 13083-859, Campinas, SP, Brazil
f Grupo de Spectroscopia Óptica e Fototermica, Universidade Estadual de Mato Grosso do Sul-UEMS, Dourados, MS, C. P. 351, CEP 79804-970, Brazil

HIGHLIGHTS

- Eu²⁺-doped OH⁻-free calcium aluminosilicate glass as a new source for white lighting.
- Correlation between emission tunability and local environment of europium ions.
- Significant reduction of Eu³⁺ to Eu²⁺ by melting the glasses under vacuum atmosphere.
- Broad, intense and tunable luminescence ranging from blue to red.

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ABSTRACT

The relationship between emission tunability and the local environment of europium ions in OH⁻-free calcium aluminosilicate glasses was investigated, focusing on the development of devices for artificial lighting. Significant conversion of Eu³⁺ to Eu²⁺ was obtained by means of melting the glasses under a vacuum atmosphere and controlling the silica content, resulting in broad, intense, and tunable luminescence ranging from blue to red. Electron spin resonance and X-ray absorption near edge structure measurements enabled correlation of the luminescence behavior of the material with the Eu³⁺/Eu²⁺ concentration ratio and changes in the surrounding ions' crystal field. The coordinates of the CIE 1931 chromaticity diagram were calculated from the spectra, and the contour maps showed that the light emitted from Eu²⁺ presented broad bands and enhanced color tuning, ranging from reddish-orange to blue. The results showed that these Eu doped glasses can be used for tunable white lighting by combining matrix composition and the adjustment of the pumping wavelength.

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1. Introduction

A major challenge in the development of a new generation of artificial lighting is the tailoring of phosphor materials that possess efficient and tunable emissions across the whole range of the visible spectrum [1–7]. Europium and cerium are currently the most widely used phosphors in devices for lighting and displays, and both can coexist in several oxidation states in the host matrices. In addition, their emissions are known to be strongly dependent on the surrounding ions' crystal field [2].

* Corresponding author.
E-mail address: medina@dfi.uem.br (A.N. Medina).

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There are several factors that determine the luminescence properties of a material, such as the phonon energy, the presence of impurities, the nature and intensity of the surrounding crystal field, the oxidation states of the sensitizer ions, the synthesis procedure, and the morphology and geometry of the final material. From the point of view of materials science, tailoring of the surrounding ions’ crystal field is essential in order to control the luminescence behavior. Consequently, many different materials have been developed for solid-state lighting, including crystals [2], ceramics [8], polymers [3,9], and glasses [1,7,10–12]. Specifically, the amorphous nature of glasses means that their broad luminescence properties are influenced by the average local crystal field surrounding the dopant ions. Selection of the matrix composition, the phosphor doping level, and the melting procedure are important ways of obtaining the desired emission characteristics.

We recently proposed the use of OH−-free low-silica calcium aluminosilicate (LSCAS) glass as an alternative luminescent material for tunable white lighting, due to its intense and broad emission spectrum in the visible region. These studies employed materials doped with Ce [7,12], Eu [10], and Ce–Eu [11]. The results revealed a significant influence of melting under a vacuum atmosphere on the production of glasses with ions in reduced oxidation states and with minimal presence of OH− in their structures. This LSCAS glass is known to have superior thermomechanical properties, good chemical resistance, transparency from the UV-visible to the near infrared (up to 5 μm), reduced oxidation states for some dopant ions, and efficient laser emissions [13–18].

The certification of visible light sources for artificial lighting can employ parameters such as the correlated color temperature (CCT), the color rendering index (CRI), and the distance from the Planckian locus to the (u', v') color coordinates. The latter is known as Du'v' and describes how close the tested light source is in relation to the ideal lighting. These parameters, which characterize the emission, can be used to evaluate the melting procedure and the changes in composition, in order to optimize the glass luminescence. To this end, knowledge of the oxidation states of the sensitizer ions and the nature of the surrounding crystal field are important for tailoring the intended luminescent source.

Given the above considerations, this work therefore involved the development of a set of OH−-free Eu-doped aluminosilicate glasses with different silica contents. Investigation was made of the influence of the surrounding ions’ crystal field and their oxidation states on the luminescence properties. The techniques used for characterization of the samples were electron spin resonance (ESR), X-ray absorption near edge structure (XANES), and photoluminescence (PL).

### 2. Experimental

High purity reagents (>99.995%) were used to prepare the glass samples. Table 1 shows the compositions (in wt%) required in order to obtain 6 g of the glasses. The reagents were mixed in a ball mill for 12 h, after which the mixtures were melted in graphite crucibles for 2 h under a vacuum atmosphere (10⁻³ mbar), at temperatures between 1400 °C and 1600 °C, according to the specific composition. Quenching was achieved by switching off the heater and moving the crucible up to a cooled vacuum chamber. Annealing was then performed by returning the crucible to the melting position, where the temperature was around 800 °C, and allowing it cool to room temperature, while maintaining the vacuum. This procedure took about 5 h. Further details of the preparation process can be found elsewhere [15,18]. The glass samples presented good optical quality, with no evidence of crystallization. They were cut and polished for the luminescence and XANES measurements, while for ESR the samples were in the form of powders.

ESR measurements were performed at room temperature using the microwave in the X-band spectral region (ν = 9.52 GHz), with a power of 5 mW and a 10 Gauss amplitude modulated field at 100 kHz. X-ray LIII edge (6977 eV) absorption spectra were measured using the D04B-XAFS2 beamline of the Brazilian National Synchrotron Light Laboratory (Laboratório Nacional de Luz Síncrotron, LNLS) in Campinas, Brazil). The XANES spectra were collected in fluorescence mode using a Canberra GL00555 LEGe detector with an array of 15 elements [19]. The data were acquired as the averages of three scans for each sample. Excitation/emission contour maps were obtained using a Xe−lamp as the excitation source, with the wavelengths selected by a Jobin-Yvon H10D monochromator. The excitation beam was focused on the sample and the emission was collected by means of an optical fiber and detected by a CCD array connected to a microcomputer. Luminescence measurements were also carried out using an Ar+ ion laser at 355 nm - 365 nm and a diode laser at 532 nm as excitation sources, with detection by a silicon photodiode. The detection system was calibrated using an irradiance and illuminance standard source (from the National Institute of Standards and Technology, NIST) from Gooch & Housego, model 752-10 E.

### 3. Results and discussion

Fig. 1 shows the ESR spectra as a function of the silica content. The data were normalized in terms of the Eu2O3 weight. Eu3+ ions...
are not paramagnetic, so could not be detected in the ESR measurements. On the other hand, Eu$^{2+}$ has electronic spin $5/2$, enabling evaluation of the $^{8}S_{7/2}$ singlet ground state by this technique. Using 2,2-diphenyl-1-picrylhydrazyl (dpph) as a reference pattern, the ESR signal intensity was considered to be proportional to the amount of Eu$^{2+}$ ions in the sample. It can be seen (Fig. 1) that there were resonance lines with g 2, 2.7, 4.7, and 6.4, and that the signal intensity increased for higher silica contents (inset of Fig. 1). This reflected an increase in the Eu$^{3+}$/Eu$^{3+}$ ratio, because the total amount of europium in the sample remained constant.

The resonance lines at g 6, 2.7, and 2 are known as the characteristic U spectrum for Gd$^{3+}$ and Eu$^{3+}$ used as dopants in glasses or disordered polycrystalline materials [20,21]. The existence of several resonance lines, as opposed to just one at g 2.0, as observed in the ESR spectra of rare earth ions in the S state, has been used to demonstrate that Gd$^{3+}$, Eu$^{3+}$, and Tb$^{4+}$ present different interactions with the crystal matrix fields [20,22]. Iton and Brodbeck [21] simulated the ESR spectra of Gd$^{3+}$ and Eu$^{3+}$ in zeolites and compared the results with experimental data obtained in the X band. They proposed that the U spectrum lines indicated the occurrence of a moderate distortion of the crystal field in relation to the cubic, tetrahedral, or octahedral basic structures. It was observed that the detected distortions were predominant when the ions were in a high coordination number (>6). On the other hand, resonance at g 4.86 is indicative of a crystal field with orthorhombic symmetry and strong interaction between the ions, as shown by Iton and Turkovich [23]. Additionally, ESR spectra of both Gd$^{3+}$ and Eu$^{3+}$ in glasses, measured at low microwave frequency, have shown reduced intensities of the g = 6 U spectra, together with the generation of a broad line at around g = 5 [24,25]. From Fig. 1, it can be seen that with the increase of silica content, the intensities of lines related to the U spectrum increased considerably, relative to that for resonance at g 4.7. This suggests that in addition to promoting the reduction Eu$^{3+}$ → Eu$^{2+}$, the addition of silica resulted in the formation of weaker crystal field interactions in the sites surrounding the Eu$^{2+}$ ions.

Fig. 2 shows the normalized X-ray absorption spectra as a function of the energy in the XANES region. Two absorption peaks were observed, with differences in energy of about 8 eV, reflecting a hybrid structure with the existence of two europium valence states in the glasses [26]. The peak at lower energy was characteristic of Eu$^{3+}$, while that at higher energy was characteristic of the Eu$^{3+}$ ions [27]. A higher silica content in the matrices resulted in increased intensity of the Eu$^{2+}$ peak and consequently a reduction in the contribution from the Eu$^{3+}$ ions. The averaging was done by performing three scans in each sample and the spectra were fitting using the Athena software (IFFEFIT package) [28]. As europium shows separate absorption edges related to the Eu$^{2+}$ and Eu$^{3+}$ ions, two arc tangent functions were used to fit the step energy observed in the spectra, and two Gaussian functions were used to fit the peaks. The inset of Fig. 2 shows the amount of Eu$^{2+}$ as a function of the silica content, from which it can be seen that the amount increased up to three-fold at higher silica concentrations. This behavior was in agreement with the ESR data. The coexistence of Eu$^{3+}$ and Eu$^{3+}$ in OH−free LSCAS glass has been reported previously [10,11,29]. In an earlier study [29], overlapping of the emission lines between 570 and 750 nm was observed, attributed to the $^{5}D_{0} \rightarrow ^{7}F_{J}$ transitions of the Eu$^{3+}$ ion. The observed broad emission band centered at 585 nm was attributed to the $^{5}D_{0} \rightarrow ^{7}F_{1}$ transition of the Eu$^{3+}$ ion. For the samples with lower silica concentrations, there were narrow emission peaks due to Eu$^{3+}$ and a broad peak in the red-orange region due to Eu$^{2+}$. For higher silica concentrations, there were increases of about 400-fold in the intensity of the emission band related to Eu$^{2+}$, and the emission peak shifted to the blue region. This significant increase was related to the greater quantities of divalent ions in the samples with higher silica contents, as indicated by the inset in Fig. 2. The Eu$^{2+}$/Eu$^{3+}$ ratio is expected to decrease considerably at low silica contents, so it becomes easy to observe the Eu$^{3+}$ emission in Fig. 3. For all the samples, the selective Eu$^{3+}$ luminescence was measured using excitation at 532 nm. The results did not show any changes in the relative intensities of the bands, according to the silica content. However, as expected, a decrease in the total intensity was observed (shown in the inset of Fig. 1). The observed behavior was like a mirror, compared to the ESR intensity. Although the use of 350 nm does not directly excite Eu$^{3+}$, energy transfer from Eu$^{2+}$ to Eu$^{3+}$ by reabsorption is possible, due to the overlap among Eu$^{2+}$ absorption bands and Eu$^{3+}$ absorption lines, as reported recently [10].
Wang et al. [30] suggested that a decrease in the Eu²⁺ emission intensity can result from an increase in the number of non-bridging oxygens (NBOs), caused by a greater number of negative charges in the glass network. This favors the formation of ions with higher valence states, maintaining the neutrality of the material. It has been shown previously that the quantity of NBOs in calcium aluminosilicate glasses decreases when the amount of silica in the matrix is increased [31]. A calculation assuming all Ca and Mg to be modifiers and all Al and Si to be network formers was used to estimate the theoretical %NBO. The results (Table 1) showed that from the lowest to highest silica contents, the %NBO changed from 48% to 23%. Therefore, the increase in the Eu²⁺ emission intensity at higher silica contents was in agreement with the data published previously.

The red shift of the emission band of Eu²⁺ is often associated with the optical basicity of the glass (Lth). Following the work of Duffy and Infram [32], a theoretical optical basicity can be calculated for the glass studied here (Table 1). The increase in the optical basicity from 0.56 to 0.74, with decrease of the silica content, suggests that the crystal field strength of the glass was higher, resulting in a splitting of the Eu³⁺ d orbital degeneracy [33,34]. This would result in separation of the two energy levels, with eg and t₂g shifting to lower and higher energy, respectively [33,34]. In other words, the separation between these two bands depends on the strength of this interaction. In addition, a large separation implies a connection between the lower level (e₉) and the ground state 4f, resulting in emission with lower energy, at longer wavelengths. A similar red shift was also observed for Ce³⁺-doped LSCAS [7,12]. According to Novatski et al. [31], and in agreement with the estimated values given in Table 1, the optical basicity decreases when the silica concentration increases, so there is a decrease in the separation of the e₉ and t₂g bands, and consequently an increase in the separation between the ground state and the e₉ band. Thus, by increasing the concentration of silica, the emission peaks shift to lower wavelengths, as shown in Fig. 3. The decrease of the crystal field interaction with the d orbitals, as suggested by the emission results, is in agreement with the increasing intensity of the U spectrum observed in the ESR data.

The coordinates in the CIE 1931 chromaticity diagram were calculated from the emission spectra, as shown in the inset of Fig. 3. It can be observed that at higher silica contents, there was a shift in the emitted color from the orange-red region to the blue region.

### Table 2

<table>
<thead>
<tr>
<th>Samples</th>
<th>CRI</th>
<th>CQS</th>
<th>CCT</th>
<th>Du'v'</th>
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<tr>
<td>l₁ex1</td>
<td>l₁ex2</td>
<td>l₂ex1</td>
<td>l₂ex2</td>
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</table>

λ₁ex 350 nm; λ₂ex 399 nm.
This blue shift in the maximum emission intensity contributes to reduce the CRI values and to increase the CCT values for glasses with high silica. The photographs in Fig. 3 illustrate the differences between the emissions.

Fig. 4 shows contour plots of the excitation/emission spectra for the samples with 7, 35, 50, and 65 wt% of silica. As described above, the luminescence intensity is very different among the samples, so that in the figure they cannot be compared in intensity. In fact, only the maximum positions for the excitation and emission wavelengths can be checked. For the sample with 7 wt% of silica, the Eu²⁺ emission was much lower than that of Eu³⁺, while the Eu²⁺ emissions became more evident at higher silica concentrations. The plots indicated that the blue shift of the emission band observed for excitation at 350 nm also occurred for other excitation wavelengths. The excitation band showed similar behavior. The information obtained from the excitation/emission contour plots indicated that it was possible to tune the emission by modifying the oxidation states of the ions and/or the crystal matrix field.

The calculated values of the CRI, CQS, CCT, and Du’v parameters are shown in Table 2 and Fig. 5. The color changes were reflected by both an increase in Du’v and a decrease in the CRI values. The Du’v values lay outside the acceptable maximum of 0.006 established by the American National Standards Institute (ANSI C-78.377). However, for silica contents lower than 20%, the values complied with the Du’v limit value of <0.02 set by the Japanese Standards Association (JIS C-8152-2). It is important to note that these comparisons were made without considering complementary emissions from blue LEDs, which could be used for the construction of WLED devices. This would make it possible to obtain parameter values within the ranges established in current technical standards. It can be seen that there were substantial differences between the CRI and color quality scale (CQS) values (Table 2). The CQS is a new color quality parameter that is now accepted by NIST. It is considered superior for the measurement of color quality because it employs 15 standard pastel colors as a reference for color reproducibility, while the CRI uses only 8 standard colors. It is known that for some deep red color spectra, the CRI can give inaccurate results. This can be seen by comparing the CQS and CRI values (Table 2) for the aluminosilicate samples with low silica contents, which presented a red color under UV and violet excitation and showed the greatest differences between the CRI and CQS values. Despite this divergence between the CRI and CQS techniques, the CRI remains the internationally accepted parameter, according to the Commission on Illumination (CIE), and for much of the visible band there were only small differences between the CRI and CQS values.

4. Conclusions

In conclusion, the results showed the coexistence of Eu²⁺ and Eu³⁺ oxidation states in the aluminosilicate glasses analyzed here. It was observed that with increasing silica concentration, the matrix promoted the conversion Eu²⁺ → Eu³⁺, which was associated with changes in the optical basicity of the matrix and the amount of NBO. The ESR and luminescence results indicated that the addition of silica induced changes in the europium valence states, as well as modifications in the Eu²⁺ sites symmetry. For excitation at 350 nm, a broad emission band from the Eu²⁺ ions was observed, with a peak shift from 585 to 480 nm, resulting in the color of the emitted light changing from reddish-orange to blue. This shift to higher energies occurred for both emission and excitation bands. The present observations indicate that these Eu-doped glasses can be tailored for tunable white lighting by combining adjustment of the matrix composition, melting environment, and pumping wavelength.

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