RESEARCH ARTICLE

Sol-gel Synthesis and Characterization of Non-Substituted and Europium-Substituted Layered Double Hydroxides $\text{Mg}_3/\text{Al}_{1-x}\text{Eu}_x$

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Abstract: The Mg/Al layered double hydroxides (LDHs) were successfully synthesized by novel aqueous sol-gel method. A series of europium-substituted layered double hydroxides (Mg/Al/Eu LDHs) were also synthesized using the same sol-gel processing. The Eu^{3+} substitution effects were investigated in the $\text{Mg}_3/\text{Al}_{1-x}\text{Eu}_x$ system by changing the amount of Eu^{3+} in the crystal lattice from 0.05 to 10 mol%. The samples obtained were characterized by X-ray diffraction (XRD) analysis, Fourier Transform Infrared spectroscopy (FT-IR), thermogravimetric (TG) analysis, scanning electron microscopy (SEM) and fluorescence spectroscopy (FLS). The results revealed that europium enters into a hydrotalcite structure containing Mg^{2+} and Al^{3+} cations in the brucite-like layers.

Keywords: Layered double hydroxides, Mg/Al/Eu, sol-gel processing, europium substitution effects, luminescent properties.

1. INTRODUCTION

Layered double hydroxides (LDHs) are compounds composed of positively charged brucite-like layers with an interlayer gallery containing charged compensating anions and water molecules. The metal cations occupy the centres of shared octahedral whose vertices contain hydroxide ions that connect to form infinite two-dimensional sheets [1-4]. With this structure the most known mineral is hydrotalcite, $[\text{Mg}_{0.75}\text{Al}_{0.25}(\text{OH})_2] (\text{CO}_3)_{0.125} \cdot 0.5\text{H}_2\text{O}$. LDHs have a well-defined layered structure within nanometre scale (0.3-3 nm) of interlayer and contain important functional groups. After calcination at temperatures from 300 to 600 °C, LDHs were converted to mixed metal oxides (MMO), which have the ability to recover the original layered structure [5-7]. LDHs have unique intrinsic advantages: their layer sizes are highly tunable and their constituents can be conveniently controlled and varied. LDHs, unlike common clay minerals such as montmorillonite, cannot swell automatically in water, and their delamination is much more difficult due to their high surface charge density as well as the strong attractions between adjacent nanosheets [8]. The formation and exploitation of new types of layered double hydroxide (LDH)/polymer NC hydrogels with high performance has also been investigated [9].

One of the most important properties is the interlayer anion exchangeability, which endows LDHs the possibility of accommodating organic UV absorbers in the interlayer space [10]. Moreover, the LDHs have an HCl absorption capacity, and may be used as PVC thermal stabilizer [11].

In the positively charged brucite-like layers of LDHs, a fraction of the trivalent cations could be substituted by other same valence cations. Rare earth doped luminescent materials have drawn increasing attention as potential phosphor materials for use in optical devices [12-14].

Recently, considerable attention has been focused on incorporating rare earth ions into LDHs hosts to develop new functional materials, which resemble designed optical properties [15]. LDHs doped with Tb^{3+} ions in the brucite-like layers were prepared by a simple one-step co-precipitation method [16, 17]. Nanosized LDHs doped with Eu^{3+} , Yb^{3+} , Tb^{3+} and Nd^{3+} were prepared through the microemulsion method [18]. The Eu^{3+} and Nd^{3+} were incorporated also into hydrocalumite and mayenite [19]. The Zn/Al/Eu and Zn/Al/Dy LDHs were also reported as perspective and efficient luminescent materials [20-22]. Cerium-doped hydrotalcite-like precursors were also recently synthesized by co-precipitation method [23].

LDHs can be prepared by a variety of synthesis methods. However, the most common preparation technique is co-precipitation method starting from soluble salts of the metals. Sol-gel synthesis route for mixed metal oxides and related

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compounds have some benefits over other methods such as simplicity, synthesis at low temperatures, effectiveness, suitability for different systems and cost efficiency [24-27].

The main aim of this study was to investigate Eu^{3+} substitution effects in the $\text{Mg}/\text{Al}_{1-x}\text{Eu}_x$ systems (the Eu^{3+} concentration in the crystal lattice was changed from 0.05 to 10 mol%) fabricated for the first time to the best of our knowledge by sol-gel synthesis route. The luminescent properties of $\text{Mg}_3/\text{Al}_{1-x}\text{Eu}_x$ LDH samples were also investigated.

2. MATERIALS AND METHOD

Aluminium (III) nitrate nonahydrate ($\text{Al}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$), 98.5% (Hempur); magnesium (II) nitrate hexahydrate ($\text{Mg}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$), 99% (Hempur); europium (III) nitrate pentahydrate ($\text{Eu}(\text{NO}_3)_3 \cdot 5\text{H}_2\text{O}$, 99.9% (Sigma-Aldrich); ethylene glycol ($\text{C}_2\text{H}_6\text{O}_2$), 99.5% (Roth); citric acid ($\text{C}_6\text{H}_8\text{O}_7$), 99.5% (Hempur).

X-ray diffraction patterns were recorded using MiniFlex II diffractometer (Rigaku) using a primary beam $\text{Cu K}\alpha$ radiation ($\lambda = 1.541838 \text{ \AA}$). The 2θ angle of the diffractometer was graduated from 8 to 80° in steps of 0.02° , with the measuring time of 0.4 s per step. FT-IR analysis of compounds was conducted using Perkin-Elmer FTIR spectrometer. All spectra were recorded at ambient temperature in the range of $4000\text{--}400 \text{ cm}^{-1}$. Thermal analysis was carried out using a simultaneous thermal analyzer 6000 (Perkin-Elmer) in air atmosphere at scan rate of $10^\circ\text{C}/\text{min}$ and the temperature range from 30°C up to 900°C . Excitation and emission spectra were recorded on an Edinburg Instruments FLS 900 spectrometer. The morphology of particles was investigated using a scanning electron microscope (SEM) Hitachi SU-70.

3. EXPERIMENTAL:

The $\text{Mg}_3/\text{Al}_{1-x}\text{Eu}_x$ LDH samples were synthesised by the sol-gel method from the solution of metal nitrates $\text{Mg}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$, $\text{Al}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$ and $\text{Eu}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, dissolved in 50 ml of distilled water. Secondly, 0.2 M citric acid was added and obtained solution was stirred for 1 h at 80°C . Next, 2 ml of ethylene glycol have been added to the resulted mixture with continued stirring at 150°C until the complete evaporation of solvent. The obtained gel was dried at 105°C for 24 h . The mixed metal oxides were obtained by heating the gels at 650°C for 4 h . The $\text{Mg}_3/\text{Al}_{1-x}\text{Eu}_x$ LDH specimens were obtained by reconstruction of MMO powders in water at 50°C for 6 h under stirring. For comparison, the Mg/Al hydrotalcite was also synthesized and analysed.

4. RESULTS AND DISCUSSIONS:

The synthesized Mg/Al and $\text{Mg}/\text{Al}/\text{Eu}$ LDHs were characterized by XRD analysis (Figs. 1-3). Evidently, only amorphous $\text{Mg}-\text{Al}-\text{O}$ gel has formed during the initial stage of sol-gel processing of LDHs (Fig. 1). After heat-treatment of $\text{Mg}-\text{Al}-\text{O}$ precursor gel at 650°C , a high crystalline mixed-metal oxide (MMO) has formed. The obtained MMO powdered samples were treated in water at 50°C for 6 h under stirring. The XRD patterns of obtained specimens confirmed the formation of monophasic Mg/Al LDH during the “reconstruction” process in water. Three characteristic peaks of hydrotalcite at 2θ of about 10° (003), 23° (006) and 35° (009) were observed in sol-gel derived LDH samples. Moreover, two very important reflections for LDHs are also clear-

ly seen at about 60.2° (110) and 61.5° (113). Based on the XRD pattern, there are slight shifts of characteristic basal plane d (003) related to the existing of different anions in the interlayer regions. The carbonate acts as a balancing anion because of its high affinity in the hydroxide layer over nitrate anions. In the simplest case, anion exchange is achieved by stirring an aqueous solution of the LDH precursor with a large excess of the anion salt. LDHs containing carbonate anions within the host material are less susceptible to anion exchange reactions due to high selectivity for this anion.

Fig. (1). XRD patterns of $\text{Mg}-\text{Al}-\text{O}$ precursor gel, mixed-metal oxide obtained at 650°C and sol-gel derived Mg/Al LDH. The crystalline phases are marked: MgO - • and hydrotalcite - ♦.

All synthesized europium-substituted (1–10 mol%) $\text{Mg}-\text{Al}-\text{O}$ gels also were amorphous powders to compare with Mg/Al LDH (Fig. 1). According to the XRD patterns presented in Fig. 2 the heat-treated $\text{Mg}/\text{Al}/\text{Eu}$ gels resulted in high crystalline MMO samples. The mixed-metal oxides transformed fully to LDH structures independent on the Eu substitutional level (see Fig. 3). The observed shift of the (110) reflections displacement signal toward lower values of 2θ containing different concentration of Eu^{3+} can be an evidence of isomorphic incorporation of lanthanide ion in LDH. The calculated d values for (110) peak (for $1 \text{ mol\% Eu}^{3+}$ $1.5288 \text{ \AA}$; $5 \text{ mol\% Eu}^{3+}$ $1.5302 \text{ \AA}$; $7.5 \text{ mol\% Eu}^{3+}$ $1.5330 \text{ \AA}$ and $10 \text{ mol\% Eu}^{3+}$ $1.5359 \text{ \AA}$), compared with non-doped Mg/Al $d(110) = 1.5274 \text{ \AA}$ monotonically increased by increasing the Eu^{3+} concentration. The monophasic $\text{Mg}/\text{Al}/\text{Eu}$ LDHs were obtained with amount of Eu less than $5 \text{ mol\%}$. With increasing concentration of europium till $7.5 \text{ mol\%}$ the negligible amount of side $\text{Eu}(\text{OH})_3$ phase has formed. The formation of $\text{Eu}(\text{OH})_3$ is mainly due to the larger ionic radius of Eu^{3+} ions ($1.08 \text{ \AA}$) in comparison of ionic radius of Al^{3+} ions ($0.53 \text{ \AA}$). The Eu^{3+} ions taking up the octahedral positions of Al^{3+} ions led to the deformation of crystal lattice, and the excess Eu^{3+} ions in high basic condition crystallized as $\text{Eu}(\text{OH})_3$.

Fig. (2). XRD patterns of Mg/Al/Eu 1–10 mol% gels calcined at 650 °C. The crystalline phase is marked: MgO - •.

Fig. (3). XRD patterns of synthesized by sol-gel method Mg/Al/Eu 1 – 10 mol% LDHs. The $\text{Eu}(\text{OH})_3$ phase is marked: *.

Thus, these results confirmed once again that highly crystalline LDHs could be synthesized during hydroxylation sol-gel derived crystalline MMO samples in aqueous media. At high temperature of calcination, crystalline mixed oxide and partial changes result in the local structure of the M_{III} cations. A partial regeneration of the layered phase takes place which is characterized by different composition and morphology.

The XRD patterns of the synthesized LDHs contain very sharp diffraction lines attributable only to LDH crystalline phase at low concentration of europium. Interestingly, the formation of sol-gel based LDHs depends on the concentration of europium in the samples.

FT-IR spectra of Mg/Al and Mg/Al/Eu LDH samples in the region of 4000–500 cm^{-1} are shown in Fig. 4. The spectra recorded for all samples are very similar to each other without any important differences. The absorption bands observed at around 3500–3000 cm^{-1} correspond to the stretching vibrations of hydroxyl ($-\text{OH}$) groups from the hydroxyl layers and from intercalated and externally adsorbed water molecules [28]. The broad band identified at 2974 cm^{-1} is assigned to the stretching mode of hydroxyl groups hydrogen-bonded to interlayer carbonate anions. The presence of adsorbed molecular water was confirmed by the medium intensity band at 1629 cm^{-1} . The strong absorption band visible at 1353 cm^{-1} is attributed to the asymmetric vibrations modes of CO_3^{2-} . The absorption bands observed at 613 cm^{-1} can be ascribed to the metal-oxygen $\text{M}^{2+}-\text{O}$ and $\text{M}^{3+}-\text{O}$ stretching mode.

Fig. (4). FT-IR spectra of Mg/Al and Mg/Al/Eu 1 – 10 mol% LDHs.

The TG-DCS curves of synthesised Mg/Al and Mg/Al/Eu 10mol% LDHs are shown in Figs. 5 and 6, respectively. As seen, the thermal behaviour of synthesized LDH is not dependent on the europium substitution in the structure. The initial mass loss about ~15% was observed in the temperature range of 30–200 °C for the Mg/Al and Mg/Al/Eu LDHs prepared by sol-gel method. These changes in mass are due to the collapse of interlayer and evolution of adsorbed water. The main decomposition of LDH samples occurs via one step in the temperature range of 200–600 °C (the mass loss ~45–48%). The second mass loss step is due to the dihydroxylation and decomposition of the impurities of interlayer ionic carbonate. The observed thermal effects in DSC

curves support the results obtained from TG measurements. Above 600 °C the brucite-type structure collapses and a solid solution of mixed spinel (MgAl_2O_4) and MgO , or Al_2O_3 and MgO is formed.

Fig. (5). TG-DSC curves of the Mg/Al LDH.

Fig. (6). TG-DSC curves of the Mg/Al/Eu 10 mol% LDH.

The microstructure for the synthesized LDH samples was determined from the SEM micrographs. The SEM micrographs of calcined Mg/Al precursor and sol-gel derived LDH sample are shown in Fig. 7. The mixed metal oxide obtained by calcination of Mg-Al-O gel at 650 °C is composed of the network of spherical nanoparticles varying in size from approximately 2 to 5 μm . A layered double structure was recovered after reconstruction procedure in water, showing the formation of plate-like particles with hexagonal shape (Fig. 7b). The surface morphology of Eu^{3+} -substituted LDH samples is very similar for all specimens independent of the substitutional level. Fig. 8 shows the representative SEM micrographs of sol-gel synthesized Mg/Al/Eu 1mol% and Mg/Al/Eu 10mol% samples. As seen, small fibrous plate-like

particles with hexagonal structure are aggregated as in the case of non-substituted LDH sample. So, all sol-gel derived Mg/Al/Eu LDHs consist of the larger hexagonally shaped particles varying in size from approximately 250 to 500 nm. The surface morphological features of sample with highest europium content are more pronounced consisting of hexagonally shaped nanostructures in LDHs. Finally, the complete regeneration LDHs samples from MMO in aqueous media has been observed.

Fig. (7). SEM micrographs of MMO (a) and sol-gel synthesized Mg/Al LDH (b).

The emission spectra obtained at room temperature of all the Mg/Al/Eu LDH samples under excitation at 320 nm are presented in Fig. 9. As seen, all samples share similar emission profiles. The emission spectra of Mg/Al/Eu LDHs shows three main emissions in the wavelength range of 500–740 nm. In all spectra, the emission bands characteristic for $^5\text{D}_0 \rightarrow ^7\text{F}_j$ ($j=1, 2, 3, 4$) transitions of Eu^{3+} ions were observed. The emission peaks are referred to the typical three $^5\text{D}_0 \rightarrow ^7\text{F}_1$ (591 nm) $^5\text{D}_0 \rightarrow ^7\text{F}_2$ (615 nm) and $^5\text{D}_0 \rightarrow ^7\text{F}_4$ (703) transitions of Eu^{3+} ion. The emission due to $^5\text{D}_0 \rightarrow ^7\text{F}_2$ transition is the strongest, indicating that Eu^{3+} ions occupy a low-symmetry

site. $^5D_0 \rightarrow ^7F_2$ transition called as a hypersensitive transition. This can be caused by the addition of another molecule (for example, a competitive ligand) by a variation of the ionic strength of the solution (addition of a salt), by a variation of the polarity of the solvent (addition of another solvent), or by a variation of the pH value of the solution (addition of an acid or a base) when working in aqueous solutions. The presence of the low-symmetry structure containing the Eu^{3+} can only be observed in the nanocrystal line products, the results allow to exclude the adsorption of Eu^{3+} on the surface of LDHs [29]. When Eu^{3+} ions occupy the sites with inversion symmetry, the $^5D_0 \rightarrow ^7F_1$ transition, typical magnetic dipole transition, should be relatively dominant; while, if there is no inversion symmetry at the sites of Eu^{3+} ions, $^5D_0 \rightarrow ^7F_2$ transition should be relatively dominant [30]. It is clear that the photoluminescence intensity in the Mg/Al/Eu LDH phase increases with increasing the Eu^{3+} concentration, and reaches the maximum when the concentration of Eu^{3+} is 7.5%. With further increasing amount of europium the intensity of emission decreases due to the concentration quenching. As the concentration of Eu^{3+} increases, the distances between Eu^{3+} ions in the layers will be shortened, which dramatically increases the interaction of ions and causes serious concentration quenching. This fact suggested that Eu^{3+} ions replaced Al^{3+} in the host lattice.

Fig. (8). SEM micrographs of Mg/Al/Eu 1mol% (a) and Mg/Al/Eu 10mol% (b) LDHs.

Fig. (9). Photoluminescence emission spectra of Mg/Al/ Eu^{3+} LDHs ($\lambda_{\text{ex}} = 320$ nm).

CONCLUSION

The Mg/Al/ layered double hydroxides (LDHs) were successfully synthesized by novel aqueous sol-gel method. In the sol-gel processing route, the LDHs were obtained after the decomposition of Mg-Al-O precursor gels and following “reconstruction” of intermediate crystalline MMO samples in aqueous media. The same synthesis parameters were successfully applied for the synthesis of europium-substituted Mg/Al/ Eu^{3+} 0.05–10 mol% LDHs. Interestingly, the formation of sol-gel based Mg/Al/ Eu^{3+} LDHs depends slightly on the concentration of europium used in the samples. With increasing the concentration of europium till 7.5 mol% the negligible amount of side $\text{Eu}(\text{OH})_3$ phase has formed. The typical LDH microstructure was determined from the SEM micrographs. The sol-gel synthesized LDH samples consisted of hexagonally shaped particles varying in size from approximately 250 to 500 nm. The luminescence of Mg/Al/ Eu^{3+} LDHs was also investigated in this study. The emission peaks are referred to the typical three $^5D_0 \rightarrow ^7F_1$ (591 nm) and $^5D_0 \rightarrow ^7F_2$ (615 nm) $^5D_0 \rightarrow ^7F_4$ (703 nm) transitions of Eu^{3+} ion. It was also suggested from these results, that Eu^{3+} replaced Al^{3+} ions in the host lattice.

CONFLICT OF INTEREST

The authors confirm that this article content has no conflict of interest.

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