Effective Doping of Rare-earth Ions in Silica Gel: A Novel Approach to Design Active Electronic Devices

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Abstract: Eu$^{3+}$ luminescence spectroscopy has been used to investigate the effective doping of alkoxide-based silica (SiO$_2$) gels using a novel pressure-assisted sol-gel method. Our results pertaining to intense photoluminescence (PL) from gel nanospheres can be directly attributed to the high specific surface area and remarkable decrease in unsaturated dangling bonds of the gel nanospheres under pressure. An increased dehydroxylation in an autoclave resulted in enhanced red (∼611 nm) PL emission from europium and is almost ten times brighter than the SiO$_2$ gel made at atmospheric pressure and ∼50°C using conventional Stöber-Fink-Bohn process. The presented results are entirely different from those reported earlier for SiO$_2$:Eu$^{3+}$ gel nanospheres and the origin of the enhanced PL have been discussed thoroughly.

Keywords: Luminescence; Nanophosphor; Short-range order; Electron microscopy

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Introduction

Luminescent materials have been utilized widely in applications involving lighting to sensing [1]. The photoluminescence (PL) properties of silica have also been an important topic of research for a long time, but the difficulty in the incorporation of rare-earth (RE) ions attached covalently to the silica (SiO$_2$) network is still considered a great challenge. The weak PL bands with peak energies ∼1.9-4.3 eV for both bulk and thin films of SiO$_2$ have been reported [2]. The sol-gel method has been used to prepare nanomaterials in the form of powders, films, fibers and monoliths that are based on various metal alkoxides [3]. It has been observed that monodisperse silica nanospheres formed by hydrolysis and condensation of alkoxides using Stöber-Fink-Bohn (SFB) process gives negligible luminescence. Incorporation of inorganic luminescent centres into SFB spheres has been demonstrated by some research groups [4] using RE ions, quantum dots and organic fluorophores, but the procedure requires multiple processing steps and use of expensive and toxic ligands. During the last few years, there have been few reports [5-10] on intense PL emission in the visible region of the electromagnetic spectrum, by several different nanostructured materials that are highly disordered such as nanowires, porous silicon, silica-based mesoporous fractals, ferroelectrics with ABO$_3$ type and AWO$_4$-type perovskite structures (A=Ca, Ba, Sr); and many more. The origin of this type of luminescence is always attributed to unsaturated chemical bonds in these nanostructures. In case of ferroelectrics, as their energy band gaps are located ∼3-4 eV, the PL emission in the visible region was correlated to their highly disordered states and many localized electronic levels present within the optical gap, causing luminescence [6]. Since SiO$_2$ gel is a well-studied system [11] having visible transparency and significant absorption peak lying in the ultra-violet...
region, the introduction of dopant ions act as a perturbation to the well-studied system leading to interesting optical properties. The RE ions are used as probes in the sol-gel method due to their sensibility to change with the surrounding matrix probing the local structure [12,13]. Thus derived SiO$_2$ and organically modified matrix composites could be the main precursor to prepare many RE based smart optical materials [14].

In this paper, we propose a novel methodology to prepare alkoxide-based silica gel nanospheres doped with Eu$^{3+}$ ions that show enhanced PL brightness, uniform size distribution and improved quantum efficiencies. This is a process by which highly disordered but doped silica gels could be effectively made useful for practical applications involving luminescence. It has already been evidenced that high pressure and temperature leads to more closely packed structures [14]. The present analogy is unique and could be extended to many crystalline and non-crystalline phosphor based systems to design new family of sol-gel based nanocomposites, having a wide variety of applications in lasers, chemical sensing, waveguides, bioanalytical assays, blood flow monitoring and effectively harvesting the solar energy for improving the solar cell efficiency.

**Experimental**

Silica nanospheres were prepared with its surface modified by sol-gel method and studied through the incorporation of Europium (III) ions in two different ways. The europium nitrate (EuNO$_3$) was mixed with sufficient ethanol (EtOH) and this was added to tetraethylorthosilicate (TEOS). The stoichiometric amount of water, which is essential to carryout hydrolysis reaction, was added drop wise under continuous stirring. Prior to gelling the low and high resolution transmission electron microscopy (TEM) images of colloidal silica solution (also called silica sol) were taken at a magnification of 125 and 400 kX, respectively as shown in Fig. 1. The silica particles observed are almost spherical with an average cluster size of $\sim$5 nm. The schematic of the SiO$_2$ gel network [11] and the probable description of pore and particle sizes were illustrated in Fig.1(c). The silica sol was then divided in two halves for a systematic experimentation. For the first experiment, one of the solution containing vials was allowed to gel at atmospheric pressure (1 bar) and a controlled oven temperature of $\sim$50°C ($\pm$0.1°C). In the second case the solution was kept in an autoclave, subjected to high temperature and pressure at $\sim$150°C and 120 bars, respectively for about 5 hours. It was intentional to keep the processing time same for both the cases. Once the wet silica gels were obtained, SiO$_2$:Eu$^{3+}$ nanospheric powders were obtained by drying the gels overnight in vacuum oven at $\sim$55°C. All the samples were studied using morphology, composition evaluation, UV-VIS absorption and luminescence spectroscopy techniques.

For TEM observations, the samples were redispersed in methanol by ultrasonic treatment and dropped on carbon-copper grids. TEM images were collected using a Tecnai G$^2$ F30 S-Twin (FEI; Super Twin lens with C$_S$=1.2 mm) instrument operating at an accelerating voltage at 300 kV, having a point resolution of 0.2 nm and lattice resolution of 0.14 nm) with an EDAX attachment. Program Digital Micrograph (Gatan) was used for image processing. Scanning electron microscopy was performed using Zeiss EVO MA10.

X-ray diffraction (XRD) of the powder sample was performed using a Tecnai G$^2$ F30 S-Twin (FEI; Super Twin lens with C$_S$=1.2 mm) instrument operating at an accelerating voltage at 300 kV, having a point resolution of 0.2 nm and lattice resolution of 0.14 nm) with an EDAX attachment. Program Digital Micrograph (Gatan) was used for image processing. Scanning electron microscopy was performed using Zeiss EVO MA10.

X-ray Photoelectron Spectroscopy (XPS) studies have been carried out using a Perkin Elmer 1257 model, at 300 K with a non-monochromatic AlK$_\alpha$ line at 1486.6 eV. During photoemission studies, small specimen charging was observed which was later calibrated by assigning the C 1s signal at 285 eV.

The room temperature photoluminescence (PL) spectra were recorded using an Edinburgh Luminescence Spectrometer (Model F900) equipped with a xenon lamp. The excitation and emission spectra were recorded in the fluorescence mode over the range of 300-700 nm.

**Results and discussion**

It is important to highlight the pressure-assisted hydrothermal/solvothermal process for the preparation of variety of nanoparticles of oxides and chalcogenide materials. The solution based nanomaterial synthesis often involves reactions carried out near the boiling point.
of the solvent. This may lead to poor quality of the nanomaterial and less yield. In order to obtain crystalline, monodisperse nanoparticles, it is always necessary to work at relatively high temperatures and pressures. Use of autoclave synthesis of intrinsic silica gels have been reported extensively by Haranath et al. since 1996. In the current case, rare-earth (RE) doped silica gels were made under pressure at elevated temperatures as described in latter sections. This method of preparation, which is based on pressure-assisted sol-gel method, has compatibility in modifying the coordinating environment of RE (dopant) ions so that the loss in energy of the excited states via non-radiative mechanism is minimum. The x-ray diffraction pattern of Eu$^{3+}$ doped SiO$_2$ gel powder depicted in Fig. 2 clearly shows a broad hump at $\sim$23$^\circ$ indicating the complete amorphous nature of the SiO$_2$ nanospheres. SEM images shown in inset of Fig. 2 illustrate the quality of the nanospheres with respect to their size and shapes. In other words, the conventional sol-gel method leads to a broad distribution of particle sizes in the range 10-50 nm (Fig. 2(a)) whereas the pressure assisted sol-gel method has resulted in almost uniform particles ($\sim$15 nm) with spherical shape (Fig. 2(b)). This establishes the fact that high pressure and temperature leads to more closely packed structures and increased charge transfer energies that are efficiently transferred to the Eu$^{3+}$ ions. The TEM and scanning electron micrographs (SEM) reveal that the optimization of experimental and processing parameters allow microscopic preparation of uniform silica nanospheres.

Moreover, the chemical composition and the relevant surface chemistry of SiO$_2$:Eu$^{3+}$ nanospheres were analyzed using x-ray photoelectron spectroscopy (XPS). The XPS observations revealed three peaks corresponding to the elements Si, O and Eu, respectively. Pass energy for general survey scan and core level spectra was kept at 143.05 and 71.55 eV respectively. Surface contamination was removed by Ar ions with 4 keV beam energy. Sputtering performed in raster mode with emission current of 20 mA for 5 min at base pressure $4.5 \times 10^{-7}$ torr. Figure 3 shows the survey spectra of SiO$_2$:Eu$^{3+}$ nanospheres acquired in the range of 0-1200 eV. During photoemission studies, small specimen charging was observed, which was later calibrated by assigning the C 1s signal at 285 eV. Survey spectra after sputtering show sharp peaks of C 1s (285 eV), O 1s (537 eV). Two clear distinct state of Eu were observed at 1171 and 1141 eV for Eu 3d$^5_{3/2}$ and 3d$^5_{5/2}$ respectively. The separation between two states is because of spin-orbit splitting. Inset of Fig. 3 shows the Si(2p) core level spectra. The value of elemental Si(2p) is 99.15 eV, since, appearance of Si(2p) at 104 eV confirms the Si exist in SiO$_2$ state. The binding energies of various elements match very well with the peaks observed for standard SiO$_2$:Eu$^{3+}$. The presence of C is due to the air atmosphere and the organics used for the preparation of Eu$^{3+}$ doped in silica matrix.

Figure 4 shows the UV-VIS absorbance spectra of Eu$^{3+}$ doped SiO$_2$ gel samples prepared at 50°C, 1 bar; and 150°C, 120 bars, respectively. The spectra are the strong indicative two findings. One is being the reduction of surface states of the gel nanospheres and other being the maintenance of almost the same size for the gel particles even after subjecting to high pressure and temperature cycles. The same has been evidenced by the absorption peaks indicated in the inset of Fig. 4.

![Fig. 2 X-ray diffraction (XRD) pattern of SiO$_2$ gel powder at room temperature ($\sim$25°C) indicating the amorphous nature of the gel under study. The insets shows the SEM micrographs of SiO$_2$:Eu gels prepared at (a) atmosphere pressure (1 bar) and 50°C; and (b) 120 bars and 150°C, respectively.](http://dx.doi.org/10.3786/nml.v3i3.p141-145)

![Fig. 3 XPS survey scan spectra of sputtered and nanoparticles of SiO$_2$:Eu$^{3+}$ recorded with photon energy of AlK$_\alpha$ $(h\nu=1486.6$ eV), Inset shows the Si(2p) core level.](http://dx.doi.org/10.3786/nml.v3i3.p141-145)
The final product. The most intense line at a remarkable decrease in unsatisfied chemical bonds in clave treated gel sample, which indicate that there is the PLE spectrum also became narrower under auto-
and pressures (120 bars) inside an autoclave. Moreover,
an asymmetry parameter for the Eu
3+
ion corresponds to the hypersensitive transition between face states and unsaturated dangling bonds play a crit-
ical role in determining the overall PL characteristics
of nanostructures. If the silica gel is prepared with a
rare-earth dopant (Eu
3+) as shown in the
inset of Fig. 5. The emission spectra of Eu
3+
doped SiO
2
gels prepared at 50
°C, 1 bar; and 150
°C, 120 bars,
were shown in Fig. 5. For recording the PL spectra
the excitation energy has been fixed as 395 nm which
is the
5D0 level of Eu
3+ ligand band. The PL from the
Eu
3+ doped SiO
2 gel prepared at atmospheric pressure (1 bar) and 50
°C was found to be weak and inefficient for any practical application, whereas the PL intensity from the gel turn out to be much stronger (>10 times) when the sol was gelled under high temperature (150
°C) and pressures (120 bars) inside an autoclave. Moreover, the PLE spectrum also became narrower under auto-
clave treated gel sample, which indicate that there is a remarkable decrease in unsatisfied chemical bonds in the final product. The most intense line at ∼611 nm corresponds to the hypersensitive transition between the
5D0 and
7F2 level of the Eu
3+ ions and will be relatively strong if the surrounding symmetry is low. In the sense, it is generally admitted that the ratio of the emission intensities R=I(6D0 →7F2)/I(6D0 →7F1) is an asymmetry parameter for the Eu
3+ sites and a measure of the extent of its interaction with the surrounding ligands [20]. This indicates that the environment of the Eu
3+ is dictated by the nano-SiO
2 host under high pressure and temperature conditions. In addition, broad excitation peaks observed between 250-500 nm became distinct and sharp for the sample made under high pressures. The sharp peaks in the PL and PLE spectra of SiO
2:Eu nanospheres may be due to quantum confinement effects related to size restrictions. A combination of unique features of high surface-to-volume ratios, monodispersion and strong photoluminescence suggest that these silica nanospheres will find many inter-
esting applications in semiconductor photophysics, inorganic light emitting diodes, solar cells, environmental
remediations and optoelectronic devices.

**Fig. 4** Absorbance spectra of Eu doped-SiO
2 samples pre-
pared at atmosphere pressure (1 bar) and 120 bars and
150
°C, respectively. Inset shows the expanded version of the absorption peaks.

The observation of weak photoluminescence (PL) at
room temperature in various amorphous silica nano-
structures has been reported in the literature [15-18] but
not sufficient to use for any fundamental or potential
application. Figure 5 shows the PL excitation (PLE)
and PL spectra from Eu
3+ doped and baked SiO
2 gel
in an autoclave respectively. It is known that the sur-
face states and unsaturated dangling bonds play a crit-
ical role in determining the overall PL characteristics
of nanostructures. If the silica gel is prepared with a
rare-earth dopant (Eu
3+ in the present case), the PL
is dominated by the radiative transitions (5D0 →7F
j, j=0-3) from the levels of Eu
3+ ions [19] as shown in the
inset of Fig. 5. The emission spectra of Eu
3+ doped
SiO
2 gels prepared at 50
°C, 1 bar; and 150
°C, 120 bars,
were shown in Fig. 5. For recording the PL spectra
the excitation energy has been fixed as 395 nm which
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5D0 level of Eu
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clave treated gel sample, which indicate that there is a remarkable decrease in unsatisfied chemical bonds in the final product. The most intense line at ∼611 nm corresponds to the hypersensitive transition between the
5D0 and
7F2 level of the Eu
3+ ions and will be relatively strong if the surrounding symmetry is low. In the sense, it is generally admitted that the ratio of the emission intensities R=I(6D0 →7F2)/I(6D0 →7F1) is an asymmetry parameter for the Eu
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esting applications in semiconductor photophysics, inorganic light emitting diodes, solar cells, environmental
remediations and optoelectronic devices.

**Fig. 5** PLE and PL spectra of SiO
2:Eu gels made at (a) 50
°C, 1 bar and (b) 120 bars and 150
°C, respectively. Inset shows the mechanism of Eu
3+ transitions.

**Conclusions**

In conclusion, we have demonstrated a mechanism by
which photoluminescence enhancement in SiO
2:Eu
3+
phosphor nanospheres could be successfully achieved
using pressure-assisted sol-gel method. The observed red emission is from typical
5D0−7F2 transition of Eu
3+ and is found to be almost ten times brighter than the gel made at atmospheric pressure (1 bar) and ∼50
°C using Stöber-Fink-Bohn process. This kind of process is highly desirable for many crystalline and non-
crystalline materials system wherein the doping is inap-
propriate. This may lead to design many fundamental and novel optoelectronic applications.

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