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Introduction

Nephelauxetic effect of the hydride ligand in Sr₂LiSiO₄H as a host material for rare-earthactivated phosphors†

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Strontium lithium orthosilicate hydride Sr_2LiSiO_4H was synthesized by the reaction of Sr_2SiO_4 with LiH at 700 °C in a H₂ rich atmosphere. Rietveld refinement of the neutron powder diffraction pattern revealed that Sr_2LiSiO_4H is isostructural to Sr_2LiSiO_4F (space group $P2_1/m$) and its channel-like structure preferentially accommodates H⁻ ions over F⁻ ions. In addition, Sr_2LiSiO_4H is stable in air and its Eu²⁺-doped analog exhibits yellow photoluminescence with an emission band at 544 nm and a broad excitation band ranging from 250 to 450 nm. These bands were observed in the longer wavelength region when compared with those displayed by $Sr_2LiSiO_4F:Eu^{2+}$. The red shift, which is induced by H⁻ substitution, is consistent with the constrained density functional theory calculations, predicting the photo-excitation and emission energies of 4f–5d transitions. The present study reports the synthesis of stable oxyhydrides acting as phosphor hosts for rare earth ions. The phosphor hosts exhibit large nephelauxetic effects owing to the presence of H⁻ ligands.

The hydride ion H^- is a monovalent anion composed of one proton and two electrons with no p orbitals in the outer shell.¹ The unique atomic structure of H^- strongly influences its chemical bonding.² Owing to its multivalent property and rather low electronegativity, which is lower than that of oxygen, hydrogen can work both as an electron donor and acceptor in oxide-based materials. The unique properties of hydrogen offer promising applications for oxyhydrides, in which both oxide and hydride ions are presented, toward energy storage in semiconducting as well as superconducting materials.³⁻⁷ However, knowledge surrounding oxyhydrides with exclusive properties is limited, thereby generating much interest in pursuing further exploration of such oxyhydride materials.

Over recent decades, lanthanide ion-doped compounds have been typically examined in phosphor-converted white lightemitting diodes. Examples of such compounds include YAG:Ce³⁺ and Eu²⁺-doped SiAION phosphors on blue GaInN chips.^{8,9} Among them, Eu²⁺-doped host phosphor materials have received the most attention owing to its parity-allowed electronic dipole 4f \rightarrow 5d transitions with comparable strong emitting intensities.¹⁰ Unlike the 4f electrons, the binding energy of the unshielded 5d electrons strongly depends on the crystalline environment, which can be described as the centroid shift and crystal field splitting of 5d levels that reflect on absorption energies.11 Therefore, to design phosphors with different emitting color requirements, oxides, nitrides, halides, and mixed ligand systems have been used as host materials.12 Compared to conventional host materials, hydride and hydridebased mixed anion compounds are expected to induce larger centroid shifts due to the expansion of electron cloud of the 5d electron. This so-called nephelauxetic effect is generally based on the covalent bonding between cation and anion ligand. Since hydrogen has a lower electronegativity (2.20 in Pauling scale) than those of oxygen (3.44), nitrogen (3.04), and fluorine (3.98),¹³ the stronger covalence bonding can be formed and then the large red shift of the excitation band is expected. For the past decades, hydrides and mixed anion hydrides containing Eu²⁺ dopant have been reported to exhibit green-red luminescence.10,14-20 However, their poor stability in air and moisture limits their application. In contrast, we have recently demonstrated that oxyhydride GdHO is stable in ambient air and can act as a phosphor host for Tb³⁺. Then, the green luminescence corresponding to ${}^{5}D_{4} \rightarrow {}^{7}F_{J}$ transition was observed upon excitation in near-UV light.²¹ Therefore, in the present study, we aim to evaluate stable oxyhydrides that can be used as phosphor hosts for Eu²⁺ with strong photoluminescence.

In the preliminary of this study, we investigated the conditions required to obtain oxyhydride-based phosphor from air-

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Paper

stable oxyhydrides. To avoid optical absorption in the visible region, which is typically due to the presence of the d electrons of transition metals,^{4,7,22} the host lattice of the phosphor should not contain transition-metal. Furthermore, hydride ions can be stably incorporated in transition-metal-free anion-encaging compounds with a cage and channel structure. Examples of compounds such include H-doped mayenite $[Ca_{12}Al_{14}O_{32}]^{2+}(O_{1-x}^{2-}H_{2x}^{-})$ and apatite $[Ca_{10}(PO_4)_6]^{2+}$ $(O_{1-x}^{2}X_{2x})$ (X = H, OH).^{3,6,23} Using this approach, we focused on strontium lithium orthosilicate fluoride Sr₂LiSiO₄F doped with Eu²⁺ and/or Ce³⁺. This material has shown potential as phosphor material in white light-emitting diodes with a broad emission range from blue to green.²⁴⁻²⁶ Due to the same valence and comparable ionic radii of H⁻ and F⁻,²⁷ we expected complete substitution of F⁻ in Sr₂LiSiO₄F with H⁻. In the hydride analog Sr₂LiSiO₄H, only H⁻ occupies the anion site in the channel space similarly to H-substituted apatite. In contrast, the anion sites in may enite and apatite are partially occupied by both O^{2-} and H^{-} . The former arrangement is advantageous to suppressing optical absorption by electrons in anion vacancies generated by photodissociation of H⁻ (O²⁻ + H⁻ \rightarrow OH⁻ + 2e⁻). Thus, Sr₂LiSiO₄H is expected to be stable in air and suitable as a host material for phosphor agents. As Eu²⁺ ion has an ionic radius similar to that of Sr²⁺ ion, Eu²⁺ ion can easily be incorporated into Sr sites. Because of aforementioned, Eu^{2+} is an appropriate dopant for investigating the influence of hydride substitution on the coordination structure and optical properties. In Sr₂LiSiO₄H:Eu²⁺, we expect a larger redshift in the Eu 5d energy levels, corresponding to smaller absorption energies, when compared with Sr₂LiSiO₄F:Eu²⁺.

In the present study, we report the synthesis of oxyhydride Sr₂LiSiO₄H by heating a mixture of LiH and Sr₂SiO₄ in H₂ gas and its photoluminescence (PL) property activated by partial substitution of Eu²⁺ for Sr²⁺. Neutron powder diffraction (NPD) measurements of the deuterium-enriched sample demonstrated the formation of a crystalline phase isostructural to Sr₂LiSiO₄F and full occupation of the F sites by deuterium. Since the synthesized crystalline phase is stable in ambient air, all measurements were taken in ambient environment. The Eu-free Sr_2LiSiO_4H sample, with an optical bandgap larger than 5.2 eV, is light grey under daylight. With Eu incorporation, the sample is light yellow and emits intense yellow luminescence under near-UV light (375 nm). To investigate the increase in centroid shift of the Eu 5d level upon substitution of F⁻ with H⁻, we performed constrained density functional theory (cDFT) calculations on the Eu²⁺-doped Sr_2LiSiO_4X systems (X = F, H).

Experimental section

Unlike most oxyhydrides that are synthesized by topochemical reaction of oxide precursors with a hydride ion source, polycrystalline $Sr_{2-x}Eu_xLiSiO_4H$ was directly synthesized by reaction of $Sr_{2-x}Eu_xSiO_4$ with LiH at a high temperature. LiH is a highly thermally stable hydride, with a melting point of 689 °C and a decomposition temperature of >900 °C under ambient pressure condition. The stability of LiH is considered to be the key factor for the direct synthesis of the oxyhydride Sr_2LiSiO_4H . The precursor compounds were prepared by solid-state reaction of SrCO₃, Eu₂O₃ and SiO₂ under the reduction gas environment (95% Ar/5% H₂) and hydrogenation of Li metal, respectively. The precursor compounds were then mixed and ground into fine powders with a molar ratio of 1 : 1.05 and heated up to 700 °C in H₂ gas at 0.9 MPa. For the powder X-ray diffraction (XRD) measurements and PL study, 2% Eu²⁺-doped sample *i.e.*, Sr_{1.96}Eu_{0.04}LiSiO₄H (Sr₂LiSiO₄H:Eu²⁺) was used. As for the NPD measurements, 100% deuterium-enriched sample (Sr₂LiSiO₄D) was synthesized using Sr₂SiO₄ and LiD. Detailed descriptions of the synthesis of the samples and characterization techniques including thermal desorption spectroscopy (TDS), XRD, NPD, diffuse reflectance spectroscopy, photoluminescence spectra (PL), magnetization measurements and DFT calculations are provided in the ESI.[†]

Result and discussion

The XRD patterns of Sr₂LiSiO₄H:Eu²⁺ measured with Cu Ka radiation, (Fig. S1(a) in ESI[†]) could be indexed to the space group $P2_1/m$, with lattice parameters a = 6.5854(2), b =5.4206(2), and c = 6.9458(2) Å, and $\beta = 112.568(2)^{\circ}$. These parameters are comparable with those of Sr_2LiSiO_4F (a = 6.5825(9), b = 5.4158(8), and c = 6.9266(6) Å, and $\beta =$ 112.525(8)°), implying the formation of isostructural compound accommodating H⁻ in place of F⁻. After exposing to ambient air for 16 h, no change was observed in the XRD pattern, indicating the air and moisture stability of Sr₂LiSiO₄H:Eu²⁺. Thus, subsequent measurements were taken in ambient environment. The refinement of the XRD pattern indicated the presence of small amount of Li₂SrSiO4 (1.36 wt%), which could not be reduced even after the optimization of synthesis temperature and atmospheric condition. To determine the occupancy of the hydrogen site, NPD and thermal desorption spectroscopy (TDS) measurements were respectively performed for Sr₂LiSiO₄D and Sr₂LiSiO₄H samples. Fig. 1 shows the NPD pattern of Sr₂LiSiO₄D collected on a neutron total scattering spectrometer (NOVA; beam-line BL21) at the Japan Proton Accelerator Research Complex (J-PARC), and Rietveld refinement was performed using the Z-Rietveld code.28 Except for the weak peaks originating from Sr₂LiSiO₄ (1.97(1) wt%) and SrO impurities (0.42(1) wt%), the major reflections could be indexed to the space group $P2_1/m$, with lattice parameters a = 6.5820(5), b =5.4197(4), and c = 6.9475(5) Å, and $\beta = 112.5628(2)^{\circ}$. Rietveld refinement of the NPD pattern was easily converged by using the Sr₂LiSiO₄F-type initial structure in which the fluorine atoms were fully replaced by deuterium atoms without noticeable vacancies. The hydrogen stoichiometry determined by TDS was $H/Sr_2LiSiO_4 = 0.97$, confirming complete occupation of fluoride sites in Sr₂LiSiO₄F by hydrogen without crystal structure transformation.

The structure model of Sr_2LiSiO_4H along the *b* axis is shown in the inset of Fig. 1, and the interatomic distances in Sr_2 - $LiSiO_4D$ (Sr_2LiSiO_4D data obtained from NPD) compared with those in Sr_2LiSiO_4F are listed in Table S6.[†] In this crystal, hydrogen or deuterium occupies the anion site that is octahedrally coordinated by four Sr and two Li atoms, forming a chain



Fig. 1 Time-of-flight NPD pattern of Sr₂LiSiO₄D and Rietveld fit (red line). The blue line represents the difference between the pattern and the fitted pattern. Orange, green, and purple markers represent the Bragg diffraction positions calculated for Sr₂LiSiO₄D, SrLi₂SiO₄, and SrO, respectively. Inset shows the structure model of Sr₂LiSiO₄H visualized by VESTA.⁴³

of face-sharing octahedra. The unit cell contains two types of 10coordinated Sr sites, 4-coordinated Si atoms, and 6-coordinated H atoms. The Sr1 site is surrounded by one O1, four O2, and three O3 atoms and two H1 sites, whereas the Sr2 site is surrounded by three O1, four O2, and one O3 atoms, and two H1 sites. This monoclinic crystal structure is composed of faceshared $(Sr_4Li_2H)^{4+}$ octahedra with isolated orthosilicate $(SiO_4)^{4-}$ groups. All oxygen ions are shared at the vertices of the SiO₄ tetrahedra. The H-centered octahedra provide onedimensional (1D) chains of hydride ions with a H-H distance of 2.73 Å. Since the substitution of F⁻ by H⁻ does not cause any structural changes, hydride ion is highly stable in this 1D channel structure and its presence offers chemical stability to the corresponding oxyhydride material against air and moisture. Because of their similarity in ionic radius, Sr sites will be substituted upon introduction of divalent europium ions into the system.

To examine the effect of F⁻ substitution by H⁻, DFT calculations were performed using the VASP code with projectoraugmented plane-wave method and Perdew-Burke-Ernzerhof functional (PBE) functional.²⁹⁻³¹ The calculated electronic band structure and density of state (DOS) are shown in Fig. 2. To emphasize the domination of hydride ions on the top of valence band, the energy axes were further adjusted by the O 2s orbital which located below Fermi level around 18-19 eV. The bandgaps of Sr₂LiSiO₄H and Sr₂LiSiO₄F estimated from the band structure, were 4.32 eV and 4.82 eV, respectively. However, according to the tendency that well-established in the literature, PBE calculations underestimate the actual bandgap energy.^{32,33} To obtain more realistic gap values, the calculations were performed using G₀W₀.³⁴ Indeed, we obtained bandgap values of 6.29 eV and 6.87 eV for Sr₂LiSiO₄H and Sr₂LiSiO₄F, respectively. For Sr₂LiSiO₄F, the conduction band minima are primarily composed of Sr 3d orbitals, whereas the valence bands maxima are composed of O 2p orbitals. The 2p orbitals of F⁻ ion have

lower energies and overlap on the O 2p orbitals, thus have no effect on the bandgap of Sr_2LiSiO_4F . In contrast, the smaller bandgap of Sr_2LiSiO_4H is attributed to the hydride ions. Since its the valence band is composed of hydride 1s orbital which has a higher energy than the O 2p orbitals, a narrower bandgap was observed.

The diffuse reflectance spectra of the Sr₂LiSiO₄H and Sr₂-LiSiO₄F samples were recorded at room temperature within the wavelength range of 200-1500 nm. The optical absorption spectra transformed from the relative diffuse reflectance R using the Kubelka–Munk function $F(R) = (1 - R)^2/2R$ are shown in Fig. 3.³⁵ For Sr₂LiSiO₄H, the strong optical absorption edge located around 250 nm could be associated with electron transitions from the valence band to the conduction band of the host lattice. This absorption edge was red-shifted relative to that of Sr_2LiSiO_4F (~230 nm). This result further confirmed the narrower bandgap of Sr₂LiSiO₄H when compared with that of Sr_2LiSiO_4F . More intuitive bandgap (E_{α}) values can be calculated using the following equation:³⁶ $F(R)h\nu \propto (h\nu - E_{\sigma})^{n/2}$, where $h\nu$ is the photon energy and the value of *n* is dictated by the type of transition (n = 4 for indirect transition and n = 1 for direct transition). In the present study, $(F(R)h\nu)^2 - h\nu$ plots were used to determine E_{g} , as shown in the inset of Fig. 3. This relationship gave a better linearity than the $(F(R)h\nu)^{1/2}-h\nu$ relationship (Fig. S2[†]). By extrapolating the former plot to 0, the direct bandgap of Sr₂LiSiO₄H was estimated to be \sim 5.2 eV, whereas that of Sr_2LiSiO_4F was estimated to be larger *i.e.*, \sim 5.5 eV. These values are 1.1–1.4 eV smaller than those predicted by G_0W_0 calculations, indicating that the observed absorption below 6 eV is induced by bulk excitons or defects such as O²⁻ and OH⁻ in $X^{-}(X^{-} = F^{-}, H^{-})$ sites. Accordingly, Sr_2LiSiO_4H and Sr_2LiSiO_4F powder samples appear white-grey and white under daylight, respectively.

The photoluminescence excitation (PLE) and PL spectra of Sr₂LiSiO₄H:Eu²⁺ recorded at room temperature are depicted in Fig. 4. As a reference, the spectra of Sr_{1.96}Eu_{0.04}LiSiO₄F (Sr₂- $LiSiO_4F:Eu^{2+}$) were also recorded under the same conditions. Under an excitation light of 293 nm, Sr₂LiSiO₄H:Eu²⁺ displayed a single emission peak centered at 544 nm with a full width at half maximum (FWHM) of 0.42 eV, whereas Sr₂LiSiO₄F:Eu²⁺ displayed an emission peak at 506 nm with a FWHM of 0.7 eV under an excitation light of 324 nm. The absence of sharp emission lines characteristic of 4f-4f transitions in Eu³⁺ confirmed a complete reduction from Eu³⁺ to Eu²⁺ for both samples. This result is consistent with the magnetization data indicating that the most of Eu atom form divalent states with ⁸S_{7/2} configuration (see Fig. S3†). The broad emission band was attributed to the parity-allowed electronic dipole transitions in Eu^{2+} from 4f⁶5d to 4f⁷ configuration. The band could be deconvoluted into two Gaussian peaks, which could be assigned to Eu occupying two different Sr sites. Sr₂LiSiO₄H:Eu²⁺ possesses an emission band redshifted from that of Sr₂LiSiO₄-F:Eu²⁺ and emits intense yellow luminescence under illumination by near-UV (375 nm) LED as shown in the inset of Fig. 4.

In each PLE spectrum recorded, there was only one broad absorption band consisting of at least five or more overlapping Gaussian peaks. Each broad band could be ascribed to the



Fig. 2 Band structures and DOS of (a) Sr_2LiSiO_4F and (b) Sr_2LiSiO_4H . The energy axes are adjusted by the O 2s levels located at -18 to -19 eV below each E_{VBM} .



Fig. 3 Diffuse reflectance spectra of Sr₂LiSiO₄F and Sr₂LiSiO₄H. Inset shows $(F(R)hv)^2 - hv$ plots to estimate band gap energies. The $(F(R)hv)^2$ data of Sr₂LiSiO₄F were multiplied by 10 so that both sets of data could be observed within the same coordinate axis range.

crystal field splitting of Eu 5d levels of $4f^7 \rightarrow 4f^{6}5d^{1}$. For Sr₂-LiSiO₄H:Eu²⁺, the broad band was observed within the wavelength region of 250–450 nm, whereas in Sr₂LiSiO₄F:Eu²⁺, the broad band was observed within the higher photon energy region of 230–410 nm. An energy level scheme was established accordingly, with each peak within the broad excitation band corresponding to the Eu 5d levels and the position of the lowestexcitation peak (as marked "5") was considered as the Eu lowest



Fig. 4 Photoluminescence properties of (a) $Sr_2LiSiO_4F:Eu^{2+}$ and (b) $Sr_2LiSiO_4H:Eu^{2+}$. PL and PLE spectra and corresponding photographs of the materials under illumination with near-UV light of 375 nm. $Sr_2LiSiO_4F:Eu^{2+}$ was excited at 324 nm and monitored at 506 nm. $Sr_2LiSiO_4H:Eu^{2+}$ was excited at 293 nm and monitored at 544 nm. Both PLE and PL spectra were deconvoluted by Gaussian fitting.

5d energy level. In more details, the absolute position of the lowest 5d level depends on the redshift which typically consists of centroid shift and crystal field splitting. Centroid shift is mainly influenced by nephelauxetic effects, whereas crystal field splitting is dependent on the site symmetry.³⁷ The lowestexcitation peak positions for $Sr_2LiSiO_4F:Eu^{2+}$ and $Sr_2LiSiO_4 H:Eu^{2+}$ were 3.22 eV and 3.02 eV, indicating the downshift of Eu lowest 5d energy level and further confirming the high likelihood that the replacement of fluoride with hydride leads to the red shift of the absorption band due to large nephelauxetic effects exerted by H^- ligands.

Subsequently, we theoretically investigated the Eu²⁺ excitation and emission energies for Sr₂LiSiO₄H:Eu²⁺ and Sr₂LiSiO₄-F:Eu²⁺ using cDFT calculations with the configurational coordinate diagram. To introduce Eu into the host lattices, 2 \times 2×1 supercells were applied with one of the Sr atoms replaced by Eu, namely Sr₃₁Eu₁Li₁₆Si₁₆O₆₄H₁₆ and Sr₃₁Eu₁Li₁₆Si₁₆O₆₄F₁₆. Here, we considered two scenarios of Eu substituting into Sr1 or Sr2 site, respectively. According to the configurational coordinate diagram, excitation and emission processes occur among four electronic states that are ground state A0, excited state without structural relaxation A₀^{*}, excited state after relaxation A*, and ground state with the relaxed structure A. For the excitation process, one electron located at Eu 4f level was excited to the 5d level by absorbing a photon with energy larger than the energy difference between the A_0 and A_0^* states. Lattice relaxation corresponding to non-radiative processes from A_0^* to A*, proceeded owing to imbalance of the electronic configuration in Eu²⁺. The excited electron in the 5d level then returned to the 4f level by emission of a photon resulting in transition from an A* to an A state. On the basis of the total energy of each state, we calculated the absorption energy $E_{abs}(E_0^* - E_0)$, emission energy $E_{\rm em}(E^* - E)$, and the Stokes shift $\Delta S (E_0^* - E_0)$, which are listed in Table 1. The supercell structure was first optimized to obtain the A_0 state by standard DFT calculation using the +U method with U = 6 eV for both Sr₂LiSiO₄H and Sr₂LiSiO₄F to localize the Eu 4f bands within the bandgap. In the ground state, narrow Eu 4f bands occupied by seven spin-up electrons, were located above the valence band maximum (VBM) of the host lattice. The A₀^{*} state was calculated by cDFT method which constrains the electronic occupancy of the highest-occupied band to be 0 and that of the lowest unoccupied band to be 1. Thus, two new bands were formed by the core-hole interaction. One is unoccupied 4f bands which shifts above the valence band and the other is occupied 5d band which shifts below the conduction band. The cDFT method was continuously utilized to maintain electronic configuration during the second geometry optimization while the energy of the Eu 5d band was reduced. For the A state, without constraining the electron occupancy, the excited electron automatically returned to the

Table 1 Experimental (Exp.) and calculated (Cal.) Stokes shifts (ΔS) and absorption (E_{abs}) and emission (E_{em}) energies of Sr₂LiSiO₄F and Sr₂LiSiO₄H^a

	Sr ₂ LiSiO ₄ F		Sr ₂ LiSiO ₄ H	
	Exp. (eV)	Cal. (eV)	Exp. (eV)	Cal. (eV)
Eabs	3.22	4.49(Eu1) 4.78(Eu2)	3.02	4.12(Eu1) 4.20(Eu2)
E _{em}	2.40	3.16(Eu1)	2.18	2.97(Eu1)
	2.46	3.44(Eu2)	2.34	3.40(Eu2)
ΔS	0.82	1.34(Eu1)	0.84	1.15(Eu1)
	0.76	1.33(Eu2)	0.68	0.81(Eu2)

^{*a*} For the calculations, standard DFT+*U* and cDFT were used.

Eu 4f band with the same equilibrium configuration coordinates of the A* state. The band structures in each electronic state were assessed further for both Sr_2LiSiO_4F and Sr_2LiSiO_4H (Fig. S3†). Noteworthy, the cDFT calculations overestimate the absorption/emission energy relative to the experimental results. In fact, the cDFT method cannot be used to accurately demonstrate each splitting band position but only the tendency of energy transformations. However, the smaller absorption energy (E_{abs}) which indicates the downshifting of the Eu lowest 5d energy level in Sr_2LiSiO_4H from the calculated data wellagreed with experimental results. Compared with Sr_2LiSiO_4F , Sr_2LiSiO_4H showed lower absorption and emission energies irrespective of the Sr site substitution of Eu. This result is consistent with the observed red shifts in both absorption band and emission band induced by the substation of F^- by H^- .

Herein, we mainly discuss the absorption energies difference between Eu 5d levels in Sr₂LiSiO₄H:Eu²⁺ and Sr₂LiSiO₄F:Eu²⁺. According to the semiempirical model reported by Dorenbos,38-41 both centroid shift and crystal field splitting can separately cause downshifting of the lowest Eu 5d level. In particular, crystal field splitting ε_{cfs} can be described as $\varepsilon_{cfs} =$ β/R_{av}^2 , where β is a measure of the shape and size of the Eucentral polyhedron and R_{av} is the average bonding length of the Eu site coordination.41 As shown in Tables S7 and S8,† the bond lengths and bond angles of the coordinate structures around Eu²⁺ in A₀^{*} states were almost same in Sr₂LiSiO₄H and Sr₂LiSiO₄F, indicating that the degree of crystal field splitting for excitation energy was comparable in these two structures. This similarity is due to the identical charge and similar ionic radii of F⁻ and H⁻. Therefore, the downshifting of the Eu 5d level corresponding to the redshift of PLE band in Sr₂LiSiO₄H should be attributed to the difference in the degree of centroid shift. As stated in introduction section, hydrogen has a lower electronegativity than fluoride and can give the strong nephelauxetic effect inducing large centroid shift which presents as the right shift of absorption band redshift in Sr₂LiSiO₄H:Eu²⁺. The present findings illustrate the potential of hydride ion substitution in oxyfluoride phosphor in inducing a red shift of the absorption bands. In a very recent study by Gehlhaar et al.42 on the synthesis and photoluminescence property of Sr₂LiSiO₄H:Eu²⁺, the authors also observed red shifts of the PL band induced by substitution of F⁻ with H⁻. However, our research demonstrates that the red shifting of absorption band due to the strong nephelauxetic effect giving by hydride ligands in hydride analogue of oxyfluoride compounds are possibly predicted by cDFT calculation.

Conclusions

 Sr_2LiSiO_4H was synthesized by high-temperature reaction of oxide and hydride precursors in H_2 gas at 0.9 MPa. The synthesized compound was stable in air and the Eu^{2+} -doped analog displayed strong yellow photoluminescence upon excitation by near-UV light. The cDFT calculations successfully simulated the lowering of the Eu 5d level upon H⁻ substitution, and the calculated red shifts of the PL and PLE energies were consistent with the experimentally observed red shifts. In this

oxyhydride, the mixed ligand sphere containing hydride gives the nephelauxetic effect stronger than that in oxyfluoride analogue, which influences on the centroid shift of the Eu 5d levels to lower energy region. The present findings demonstrate that the hydride ligand can be used to design novel phosphor materials and modify PL properties of conventional oxide-based materials.

Conflicts of interest

There are no conflicts to declare.

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