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Towards white light emission from a hybrid thin film of a self-assembled ternary samarium(III) complex†

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A new samarium complex [Sm(hfaa)₃(Py-Im)] (ML) was synthesized by utilizing hexafluoroacetylacetone (hfaa) and 2-(2-pyridyl)benzimidazole (Py-Im) as coordinating ligands. Single crystal X-ray analysis shows that in the solid-state the complex dimerizes through intermolecular hydrogen-bonding with N–H···O (2.127 Å) and N–H···F (2.576 Å) interactions. The complex in the solid-state displayed highly monochromatic brilliant red emission with Commission International de l'Éclairage (CIE) color coordinates of 0.6532; 0.3336, with a remarkably long luminescence lifetime ($\tau_{\text{exp.}} \approx 204.47 \pm 4.043 \mu\text{s}$) and a high intrinsic quantum yield ($Q_{\text{Sm}}^{\text{Sm}} \approx 6.60\%$). Furthermore, the complex in different organic media displayed color tunability from orange (CIE; 0.6071; 0.3568) to violet (CIE; 0.3634; 0.2144) with $\tau_{\text{exp.}}$ from 105.50 ± 0.143 to $14.27 \pm 0.033 \mu\text{s}$. Hybrid and flexible thin films obtained by doping different concentrations (1%, 2%, 4% and 6%) of the complex into the poly(urethane) (PU) polymer matrix also exhibited color tunability from violet (CIE; 0.3380; 0.2162) to light orange (CIE; 0.5666; 0.3408). White light emission from a 1% doped thin film of the complex was realized by changing the excitation wavelength. Thus, the new material could be a potential candidate for the fabrication of full-color display devices and for solid-state lighting (SSL) applications.

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1. Introduction

Highly photoluminescent lanthanide (Ln) complexes incorporating organic ligands that utilize the antenna effect to generate efficient luminescent materials continue to be of interest to academics and industrialists because of their applications. A range of intriguing technological applications incorporating organo-lanthanide complexes has been developed such as organic light emitting diodes (OLEDs),¹ laser materials,² sensitizers to improve the electroluminescence (EL) of red and green emitting iridium complexes,^{1a,3} luminescent thermometers,⁴ or sensors.⁵ The extensive research efforts are due to the unique optical properties of trivalent lanthanide ions [Ln(III)], which feature fingerprint emissions for each Ln(III) ion spanning through the visible to near-infrared (NIR) region of the spectrum, exhibit highly

monochromatic emissions, and have long-lived excited lifetimes. However, most of the work reported to date deals with either Eu(III) or Tb(III) that emits intense red and green light. Work related to luminescent Sm(III) complexes is very limited by comparison.

Generally, an efficient Sm(III) complex would display a deep red emission⁶ because of the presence of the electric-dipole $^4G_{5/2} \rightarrow ^6H_{9/2}$ transition at 648 nm and thus is beneficial for constructing the red element in a full-color display. Moreover, the emission spectrum of Sm(III) ion displays two more medium intensity transitions at 563 nm (greenish-yellow emission) and 606 nm (yellowish-orange emission). The presence of these two medium intensity emission transitions in the spectrum offers emission color tunability; if the emission is not very efficient that would lead to residual ligand fluorescence (RLF) in the blue region of the spectrum. In fact, proper amalgamation of these colors in Sm(III) complex(es) could result in white light emission from a single molecule platform.⁷ This is achievable by change in concentration, excitation, temperature and solvent medium or by embedding the complex into a polymer matrix.^{7b,d,e} Motivated by these ideas, and following on from some previous luminescence studies of Sm(III) hexafluoroacetylacetone complexes,⁸ we have synthesized a new ternary Sm(III) complex using hexafluoroacetylacetone (hfaa) as the primary antenna and 2-(2-pyridyl)benzimidazole (Py-Im) as the secondary ligand (Scheme 1). The primary hfaa ligand was chosen because of the presence of

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Scheme 1 Synthesis of [Sm(hfaa)₃(Py-Im)] complex.

low vibrational bonds (C–F) that suppress radiationless transitions *via* vibrational relaxations.⁹ The bidentate secondary Py-Im ligand was chosen because of its compatible triplet state (${}^3\pi\pi^* \approx 20576 \text{ cm}^{-1}$)^{1a} and its asymmetrical *N,N*-chelating coordination mode *via* the five-membered imidazolyl nitrogen and the six-membered pyridyl nitrogen atoms, and thus could be useful in enhancing the radiative decay rate.¹⁰ Moreover, the NH proton of the imidazolyl ring is usually involved in hydrogen bond formation and self-assembly, resulting in fascinating structures and arrangements in the solid-state. The complex was characterized by analytical and spectroscopic methods and the solid-state structure was determined by single crystal X-ray diffraction (SC-XRD). It was found that NH proton in the complex contributes to the formation of a hydrogen bonded dimer in the solid-state, which is not common in this class of complexes.¹¹ The photophysical properties of the complex were analyzed in detail by steady-state excitation and emission spectroscopy as well as by time-resolved spectroscopy in the solid-state and in different organic media (dichloromethane, acetonitrile, acetone, tetrahydrofuran, ethanol and methanol). The complex exhibits color tunability and shows a brilliant red emission in the solid-state with a remarkably long luminescence lifetime ($\tau_{\text{exp.}} \approx 204.47 \pm 4.043 \text{ }\mu\text{s}$) and an intrinsic quantum yield of $Q_{\text{Sm}}^{\text{Sm}} \approx 6.60\%$. Upon changing the medium, the complex displays color tunability from orange to violet with an obvious change in the luminescence lifetime, thus hinting towards white light generation. Considering these interesting photophysical properties and the possibility of generating white light emission at a single component level, the complex was further embedded into a PU polymer matrix with different doping concentrations to develop color-tunable flexible hybrid thin films with a view to developing high-performance luminescent materials for optoelectronic application.

2. Experimental section

2.1 Materials, syntheses and analyses

$\text{SmCl}_3 \cdot 6\text{H}_2\text{O}$ was purchased from Strem Chemicals, Inc. USA, while other chemicals were obtained from Sigma Aldrich and used as received. Solvents were pre-dried and distilled before use according to standard procedures.¹² Elemental analysis was performed on Euro EA – CHN Elemental Analyser in the Department of Chemistry, Sultan Qaboos University. Attenuated total-reflectance

(ATR) infrared (IR) spectra of pure solid samples were recorded on diamond using a Cary 630 FT-IR spectrometer. Mass spectra were obtained using a VG Autospec magnetic sector instrument using electrospray ionization (ESI).

2.2 Synthesis of the [Sm(hfaa)₃(Py-Im)] ternary complex

The samarium complex was synthesized by a one-step reaction at room temperature (RT) (Scheme 1). To an ethanolic hfaa (0.73 g; 3.50 mmol) solution, 25% ethanolic ammonia (0.27 mL) solution was added dropwise with constant stirring. After the mixture was stirred for a further 30 min, ethanolic solutions of Py-Im (0.226 g, 1.16 mmol) and $\text{SmCl}_3 \cdot 6\text{H}_2\text{O}$ (0.422 g, 1.16 mmol) were added sequentially, and the reaction mixture was left with stirring for a further 6 h. The solution was left to allow slow solvent evaporation. After a week, a yellow powder precipitated. The precipitate was filtered, washed with ice-cold ethanol followed by hexane, and dried in the air to obtain a yellow solid in 65% yield. Calculated for $\text{C}_{27}\text{H}_{12}\text{F}_{18}\text{N}_3\text{O}_6\text{Sm}$: C, 33.55; H, 1.25; N, 4.35; found C, 33.88; H, 1.20; N, 4.29%. ESI-MS – $[\text{M} + \text{K}(\text{CH}_3)_2\text{CO}]$; $m/z = 948.1$; melting point (T_m) = 195 °C.

2.3 Single crystal X-ray diffraction analyses

A single crystal of [Sm(hfaa)₃(Py-Im)] with dimensions of $0.77 \times 0.57 \times 0.43 \text{ mm}$ was selected for X-ray diffraction. The intensity data was collected at 296 K on a STOE IPDS 2 diffractometer equipped with graphite monochromatic Mo-K α radiation ($\lambda = 0.71073 \text{ \AA}$). The structure was solved by direct methods using SHELXT¹³ and refined by full-matrix least-squares methods on F^2 using SHELXL17/1.¹⁴ All the hydrogen atoms were placed in calculated positions and refined using a riding model, with C–H = 0.93 Å for CH H-atoms, and $U_{\text{iso}}(\text{H}) = 1.2U_{\text{eq}}(\text{C})$. The N-bound H-atom was positioned geometrically (N–H = 0.86 Å) and refined with a riding model, with $U_{\text{iso}}(\text{H}) = 1.2U_{\text{eq}}(\text{N})$. The molecular geometry calculations and drawings were performed with Mercury for Windows¹⁵ and PLATON.¹⁶ WinGX¹⁷ was used to prepare material for publication. The crystal data, data collection and refinement parameters for [Sm(hfaa)₃(Py-Im)] are presented in Table S1, CCDC 1943973.†

2.4 Hirshfeld surface analysis

Hirshfeld surface (HS) analysis (including fingerprint plots) was used to understand the interactions involved in the crystal packing. The Hirshfeld surface analysis¹⁸ and the associated





Fig. 1 (a) A view of the asymmetric unit of $[\text{Sm}(\text{hfaa})_3(\text{Py-Im})]_2$ and (b) comparative view of $[\text{Sm}(\text{hfaa})_3(\text{Py-Im})]$ represented via $\text{N-H}\cdots\text{O}$ and $\text{N-H}\cdots\text{F}$ intermolecular interactions. Dotted pink and blue lines depict the intermolecular interactions (c) packing diagram of the crystal exhibiting $\text{F}\cdots\text{F}$ interaction between hfaa ligands.



Fig. 2 (a) The d_{norm} -mapped Hirshfeld surface for visualizing the intermolecular contacts for the complex (b) 2D fingerprint plots and (c) π -chart for complex showing the percentage contribution of interactions.

CH_3 groups by CF_3 groups in the ligands, compared to other studies in the literature,^{2c,22} in this complex, the contribution of $\text{H}\cdots\text{H}$ interactions (4%) to the total HS is quite low.

3.2 Thermal analysis

An important factor for the use of lanthanide-based hybrid materials in optoelectronic devices is their high thermal stability.

The thermal stability of the $\text{Sm}(\text{III})$ complex was determined simultaneously by thermogravimetric analysis (TGA) and differential thermal analysis (DTA) as shown in Fig. 3. The thermogram exhibits a one-step weight loss due to the removal of the coordinated organic ligands (hfaa and Py-Im) with the decomposition temperature (T_d) ≈ 224 °C. The DTA curve shows one sharp endothermic peak at 195 °C representing the melting point (T_m) of the complex.



Table 1 Photophysical parameters of the complex in different media at RT

Medium	FWHM ^a (nm)	R _{Sm} ^b	τ _{exp.} (μs)	CIE (x; y)	Color	Q _{Sm} ^{Sm c} (%)
Solid-state	11.83	6.00	204.47 ± 4.043	0.6532; 0.3336	Brilliant red	6.60
DCM	12.25	2.75	105.50 ± 0.143	0.6071; 0.3568	Orange	3.40
ACN	11.13	2.70	42.75 ± 0.055	0.5607; 0.3167	Dark-coral	1.37
Acetone	7.61	2.80	37.80 ± 0.187	0.4347; 0.2328	Rose-pink	1.22
THF	6.55	2.47	35.01 ± 0.187	0.3451; 0.2355	Light-violet	1.13
EtOH	7.96	2.20	20.39 ± 0.041	0.4409; 0.2623	Pale pink	0.65
MeOH	8.02	2.39	14.27 ± 0.033	0.3634; 0.2144	Violet	0.46

^a FWHM of ${}^4G_{5/2} \rightarrow {}^6H_{9/2}$ ED transition. ^b Ratio of corrected emission intensity of ED to MD transition, *i.e.*, $\frac{{}^4G_{5/2} \rightarrow {}^6H_{9/2}}{{}^4G_{5/2} \rightarrow {}^6H_{7/2}}$. ^c $Q_{Sm}^{Sm} = \frac{\tau_{exp.}}{\tau_{rad.}}$.

hypersensitive transition dominates with the full width at half maxima (FWHM) ≈ 11.83 nm (Table 1) followed by the MD transition. It is well-established that the ratio (R_{Ln}) of the corrected ED to MD emission transition gives clues regarding the coordination symmetry and polarizability of the local environment around the Ln(III) ion in a given complex *i.e.*, $R_{Sm} = ED({}^4G_{5/2} \rightarrow {}^6H_{9/2})/MD({}^4G_{5/2} \rightarrow {}^6H_{7/2})$. The higher intensity of the ED transition and high R_{Sm} value of 6.00 for the present complex suggests a low molecular symmetry of the coordination sphere and that Sm(III) ion in the complex sits in a highly polarizable environment. The R_{Sm} value is comparable to values reported for similar Sm(III) complexes.^{2c,8a} It worth noting that there is no residual ligand fluorescence in the 400–500 nm range suggesting that the energy absorbed by the organic ligands is efficiently transferred to Sm(III) emitting state as can be seen in Fig. 4b and colorful emission spectrum in Fig. S5, ESI† as expected by the energy gap law. To get information about the emitted color from the solid complex, the CIE color coordinates (x,y) were calculated from the steady-state emission spectrum and the results are summarized in Table 1. As can be seen from the magnified CIE color diagram in Fig. 5, the complex emits intense red emission with color coordinates $x = 0.6532$; $y = 0.3336$, which are similar to National Television System Committee (NTSC) ($x = 0.67$; $y = 0.33$). It is important to emphasize that there are only few reported [Sm(β-diket.)₃(ancillary ligand)] complexes that display an intense red emission.^{6,8a,b}



Fig. 5 A magnified view of the CIE color diagram showing the intense red emission from the complex in the solid-state at RT.

In addition to the steady-state emission, we also carried out the time-resolved PL decay dynamics. The PL lifetime (τ_{exp}) of the complex in the solid-state was measured at RT (Fig. S6 (ESI†) and Table 1) by monitoring the emission decay curve within ${}^4G_{5/2} \rightarrow {}^6H_{9/2}$ transition at 648 nm. The decay curve fits very well with mono-exponential behavior, indicating the presence of only emitting Sm(III) center in the solid-state. The complex displays remarkably long $\tau_{exp.} \approx 204.47 \pm 4.043$ μs, which could be attributed to optimum ΔE between Sm(III) emitting state and prime hfaa and ancillary Py-Im ligands. Furthermore, the minimization of high energy oscillator strength of the N–H by intermolecular interactions *i.e.*, N–H...O and N–H...F as shown in Fig. 1b and was complimented by the FTIR results, which show a relatively less strong sharp peak for N–H oscillator (Fig. S2, ESI†). Moreover, as determined by HS maps, the F...H/H...F largest relative contribution, amounting to 39.4% in the present complex further diminishes the role of another high energy oscillator enhancing the PL lifetime. Thus, HS analysis and 2-D fingerprint were very useful for understanding the contribution of intermolecular interactions in determining the PL properties. The PL lifetime of the present complex is higher than most of the known highly efficient [Sm(β-diket.)₃(ancillary ligand)] complexes such as [Sm(hfaa)₃(TPPO)₂] ≈ 97 μs,^{8a} [Sm(hfaa)₃(phen)₂] ≈ 37 μs^{8a} and 54–56 μs,⁶ [Sm₂(BTP)₃(phen)₂] ≈ 100 μs,²⁸ [Sm(hfaa)₃(*t*Bu-xantpo)] ≈ 150 μs^{8c} and is lower only two longest and efficient complexes [Sm(hfaa)₃(pybox)] ≈ 390 μs^{8b} and [Sm(hfaa)₃(dpepo)] ≈ 280 μs^{8c} until now, where TPPO = triphenylphosphine oxide; phen = 1,10-phenanthroline; BTP = 1,3-bis(4,4,4-trifluoro-1,3-dioxobutyl)phenyl; pybox = bis(oxazolonyl)pyridine; dpepo = bis[(2-diphenylphosphoryl)phenyl] ether; *t*Bu-xantpo = 4,5-bis-(di-*tert*-butylphosphoryl)-9,9-dimethylxanthene. The intrinsic quantum yield (Q_{Sm}^{Sm}) of the complex is calculated by the following general expression 2 and the obtained data is summarized in Table 1:

$$Q_{sm}^{sm} = \frac{\tau_{exp.}}{\tau_{rad.}} \quad (2)$$

where τ_{rad} is the calculated natural radiative lifetime of Sm(III) ion (3100 μs) reported in the literature.²⁹

3.4 Effect of medium on PL properties and color.

ΔE between the emitting level and the next lower energy level in Sm(III) is ≈ 7400 cm⁻¹.³⁰ Therefore PL quenching is inevitable by the high energy oscillators (O–H ~ 3300 –3500 cm⁻¹,



thus the Franck–Condon factor becomes more favorable. This results in higher quenching compared to DCM and other solvents and less efficient ET, which could be seen (Fig. 5 and separate emission diagram Fig. S8–S13, ESI†) with concomitant decrease in the intensity of hypersensitive $^4G_{5/2} \rightarrow ^6H_{9/2}$ (I_{Red}) in the red

region and simultaneous increase in intensity of RFL (I_{RFL}) in the blue region. The change in the I_{Red} and I_{RFL} will subsequently effect the emitted color of the complex. In order to see the emitted color in different solvents, we calculated the CIE color coordinates (x, y) from the emission spectra and the obtained color coordinates are shown in Table 1 and Fig. 7. It is clear from the CIE color diagram that the present complex displays different colors in different solvents such as orange color in DCM and violet in MeOH (Fig. 7b) due to increase of I_{RFL} .

The $Q_{\text{Sm}}^{\text{Sm}}$ of the complex in each solvent was calculated (Table 1) and follows the trend: DCM \gg ACN $>$ Acetone $>$ THF $>$ EtOH $>$ MeOH, respectively, which is directly related to τ_{exp} . Looking at these interesting PL properties, *i.e.*, remarkably long τ_{exp} in the solid-state and color tunability from orange to violet of the present complex in different solvents, we decided to develop hybrid thin films by encapsulating it in a PU polymer matrix. The encapsulation of lanthanide complex into polymer matrix usually increases the thermal and photostability coupled with the enhanced mechanical strength of the developed hybrid material, a necessary prerequisite for optoelectronic applications.^{2a,33} The hybrid and flexible thin films were developed by doping different concentrations such as 1%, 2%, 4% and 6% of the complex into the PU polymer matrix assuming that it would give us color-tunable materials. The PL properties of the flexible



Fig. 8 Corrected emission spectra of $[\text{Sm}(\text{hfaa})_3(\text{Py-IIm})]@PU$ at different doping concentrations at RT.

Table 2 Photophysical parameters of hybrid thin films of $[\text{Sm}(\text{hfaa})_3(\text{Py-IIm})]$ at RT

Medium (%)	FWHM ^a (nm)	R_{Sm}^b	τ_{exp} (μs)	CIE (x, y)	Color	$Q_{\text{Sm}}^{\text{Sm}c}$ (%)
1	5.66	1.99	43.55 ± 0.274	0.3380; 0.2162	Violet	1.40
2	5.90	2.45	46.85 ± 0.240	0.5028; 0.3080	Coral pink	1.51
4	5.72	2.40	51.81 ± 0.149	0.5295; 0.3203	Light coral	1.67
6	5.84	2.41	54.14 ± 0.131	0.5666; 0.3408	Light orange	1.74

^a FWHM of $^4G_{5/2} \rightarrow ^6H_{9/2}$ ED transition. ^b Ratio of corrected emission intensity of ED to MD transition, *i.e.*, $\frac{I_{450-500}}{I_{600-700}}$. ^c $Q_{\text{Sm}}^{\text{Sm}} = \frac{\tau_{\text{exp}}}{\tau_{\text{rad}}}$.



Fig. 9 (a) 1931 CIE Chromaticity diagram and (b) a magnified CIE showing the exact emitted color of $[\text{Sm}(\text{hfaa})_3(\text{Py-IIm})]@PU_x\%$ in different doping concentration at RT.



thin films were analyzed by the steady-state excitation and emission spectra. The excitation spectra of thin films displayed similar broadband as observed in the solid-state and in solution (Fig. S20, ESI†).

The emission spectra of the hybrid thin films obtained by exciting at $\lambda_{\text{Ex}}^{\text{max}}$ displayed typical metal centered emission transitions (discussed above) as observed in the solid-state and in solution as shown in Fig. 8 and the data obtained is shown in Table 2. Hypersensitive $^4G_{5/2} \rightarrow ^6H_{9/2}$ emission transition dominates the spectra and is narrower in hybrid thin films with

FWHM in 5.66–5.90 nm indicating the higher potential of the developed hybrid thin film for optoelectronic applications. R_{Sm} values remain in the range of 1.99–2.45 (Table 2) as observed in different solvents suggesting that incorporation of the complex in the PU polymer matrix does not alter the symmetry and coordination sphere of the bare complex, so it does not lose its structural identity. The τ_{exp} of hybrid thin films were determined as discussed above and the obtained data is gathered in Table 2 while the decay curves together with fitted curves are shown in Fig. S21–S24, ESI.† The lifetime of the hybrid thin



Fig. 10 Corrected emission spectra of hybrid thin films at different excitation wavelengths at RT. The values are CIE (x,y) color coordinates obtained.



Fig. 11 1931 CIE Chromaticity diagram of hybrid [Sm(hfaa)₃(Py-Im)]@PU_1% thin film at different excitation wavelength displaying near white-light emission.



films is in the range of 43.55–54.14 μs which is almost 4- and 2-fold shorter than in the solid-state and in DCM solution, this decrease could be attributed to the presence of a high-energy oscillator in the PU polymer matrix, nevertheless it remains at higher end of efficient emitting samarium complexes. To obtain the information for the emitted color by the different hybrid thin films, the CIE color coordinates (Table 2) and diagram were obtained from the steady-state emission spectra and are shown in Fig. 9a and a magnified view of the exact color is also shown in Fig. 9b. As can be seen and as per our assumption from Fig. 9, the developed material displayed color-tunable emission from violet to orange as the doping concentration of the complex was increased from 1–6%. The change in the color from violet to orange is due to the decrease in the I_{RFL} and can be clearly seen in Fig. S25–S28, ESI.†

Moreover, to see more color tunability and to increase the chance to get the near-white light or white light emission from the developed hybrid thin films, the excitation wavelength was varied in the range of 270–340 nm at intervals of 10 nm. 1% and 6% doped hybrid materials were taken as reference models, as shown in Fig. 10, the steady-state emission spectra of both thin films exhibit increased emission intensity of the $^4\text{G}_{5/2} \rightarrow ^6\text{H}_{9/2}$ emission transition with I_{RFL} decreasing simultaneously. The corresponding CIE color coordinates is shown in their respective diagram while CIE chromaticity are shown in Fig. 11 and Fig. S29, ESI.† It can be seen from Fig. S29, ESI,† that 6% hybrid thin film displayed orange to light pink. Interestingly, the emission of 1% doped hybrid thin film falls mostly in the white region 1931 CIE chromaticity diagram as can be seen clearly in Fig. 11. On excitation at 300 and 310 nm, the 1% doped hybrid thin film exhibited a near white light emission with CIE coordinates of $x = 0.3673$; $y = 0.2843$ and $x = 0.3595$; $y = 0.2814$, respectively, which are very close to that for pure white light ($x = 0.33$, $y = 0.33$) according to the 1931 CIE diagram. Therefore this material could be a potential candidate to be used as a single-component material to generate white light or near-white light emission for SSL application.³⁴

4. Conclusion

We have synthesized a medium-dependent color-tunable $\text{Sm}(\text{III})$ complex that dimerizes in the solid-state through $\text{N-H}\cdots\text{O}$ (2.127 Å) and $\text{N-H}\cdots\text{F}$ (2.576 Å) hydrogen bonds. White light emission was achieved by adjustment of doping concentration into a PU polymer and excitation wavelength (as shown in Fig. 10a and 11). ΔE value of $\approx 7400\text{ cm}^{-1}$ between the emitting $^4\text{G}_{5/2}$ level and the next lower energy level in the complex results in effective quenching by high energy oscillators and reduces the overall ET efficiency. This leads to RFL in the range between 350–500 nm, which plays a vital role in obtaining white light emission. The present work opens the door to exploit similar color-tunable $\text{Sm}(\text{III})$ based white light materials for the fabrication of full-color display devices and SSL applications.

Conflicts of interest

There are no conflicts of interest to declare in this work.

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