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## **Supporting Information**

Electrochemically controllable emission and coloration using a modified electrode with a layered clay compound containing viologen molecule and Europium(III) complex

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**Fig.S1** (a) Chemical structure diagram of  $HV^{2+}$  molecules obtained using MM3 calculation in Chem 3D application. Interatomic distances of the  $HV^{2+}$  molecules ranging from approximately 0.43 nm to 2.55 nm. (b) Molecular structure diagram of Eu(hfa)<sub>3</sub>(TPPO)<sub>2</sub> complex drawn using single crystal X-ray measurement data from ref <sup>52</sup> in main text. Interatomic distances of the Eu(hfa)<sub>3</sub>(TPPO)<sub>2</sub> complex ranging from approximately 1.08 nm to 1.6 nm.



**Fig.S2** (a) Absorption spectra of the  $HV^{2+}$  solution,  $STN/HV^{2+}$  solution, and STN solution; (b) Absorption spectra of  $Eu(hfa)_3(TPPO)_2$  solution,  $STN/Eu(hfa)_3(TPPO)_2$  solution, and STN solution.



**Fig.S3.** Emission spectra of  $HV^{2+}$ ,  $STN/HV^{2+}$  and  $STN/HV^{2+}/Eu(hfa)_3(TPPO)_2$  solutions with excitation wavelength of 260 nm. Inset: photographs of the solutions under UV irradiation.



**Fig.S4**. Absorption spectra of the STN/HV<sup>2+</sup> hybrid solutions and supernatants (left); (b) Adsorption ratios of  $HV^{2+}$  at different ratios (right).



**Fig.S5**. Absorption spectra of the STN/Eu(hfa)<sub>3</sub>(TPPO)<sub>2</sub> hybrid solutions and supernatants (left); (b) Adsorption ratios of Eu(hfa)<sub>3</sub>(TPPO)<sub>2</sub> at different ratios (right).



**Fig.S6** (a) Normalized emission spectra of Eu(hfa)<sub>3</sub>(TPPO)<sub>2</sub>, STN/Eu(hfa)<sub>3</sub>(TPPO)<sub>2</sub> and STN/HV<sup>2+</sup>/Eu(hfa)<sub>3</sub>(TPPO)<sub>2</sub> in the solution. (b) Emission decay curves of Eu(hfa)<sub>3</sub>(TPPO)<sub>2</sub>, STN/Eu(hfa)<sub>3</sub>(TPPO)<sub>2</sub> and STN/HV<sup>2+</sup>/Eu(hfa)<sub>3</sub>(TPPO)<sub>2</sub> in the solution at 613 nm ( ${}^{5}D_{0} \rightarrow {}^{7}F_{2}$ ). Excitation wavelength was 337 nm.



**Fig.S7**. (a) Absorption spectra of the STN/ $HV^{2+}$  film under open circuit condition (black line) and applied -2.0 V for 150 s (blue line)

The donor-acceptor distance is expressed using the following equations.

$$r_{DA} = \left(\sqrt[6]{\frac{1}{E}} - 1\right) \times R_0$$

Here,  $R_{DA}$  is the donor-acceptor distance, E is the energy transfer efficiency, and  $R_0$  is the Förster distance. The Förster distance is expressed as follows.

$$R_0^6 = \frac{9\Phi_0(ln10)k^2J}{128\pi^5 n^4 N_A}$$

where  $\Phi_0$  is the quantum yield of the donor,  $\kappa^2$  is the dipole orientation factor (2/3), J is the overlap integral, n is the refractive index of the solvent, and  $N_A$  is the Avogadro's constant.

The overlap integral of the absorption spectrum of the electrochromic (EC) material and the emission spectrum of the luminescent material is expressed as follows.

$$J = \int f_D(\lambda) \varepsilon_A(\lambda) \lambda^4 d\lambda$$

where J is the overlap integral ( $M^{-1}cm^3$ ),  $f_D$  is the normalized emission spectrum of the luminescent material, and  $\varepsilon_A$  is the molar extinction coefficient ( $Lmol^{-1}cm^{-1}$ ) of each EC material.