Ionic Rigid Organic Dual-State Emission Compound With Rod-Shaped and Conjugated Structure for Sensitive Al<sup>3+</sup> Detection

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Dual-state emission (DSE) luminogens, a type of luminescent material which can effectively emit light in both dilute solution and solid states, have attracted tremendous attention, due to their widespread applications in chemical sensing, biological imaging, organic electronic devices, and so on. They overcome the shortcomings of aggregation-induced emission (AIE)-type compounds that do not emit light in dilute solutions and aggregation-caused quenching (ACQ)-type compounds that do not emit light in a concentrated or aggregated state. This work reports a novel ionic DSE material based on rigid rod-shaped organic conjugated structure using 4,4′-bis(2-sulfonatostyryl) biphenyl disodium salt (BSBDS); the ion repulsion effect can reduce the strong π–π interaction in aggregation and achieve high-efficiency luminescence in solution and solid states. In addition to excellent DSE characteristics, BSBDS also exhibits a mechanochromic nature and sensitive detection performance for aluminum ion (Al<sup>3+</sup>).

Keywords: dual-state emission, Al<sup>3+</sup> detection, organic conjugated compound, rod-shaped structure, mechanochromism

INTRODUCTION

Organic luminescent materials have been widely used in various areas, such as organic light-emitting diodes (OLEDs) and chemical sensors (Reineke et al., 2009; Uoyama et al., 2012; Zhang et al., 2014; Meng et al., 2021). Traditional organic luminescent materials usually have large planar conjugated units, allowing them to exhibit bright emission in solution states; however, these materials often encounter the infamous aggregation-caused quenching (ACQ) effect and exhibit no or weak luminescence in the aggregated state, which extremely limits their wider applications. Fortunately, in 2001, Tang et al. discovered an opposite and interesting aggregation-induced emission (AIE)-type compounds that do not emit light in dilute solutions and aggregation-caused quenching (ACQ)-type compounds that do not emit light in a concentrated or aggregated state. This work reports a novel ionic DSE material based on rigid rod-shaped organic conjugated structure using 4,4′-bis(2-sulfonatostyryl) biphenyl disodium salt (BSBDS); the ion repulsion effect can reduce the strong π–π interaction in aggregation and achieve high-efficiency luminescence in solution and solid states. In addition to excellent DSE characteristics, BSBDS also exhibits a mechanochromic nature and sensitive detection performance for aluminum ion (Al<sup>3+</sup>).
and shows a broad application. In the past few years, organic dual-state emission (DSE) materials that can effectively emit light in both the solution and solid states have been reported sporadically and applied in OLEDs, cell imaging, and other fields. For instance, Tang et al. reported conjugation-induced rigidity and through-space conjugation in twisting molecules for designing novel DSE materials (Chen G et al., 2015; Chen L et al., 2015). Dong et al. reported a series of DSE compounds based on triphenylpyrrole (TPP) (Li et al., 2019) and triphenylquinoline (TPQ) (Dai et al., 2019). Goel et al. reported 5,6-dihydro-2h-fluorophores showing a DSE nature (Raghuvanshi et al., 2017). Ulrich et al. reported π-extended salicylaldehyde fluorophores showing a DSE nature (Stoerkler et al., 2021). Recently, a rational molecular design strategy has been presented based on triphenylamine (Stoerkler et al., 2021). Theoretical calculation and practical verification by the Gong group (Zou et al., 2022). However, although many efforts including various physical and chemical methods have been contributed to designing and exploring new DSE compounds, the research of DSE luminogens remains preliminary with relatively limited examples in the literatures (Belmonte-Vázquez et al., 2021). There still exists a serious lack of a universal molecular design strategy for DSE luminogens.

Herein, an ionic organic conjugated compound with a rigid rod shape was proposed to construct DSE luminogens. First, the rod-shaped structure can effectively inhibit the strong π–π interaction leading to the quenching of luminescence. Second, the ionic compound can avoid the formation of excimer and shows a broad application. In the past few years, organic dual-state emission (DSE) materials that can effectively emit light in both the solution and solid states have been reported sporadically and applied in OLEDs, cell imaging, and other fields. For instance, Tang et al. reported conjugation-induced rigidity and through-space conjugation in twisting molecules for designing novel DSE materials (Chen G et al., 2015; Chen L et al., 2015). Dong et al. reported a series of DSE compounds based on triphenylpyrrole (TPP) (Li et al., 2019) and triphenylquinoline (TPQ) (Dai et al., 2019). Goel et al. reported 5,6-dihydro-2h-fluorophores showing a DSE nature (Raghuvanshi et al., 2017). Ulrich et al. reported π-extended salicylaldehyde fluorophores showing a DSE nature (Stoerkler et al., 2021). Recently, a rational molecular design strategy has been presented based on triphenylamine via theoretical calculation and practical verification by the Gong group (Zou et al., 2022). However, although many efforts including various physical and chemical methods have been contributed to designing and exploring new DSE compounds, the research of DSE luminogens remains preliminary with relatively limited examples in the literatures (Belmonte-Vázquez et al., 2021). There still exists a serious lack of a universal molecular design strategy for DSE luminogens.

Herein, an ionic organic conjugated compound with a rigid rod shape was proposed to construct DSE luminogens. First, the rod-shaped structure can effectively inhibit the strong π–π interaction leading to the quenching of luminescence. Second, the ionic compound can avoid the formation of excimer and complex in the solution due to its own electrostatic repulsion. Combining the rod-shaped structure and ionic characteristics, universal molecular design strategy for DSE luminogens. However, although many efforts including various physical and chemical methods have been contributed to designing and exploring new DSE compounds, the research of DSE luminogens remains preliminary with relatively limited examples in the literatures (Belmonte-Vázquez et al., 2021). There still exists a serious lack of a universal molecular design strategy for DSE luminogens.

Herein, an ionic organic conjugated compound with a rigid rod shape was proposed to construct DSE luminogens. First, the rod-shaped structure can effectively inhibit the strong π–π interaction leading to the quenching of luminescence. Second, the ionic compound can avoid the formation of excimer and complex in the solution due to its own electrostatic repulsion. Combining the rod-shaped structure and ionic characteristics, novel DSE luminogens can be designed effectively to meet various application requirements. In our work, a typical fluorescent whitening agent named disodium 4,4′-bis (2-sulfonatostyryl) biphenyl disodium salt (C28H20Na2O6S2) was purchased from Tokyo Chemical Industry Co., Ltd. Methanol, anhydrous ethanol, and metallic salts were purchased from Xilong Scientific Co., Ltd. Tetrahydrofuran was obtained from Guangdong Guanghua Technology. Standard stock solutions of metal ions (Li⁺, K⁺, Ag⁺, Cu²⁺, Ca²⁺, Mg²⁺, Zn²⁺, Cd²⁺, Co²⁺, Ni²⁺, Cu²⁺, Fe²⁺, Fe³⁺, and Al³⁺) were prepared with ultrapure water from the respective metallic salts (LiCl, KCl, AgNO₃, CaCl₂, MgCl₂, AlCl₃, 6H₂O, Cu(NO₃)₂ 3H₂O, FeCl₂ 4H₂O, FeCl₃ 6H₂O, Zn(NO₃)₂ 6H₂O, CuCl, concentrated sulfuric acid (H₂SO₄), CdCl₂ 5/2H₂O, NiCl₂ 6H₂O, CoCl₂ 6H₂O, and K₂CO₃). All the reagents were of the analytical reagent grade and used as received. All the aqueous solutions were prepared with distilled water supplied from a Milli-Q purification water system. Tetrahydrofuran (THF) was distilled from sodium/benzophenone under nitrogen before use.

**Instruments**

Ultraviolet–visible (UV-Vis) spectra were recorded on a UV3600 spectrophotometer. Fluorescence emission spectra and the quantum efficiency analysis were performed on a Horiba FluorMax-4 spectrofluorometer. X-ray diffraction (XRD) was measured on a X’Pert Pro X-ray diffractometer. The fluorescence lifetime was obtained using a Horiba QM-8000 fluorescence spectrophotometer. The luminescence quantum efficiency in solutions was measured using quinine sulfate as the standard solution (Φₑ = 54%, dilute sulfuric acid as the solvent). The size distribution of the suspensions for BSBDS in 90/10 THF–water was determined on a Malvern Zetasizer nano ZS90 instrument at 25°C. The toxicity of cells was confirmed with a Tecan Infinite F50 Microplate Reader. The fluorescence images of cells were gained on an Olympus FV 3000 laser scanning confocal microscope (LSCM).

**Theoretical Calculation**

Geometry optimizations and frequency calculations were performed by using the density functional theory (DFT), and the time-dependent density functional theory (TD-DFT) was applied to study the low-lying excited states with the B3LYP functional and 6-31+G(d) basis set. All above calculations were carried out with the Gaussian 16 B.01 package (Frisch et al., 2016). Gaussian calculation results were analyzed by Multiwfn (Lu and Chen, 2012) and VMD (Humphrey et al., 1996) software.

**MTT Assay for Cytotoxicity Evaluation**

The toxicity of BSBDS was evaluated using conventional MTT assay. HeLa cells were seeded in a 96-well plate at a density of 1 x 10⁴ cells per well in DMEM. After 24 h of growth, the medium in the cells was replaced with a fresh solution (100 µL) containing BSBDS with different concentrations ranging from 12 (control) to 96 µM. Following the incubation of cells with the BSBDS for 24 h, the medium was replaced with a fresh medium containing MTT (0.5 mg/ml) and incubated for another 4 h. The medium and MTT solution were replaced with 150 µL DMSO in each well. The toxicity of BSBDS was evaluated using conventional MTT assay. HeLa cells were seeded in a 96-well plate at a density of 1 x 10⁴ cells per well in DMEM. After 24 h of growth, the medium in the cells was replaced with a fresh solution (100 µL) containing BSBDS with different concentrations ranging from 12 (control) to 96 µM. Following the incubation of cells with the BSBDS for 24 h, the medium was replaced with a fresh medium containing MTT (0.5 mg/ml) and incubated for another 4 h. The medium and MTT solution were replaced with 150 µL DMSO in each well. The plate was shaken well and the absorbance at 492 nm was recorded using a microplate reader (Infinite F50, Tecan). The cell viability was calculated as the ratio of average absorbance intensity of samples to that of the control.
Cell Imaging
HeLa cells were seeded and grown in DMEM supplemented by 10% fetal bovine serum (FBS) and 1% penicillin/streptomycin (PS) under ambient conditions of 5% CO2 and humidity at 37°C. At 80% confluence, these cells were trypsinized, and about 2 × 10^5 cells were added to each well of a 6-well culture plate. After 24 h of growth, the cells were incubated with a commercial dye (Nuclear Red™ LCS1) at 37°C for 20–30 min; then, Al^3+ and BSBDS were added to the culture medium for incubating (washing off the excess agent and washing cells with PBS three times before adding different reagents). Finally, the cells were fixed with 4% paraformaldehyde. HeLa cell imaging was carried out by using an LSCM (OLYMPUS, FV3000).

RESULTS AND DISCUSSION
Photophysical Properties of BSBDS in Solutions
In view of good water solubility of BSBDS, the photophysical properties of BSBDS were measured in water. The absorption spectrum of BSBDS shown in Supplementary Figure S1 exhibits the maximum absorption peak at 350 nm. The emission spectra and time-resolved emission decay curves of BSBDS in aqueous solution at different concentrations are shown in Figures 2A,B, and the results suggest that BSBDS solution exhibited bright a blue to green emission from low to high concentrations. When the concentration is less than or equal to 2 × 10^{-4} M, the emission peak of BSBDS was at 430 nm. When the concentration reaches 2 × 10^{-2} M, the emission wavelength significantly red-shifted to 470 nm, which can be attributed to the formation of local excimers in BSBDS at high concentrations. The fluorescence lifetime/quantum efficiency of BSBDS at the concentration of 2 × 10^{-2} and 2 × 10^{-5} M are 5.55 ns/62.3% and 1.45 ns/74.3%, respectively, which indicates that luminescence at low concentration (≤ 2 × 10^{-4} M) originates from the intrinsic monomer emission of BSBDS, and luminescence at high concentration (2 × 10^{-2} M) is derived from excimer emission of BSBDS. It is worth noting that the fluorescence intensity of solution at the concentration of 2 × 10^{-5} M is greater than those of at 2 × 10^{-2} M (Supplementary Figure S2), but the opposite is observed on the luminescence photos (Figure 2B). According to previous studies, this phenomenon can be attributed to the self-absorption effect. This phenomenon was reported in Henderson et al. (1977) and Yamasaki et al. (1997). When the excited light irradiates the sample at very high concentrations, there exists an overlap between the absorption spectrum and emission spectrum, and thus some emitted photons...
are absorbed by other solute molecules in the solution before reaching the detectors.

To investigate the photophysical properties of BSBDS at the aggregate state, the emission spectrum of BSBDS in THF/water mixed solution was further tested (H2O and THF are good and poor solvents of BSBDS, respectively) as shown in Figures 2C,D. As the content of THF increases (volume fraction), the luminescent intensity of BSBDS in the THF/water mixed solution shows a significant increase, while the emission peak exhibits a slight blue-shift from 430 to 427 nm. This may be due to the formation of a unique nanostructure from a single-molecule state as the THF content increases. Then, the measurement of particle size for BSBDS in the solution with a THF content of 90% was carried out, and the results indicated that nanoparticles with an average particle size of 119 nm were formed (Figure 2D). It is worth noting that the rod-shaped rigid unit in the middle of BSBDS is lipophilic, and the two ends are hydrophilic sodium sulfonate; this amphiphilic compound can form a variety of unique nanostructures. Compared with the photophysical properties of solution at the concentration of 2 × 10^{-2} M, the solution with the THF content of 90% should have no excimer formed. The slight blue-shift of the spectrum with the increasing THF content is difficult to explain because there is no specific molecular packing model in the nanoparticles. In addition, the increase of luminescent intensity with the increasing THF content can be attributed to RIM in nanoparticles, which is similar to the AIE mechanism (Wang et al., 2015).

**Photophysical Properties of BSBDS in the Solid State**

Similar to their solution, the BSBDS powder shows bright green light upon UV illumination (Figure 3A). According to our previous experience (He et al., 2019), the rod-like ion compound with highly efficient emission in the solid state may possess mechanochromic characteristics. Then, the response of BSBDS to force stimulation was carried out carefully as shown in Figures 3A,B. The crystalline powder of BSBDS recrystallized from water/methanol showed bright green emission peaking at 480 nm; however, after being ground gently with a spatula on filter paper, they exhibited a large blue shift with a deep blue luminescence peaking at 433 nm. In order to explore the mechanochromic mechanism, the XRD measurement of solid

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**FIGURE 3** Photographs of the crystalline and ground powders of BSBDS taken under 365-nm UV light irradiation (A). PL spectra (B), XRD (C), and time-resolved emission decay curves (D) of the crystalline and ground powders of BSBDS.
samples before and after grinding was performed. As shown in Figure 3C, the XRD pattern of the crystalline BSBDS powder exhibited sharp and intense diffraction peaks, which indicated their well-defined microcrystalline lattices. However, their grounded samples did not show any noticeable reflections but had broad diffused halos with low intensity, suggesting their disordered amorphous nature. Clearly, the transition between the ordered crystalline and disordered amorphous states is crucial to the mechanochromism. Besides the change in emission color, the photoluminescence (PL) efficiency of the powders after being ground also increased considerably from 65.5 to 84.5%, which might be ascribed to the destruction of π−π interaction between adjacent molecules in crystalline solid states. Normally, an increased emission efficiency would result in a prolonged lifetime, whereas the formation of excimer would decrease the lifetime. Here, the <τ> values of the crystalline powders and ground solids of DBSDB were determined as 3.03 and 2.21 ns (Figure 3D), respectively. The increased efficiency along with the decreased lifetime of the ground samples verified that the excimer was locally formed in the rigid element, and it was destroyed upon mechanical stimuli.

Theoretical Calculation
In order to deeply understand the mechanism of DSE, the density functional theory and time-dependent density functional theory were applied to study the mechanism of luminescence. First, the B3LYP/6-31 + g(d) method was used to optimize the molecular structure of BSBDS. It is worth noting that the sodium ions (Na+) in the solution had no effect on the luminescence of the BSBDS, so the optimized structure did not contain the content of Na+. The optimized anion structure of BSBDS is shown in Figure 4A. BSBDS presents a typical rod-like structure, and the dihedral angles of ∠C1-C2-C3-C4, ∠C3-C4-C5-C6, and ∠C7-C8-C9-C10 are 19.04, 2.78, and 35.03°, respectively. A smaller dihedral angle means that BSBDS has a larger degree of conjugation. The lowest unoccupied molecular orbital (LUMO) and highest occupied molecular orbital (HOMO) of BSBDS distributed on the whole molecule, and the large degree of overlap indicates that they have no internal charge transfer (ICT) effect (Figure 4B). It coincides with the basically unchanged spectral position in the THF–water mixed solution with a different THF content. In addition, the transition process from HOMO to LUMO has a larger oscillator strength (f = 0.9057). f has a certain relationship with the luminous efficiency (Hilborn, 1982); if the f is less than 0.01, it indicates that the transition is forbidden. Since BSBDS is an amphiphilic compound, it has many ways of molecular accumulation in aggregates or high concentrations. However, due to the rod-like structure and ion repulsion effect, it is possible for BSBDS to form excimers in local areas (Figures 4C,D) to make its luminescence red-shifted, and the luminous efficiency is reduced to a small extent.

Metal Cation Detection
As mentioned previously, the photophysical properties of BSBDS can significantly change after forming an excimer or a complex at a high concentration. Therefore, it is assumed that BSBDS in the dilute solution state can form an excimer or exciplex after binding with a certain cation, which significantly changes its photophysical properties, and it can maybe be used to detect certain ions in the aqueous solution. In order to verify this hypothesis, 14 kinds of metal cations, Li+, Ca2+, Zn2+, K+, Mg2+, Fe2+, Ag+, Cd2+, Co2+, Cu2+, Cu+, Ni2+, Al3+, and Fe3+, were prepared. First, 2.0 ml of different metal cation solutions with a concentration of 2.0 × 10^{-3} M were added to 3.0 ml of BSBDS aqueous solution with a concentration of 2.0 × 10^{-5} M. Surprisingly, as shown in Figure 5A, the luminescent color of
BSBDS changed instantly after the addition of Al$^{3+}$ under 365-nm UV light irradiation. In addition, when Fe$^{3+}$, Cu$^{2+}$, and Cu$^{+}$ ions were added to the BSBDS solution, the luminescence of the mixed solution quenched due to the electron or energy transfer. Furthermore, PL spectra of mixture solutions of BSBDS combined with different metal ions were studied (Figure 5B). Except for the above Cu$^{+}$, Cu$^{2+}$, Fe$^{3+}$ which led to quench the emission of BSBDS, the addition of other ions showed a little effect on the emission peak (~430 nm) and intensity. However, when Al$^{3+}$ ions were added to the BSBDS solution, the emission peak red-shifted from 430 to 475 nm, which was consistent with the emission behavior at the concentration of $2 \times 10^{-2}$ in the solution and crystalline powder. The results indicated that BSBDS complexed with Al$^{3+}$ formed an excimer (Figure 5C). Furthermore, we had further studied the detection performance of BSBDS in the solid state for aluminum ions. First, the blank filter paper is soaked in a $2 \times 10^{-5}$ M BSBDS aqueous solution for 1 min and then dried. The solution...
containing different metal ions was filtered through a filter paper containing BSBDS. After drying, it is found that BSBDS can still detect metallic aluminum ions (Supplementary Figure S3).

Furthermore, the sensitivity is also an important factor for a chemical sensor. Then, different amounts of Al\(^{3+}\) (10 \(\mu\)L) were added sequentially to the BSBDS aqueous solution to estimate the detection limit of Al\(^{3+}\). Upon the addition of Al\(^{3+}\), the fluorescence intensity of BSBDS at around 430 nm gradually declined due to electron or energy transfer (Haldar et al., 2016) (Figure 6A and Supplementary Figure S4). Based on the results, a good linear curve was obtained in the range of 0.1–1.1 \(\mu\)M (Figure 6B), and the limit of detection (LOD) was found to be 2.09 \(\times\) 10\(^{-8}\) M according to computational method of \(3\sigma/k\) (Wang et al., 2008) (\(k\) are standard deviation of intercept and slope of the linear curve, respectively). It is worth noting that the detection limits for Al\(^{3+}\) in drinking water are 7.4 \(\times\) 10\(^{-6}\) M according to the World Health Organization (WHO) regulations (Dong et al., 2016). Therefore, BSBDS can be used as a convenient and sensitive fluorescent probe for the detection of Al\(^{3+}\) in drinking water. The dynamic quenching constant (\(K_q\)) is calculated via the Stern–Volmer formula (Dhingra et al., 2020) \(F_0/F = 1 + K_q\times\tau_0\times[Q] = 1 + K_{sv}\times[Q]\), where \(K_q\) is the dynamic quenching rate constant, \(\tau_0\) is the average lifetime of the lumiongens, and \([Q]\) is the concentration of Al\(^{3+}\). We got the function of \(y = 362113.11254x+0.99312\), so the quenching constant in Supplementary Figure S5 is 362113 L/mol.

### Cell Imaging

In order to explore the biological feasibility of BSBDS, cytotoxicity of BSBDS was evaluated by MTT assay. As shown in Supplementary Figure S6, BSBDS was non-cytotoxic to living cells. It means that BSBDS has good biocompatibility, and it is safe for cell-imaging or biological applications. In view of the non-toxic nature of BSBDS, intracellular detection of Al\(^{3+}\) was performed. HeLa cells were incubated with Nuclear Red™ LCS1 (nucleus-tracker, a commercial dye by AAT Bioquest®, Inc.) for 30 min, and then treated with 20 \(\mu\)M BSBDS for another 1 h. The confocal images of HeLa cells showed a certain amount of a fluorescent signal in pseudo colors with blue and green from BSBDS, which mainly appeared in the cytoplasm of HeLa cells; it was localized in different area with BSBDS (shown in Figures 7C–E). The other one is the experimental group which was preincubated with Al\(^{3+}\); it emitted intense blue fluorescence signal (Figure 7F) in comparison with Figure 7A; and weaker red fluorescence in the nucleus (Figure 7H) compared with the group preincubated without Al\(^{3+}\) (Figure 7C). The fluorescence signal can also be precisely observed in the nucleus in Figure 7H. We thus speculate that with the entry of Al\(^{3+}\) BSBDS emits strong light in both the cytoplasm and nucleus when excited by a 405-nm laser; Al\(^{3+}\) can also help BSBDS to enter into the nucleus. The results indicated that BSBDS is cell-permeable and could be used for dynamic imaging and Al\(^{3+}\) tracking in cells.

### CONCLUSION

In summary, we propose an ionic rod-shaped organic compound for the construction of high-efficiency DSE luminescent materials. Due to the large rod-shaped conjugation structure, BSBDS can emit light efficiently in the state of dilute solution. At the same time, the repulsive effect of ions at high concentrations can lead the rod-shaped molecules only form excimers locally, avoiding the aggregation quenching effect and achieving highly efficient luminescence in concentration and aggregation states.
BSBDS was selected as a model compound to validate the molecular design strategy, and the results showed that the compound possessed high-fluorescence quantum yield in both solution and solid states. In addition, BSBDS has recognition ability for Al\(^{3+}\) in the solution state, obvious mechanochromism properties in the solid state, and good biocompatibility, which can be used for cell imaging. The unique chemical structure characteristics and excellent optical properties of this ionic rod-shaped organic compound make it exhibit broad applications in many fields, such as organic optoelectronics, chemical sensing, ion detection, and biological imaging. This work will provide new ideas for the development of novel DSE materials.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding authors.

**AUTHOR CONTRIBUTIONS**

HL, XZ, and XQ: photophysical performance measurements and analyses, density function theory calculation, and manuscript preparation; SG: manuscript preparation and writing—review, and editing; HL, FC, and LW: cell culture, cytotoxicity, and cell imaging studies; CW, BJ, and YG: proposed the initial idea, designed the experiments, and were involved in conceptualization, supervision, funding acquisition, writing—review, and editing. All authors have read and agreed to the published version of the manuscript.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fchem.2022.807088/full#supplementary-material

**REFERENCES**

Belmonte-Vázquez, J. L., Amador-Sánchez, Y. A., Rodríguez-Cortés, L. A., and Rodríguez-Molina, B. (2021). Dual-State Emission (DSE) in Organic Fluorophores: Design and Applications. *Chem. Mater.* 33 (18), 7160–7184. doi:10.1021/acs.chemmater.1c02460

Chen G, G., Li, W., Zhou, T., Peng, Q., Zhai, D., Li, H., et al. (2015). Conjugation-Mediated Rigidity in Twisting Molecules: Filling the Gap between Aggregation-Caused Quenching and Aggregation-Induced Emission. *Adv. Mater.* 27 (30), 4496–4501. doi:10.1002/adma.201501981

Chen I, L., Wang, Y.-H., He, B., Nie, H., Hu, R., Huang, F., et al. (2015). Multichannel Conductance of Folded Single-Molecule Wires Aided by Through-Space Conjugation. *Angew. Chem. Int. Ed.* 54 (14), 4231–4235. doi:10.1002/anie.201411909

Dai, W., Liu, P., Guo, S., Liu, Z., Wang, M., Shi, J., et al. (2019). Triphenylquinoline (TPQ)-Based Dual-State Emissive Probe for Cell Imaging in Multicellular Tumor Spheroids. *ACS Appl. Bio Mater.* 2 (8), 3686–3692. doi:10.1021/acsabm.9b00396

Dhingra, D., Bhawna, P., and Pandey, S. (2020). Fluorescence Quenching by Nitro Groups in Ethylenediamine. *ACS Omega* 5 (1), 344–351. doi:10.1021/acsomega.9b01971

Henderson, G. (1977). The Effects of Absorption and Self-Absorption on Fluorescence Intensities. *J. Chem. Educ.* 54 (1), 57. doi:10.1021/ed054p57

Hilborn, R. C. (1982). Einstein Coefficients, Cross Sections, F Values, Dipole Moments, and All that. *Am. J. Phys.* 50 (11), 982–986. doi:10.1119/1.12937

Humphrey, W., Dalke, A., and Schulten, K. (1996). VMD: Visual Molecular Dynamics. *J. Mol. Graphics* 14 (1), 38–46. doi:10.1021/acs.jchemed.9b00596

Lu, T., and Chen, F. (2012). Multiwfns: a Multifunctional Wavefunction Analyzer. *J. Comput. Chem.* 33 (5), 580–592. doi:10.1002/jcc.22885

Luo, J., Xie, Z., Lam, J. W., Chen, L., Qin, C., et al. (2001). Aggregation-Induced Emission of 1-Methyl-1,2,3,4,5-Pentaphenylsilole. *Chem. Commun.* (Camb), 1740–1741. doi:10.1039/b105159a

Meng, Y., Guo, S., Jiang, B., Zhang, X., Zou, L., Wei, C., et al. (2021). Boosting the Humidity Resistance of Nonconventional Luminogens with Room Temperature Phosphorescence via Enhancing the Strength of Hydrogen Bonds. *J. Mater. Chem. C* 9 (27), 8515–8523. doi:10.1039/D1TC01271A

Raghuvanshi, A., Jha, A. K., Sharma, A., Umar, S., Mishra, S., Kant, R., et al. (2017). A Nonarchetypal 5,6-Dihydro-2H-Pyrano[3,2-G]indolizine-Based Solution-
Reineke, S., Lindner, F., Schwartz, G., Seidler, N., Walzer, K., Lüssem, B., et al. (2009). White Organic Light-Emitting Diodes with Fluorescent Tube Efficiency. *Nature* 459 (7244), 234–238. doi:10.1038/nature08003

Stoerkler, T., Frath, D., Jacquemin, D., Massue, J., and Ulrich, G. (2021). Dual-State Emissive π-Extended Salicylaldehyde Fluorophores: Synthesis, Photophysical Properties and First-Principle Calculations. *Eur. J. Org. Chem.* 2021 (26), 3726–3736. doi:10.1002/ejoc.202100650

Uoyama, H., Goushi, K., Shizu, K., Nomura, H., and Adachi, C. (2012). Highly Efficient Organic Light-Emitting Diodes from Delayed Fluorescence. *Nature* 492, 234–238. doi:10.1038/nature11687

Wang, H., Zhao, E., Lam, J. W. Y., and Tang, B. Z. (2015). AIE Luminogens: Emission Brightened by Aggregation. *Mater. Today* 18 (7), 365–377. doi:10.1016/j.mattod.2015.03.004

Wang, Y.-Q., Ye, C., Zhu, Z.-H., and Hu, Y.-Z. (2008). Cadmium telluride Quantum Dots as pH-Sensitive Probes for Tiopronin Determination. *Analytica Chim. Acta* 610 (1), 50–56. doi:10.1016/j.aca.2008.01.015

Yamasaki, K., Kimura, T., Masuya, H., and Tokue, I. (1997). A New Method for Eliminating Self-Absorption Effect from Dispersed Emission Spectra. *Rev. Scientific Instr.* 68 (10), 3697–3701. doi:10.1063/1.1148013

Zhang, Q., Li, B., Huang, S., Nomura, H., Tanaka, H., and Adachi, C. (2014). Efficient Blue Organic Light-Emitting Diodes Employing Thermally Activated Delayed Fluorescence. *Nat. Photon* 8 (4), 326–332. doi:10.1038/nphoton.2014.12

Zou, L., Guo, S., Lv, H., Chen, F., Wei, L., Gong, Y., et al. (2022). Molecular Design for Organic Luminogens with Efficient Emission in Solution and Solid-State. *Dyes Pigm.* 198, 109958. doi:10.1016/j.dyepig.2021.109958

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