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5, 7778Zinc(II)-heteroligand compounds for wet
processing OLEDs: a study on balancing
charge carrier transport and energy transfer†Emmanuel Santos Moraes, ^{‡ab} Luís Gustavo Teixeira Alves Duarte, ^{‡b}
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Organic light-emitting diodes (OLEDs) are one of the most studied and utilized optoelectronic components in display technology. However, their application in lighting remains limited due to materials costs and a guaranteed feasible deposition technique. To address this challenge, we explored the use of easily synthesized organic molecules capable of complexation with abundant transition metals to enhance their optoelectronic properties, coupled with low-cost wet processing protocols. Four zinc(II) coordination compounds were synthesized and the impact of incorporating two different ligands into a metal center was evaluated in terms of their optoelectronic properties. A photophysical investigation was made, encompassing emission and absorption analyses in both solid-state and thin film configurations. Förster resonance energy transfer (FRET) processes were performed using polyfluorene (PFO) and zinc(II) compounds in a host–guest system, revealing FRET efficiencies ranging from 10 to 68%, depending on the concentration of zinc(II) compounds in the PFO matrix. Subsequently, solution-processed OLEDs were fabricated using PFO:zinc(II) homo (**ZnL11** and **ZnL22**) and heteroligand (**ZnL13** and **ZnL23**) compounds as the emissive layer at a concentration of 1%, following a straightforward architecture, ITO|PEDOT:PSS|PVK|PFO:Zn(II)-compounds|TmPyPB|Ca|Al. The OLEDs achieved external quantum efficiencies (EQE) close to the theoretical limit of these active layers, ranging from 1.2% to 1.8%, with an applicable brightness value ($L > 100 \text{ cd m}^{-2}$), coupled with low roll-off in EQE values. Notably, the heteroligand coordination compounds exhibited superior device performance, attributed to their high electrical charge-carrier mobilities, trap-state profiles, and density of free carriers, as elucidated by space-charge shallow- (SCLC) and deep-trap (TCLC) transport models.

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1. Introduction

Organic light-emitting diodes (OLEDs) synthesized by solution-processing techniques have emerged as promising alternatives to traditional thermally evaporated devices for producing low energy-consuming large area emitters.¹ For solution-processed OLEDs, the active layers can be deposited by several wettable

methods, such as spin-coating,² slot-die coating,³ blade coating,⁴ and inkjet printing⁵ methods, offering scalable and straightforward implementation. However, there are several drawbacks to achieving a reasonable device performance. One is specially related to the design of active molecules with reasonable simple synthetic processes and with good emission properties; others are related to their solubility in commercial inexpensive solvents, which aids in achieving stable and uniform morphologies (without creation of intrinsic defects acting as energetic traps for electrical carriers), chemical and photochemical stabilities and reasonable charge mobility in the solid state.⁶ The best way to produce solution-deposited OLED active layers is still based on the host-guest concept.⁷

To disperse an emissive molecule in such active layers, it is critical to choose solubility-compatible host-guest materials, ensuring proper charge-carrier injection, transport, and recombination from the host matrix to the guest emissive target.^{8,9} To improve the efficiency of host-guest systems, the energy levels

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of the components must fulfill the requirements for Förster/Dexter energy transfer, creating cascade channels for charge mobility and energy transfer processes, alongside a good molecular conformation.^{10,11}

In recent decades, the external quantum efficiency (EQE) has increased substantially for solution-processed OLEDs.¹² For instance, for thermal-evaporated OLEDs, Wang *et al.* obtained a red device based on the Pt(II) phosphorescent compound, composed of five evaporated layers between the electrodes, which resulted in a good EQE of 31.8%.¹³ Additionally, Pt(II) complex solution-processed OLEDs, which possess simpler architectures (with three or two layers), achieved EQEs of 22.9% and 15.0%, respectively.^{14,15} Therefore, the production of stable materials for high-performance OLEDs must lead to the development of strategies toward the use of low-temperature protocols and simpler device architectures.

In this context, coordination compounds are a very wide class of OLEDs' active layers. They might be composed of two categories: compounds with prevalent emission of the ligand and those with inner transition metal emission, singlet emission, or triplet emission. Because these metal–ligand properties define the emission process, there are several combinations of the same metallic center with ligands to provide unique materials and properties. Coordination compounds, encompassing ligand-based and metal-based emissions, offer a wide range of possibilities for OLED active layers.^{16–19} However, although heavy metals, such as iridium(III),²⁰ platinum(II),¹⁵ osmium(III),²¹ ruthenium²² and rhenium²³ complexes, can exhibit good performance, their cost, toxicity, and availability in nature are the main drawbacks leading to the choice of abundant metals, such as zinc(II).²⁴

Zinc(II) complexes, which have versatile coordination geometries, have been demonstrated to be a promising candidates for active layer components, due to their desirable physical-chemical properties such as solubility, thermal stability, and fluorescence quantum yields.^{24–26} Despite their limitations in terms of internal quantum efficiency, zinc(II) complexes have already proven to achieve nearly 5% EQE values (the theoretical maximum values expected for such 1st generation emitters), almost close to the limit for singlet emission.²⁷ In general, Zn(II) compounds are stable, with good solubilities, and can be easily synthesized, and the emission originated from the ligand. Improvement of the device performance can be achieved by the optimization of the charge transport mobility.¹⁹

In this sense, and seeking for solution-processed materials, we report the preparation and characterization of four zinc(II) complexes with two ligands to determine the role of their chemical structures in the figures of merit of OLEDs with active layers containing such complexes as emitters. The ligands (L1 and L2) are based on a salicylidene derivative that differs in the insertion of a bromine atom, forming **ZnL11** and **ZnL22** (homoligand) compounds and **ZnL13** and **ZnL23** are compounds with one ligand L1 or L2 and a 2-(2'-tosylaminophenyl) benzothiazole moiety (L3), forming structures L13 and L23 (heteroligand) (Fig. 1). Due to the solubility issues in solution-processed OLEDs, to improve optoelectronic properties without inserting

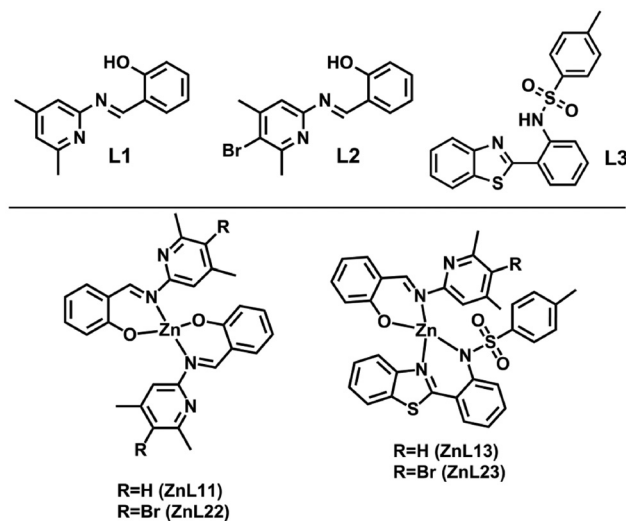


Fig. 1 Chemical structures of the ligands (L1, L2, and L3) and their corresponding complexes (**ZnL11**, **ZnL22**, **ZnL13**, and **ZnL23**).

another compound, two different ligands (L1–L3 and L2–L3) were used to improve the optoelectronic properties without changing the photophysical properties of the zinc(II) complex (L1–L1 and L2–L2). The optical properties of all these compounds were studied in polymethyl-methacrylate (PMMA), poly(9,9-di-*n*-octylfluorenyl-2,7-diyl) (PFO) thin films, and solid state. A theoretical approach based on density functional theory (DFT) further enhances the understanding of the optical properties.

By comparing both the photoluminescence (PL) and electroluminescence (EL) spectra of these four compounds, differences between the emissions in the inert PMMA and in the optically active PFO are interpreted in terms of the fluorescence resonance energy transfer (FRET) process by the Förster mechanism.¹⁶ For the PFO thin films, the molar ratios of the zinc(II) compounds to PFO were 0.1%, 0.5%, 1.0%, and 2.5% mol mol^{−1} to assess the solubility limit and processability of these host/guest systems, considering the FRET efficiency threshold. We built solution-processed OLEDs following a simple straightforward architecture. OLEDs based on PFO: Zn(II)-heteroligand coordination compounds have achieved better optoelectronic figures of merit and efficiency losses (roll-off <20%), compared to their homoligand structures, highlighting the **ZnL23** compound: $V_{on} = 8.5 \text{ V}@L_{max} = 536.8 \text{ cd m}^{-2}@EQE_{max} = 1.84\%@\eta_{Cmax} = 2.11 \text{ cd A}^{-1}@ \eta_{Pmax} = 0.49 \text{ lm W}^{-1}$. The obtained performance parameters of each Zn(II) coordination compound device exhibited EQE values close to those theoretically estimated, considering active layer internal quantum efficiencies (IQEs). The optoelectronic properties of OLEDs have been consistently correlated with photophysical and charge-carrier transport features, according to space-charge shallow- and deep-trap models. This research aimed to advance the development of stable materials for high-performance OLEDs by adopting low-temperature protocols and simpler device architectures, focusing on zinc



complexes as versatile active layer components, which can underscore their potential for achieving efficient OLEDs, contributing to the broader field of solution-processed optoelectronic devices. In summary, the achieved results show that Zn(II)-based emitters can be particularly useful for OLEDs' active layers despite the low EQE; however, the theoretical limit was reached with an appropriate Zn(II)-heteroligand structure (ZnL23), in a good trade-off between the simple device structure/efficiency and the good electrical charge balance.

2. Experimental

2.1. Synthesis of the ligands and zinc(II) complexes

All reagents were purchased from Sigma-Aldrich with >98% purity and used without further purification. Salicylidene ligands (L1–L3) were synthesized and characterized elsewhere.^{28,29}

For the complexation protocol, a mixture containing 1.0 mmol of the ligands (1 : 1) with 0.5 mmol of dihydrate zinc acetate in 30 mL of methanol was stirred under reflux for 2 hours. The product was obtained as a green-yellow precipitate, which was subsequently filtered, washed in cold methanol and hexane, and dried for 12 hours at 60 °C to afford the respective complexes ZnL11, ZnL22, ZnL13, and ZnL23 (Fig. 1). Single-crystals of the ZnL11, ZnL22 and ZnL13 compounds were obtained from the synthesis of crude products. The major difference between compounds L1 and L2 is the bromine in the *para* position.

2.2. Structural characterization

NMR spectra were recorded in DMSO-*d*₆ or CDCl₃ using a Bruker 250 MHz ¹H NMR spectrometer and ¹³C NMR spectra were recorded using a Bruker Advance 400 MHz spectrometer. Chemical shifts (δ) were given in parts per million from the peak of tetramethylsilane (δ = 0.00 ppm) as the internal standard in ¹H NMR or from the solvent peak of CDCl₃ (δ = 77.00 ppm) in ¹³C NMR. The original NMR spectra for all compounds are presented in the ESI† (Fig. S1–S8). Using an ATR mode, the FTIR spectra were acquired using a Cary 630 FTIR spectrometer from Agilent Technologies (Fig. S9, ESI†). High-resolution mass spectra in the positive mode were obtained using a Xevo G2-XS QTOF spectrometer (Fig. S10–S13, ESI†). Thermogravimetric analysis (TGA) was performed from 25 to 700 °C using a TA Instruments model 2950 instrument under a nitrogen atmosphere (100 mL min^{−1}) at a heating rate of 20 °C min^{−1} (Fig. S14, ESI†). The crystal structures were determined by single-crystal X-ray diffraction using a Bruker APEX II DUO area detector diffractometer, equipped with a low-temperature device (Oxford Cryosystems CRYOSTREAM 700). The data were collected at a crystal temperature of 150 K, using Cu Kα radiation (λ = 1.54184 Å; Incoatec microfocus X-ray source), based on a strategy of combining Ω and φ scans, a width of 0.5°, and an acquisition time of 30 s per frame. Cell refinement and data reduction were performed using SAINT and multiscan absorption correction was performed using SADABS-2014/5. Structures were solved by direct methods using

SHELXTL XT-2014/4 and refined by the least-squares method against F² using SHELXL^{30,31} with all nonhydrogen atoms refined anisotropically and all hydrogen atoms added geometrically.

ZnL11. Yield (66%). ¹H NMR (250 MHz, DMSO-*d*₆, ppm): 9.31 (s, 2H), 7.53, 7.34, 7.29, 6.87 (Ar-H, 12H), 2.66 (s, 3H), and 1.97 (s, 3H). ¹³C NMR (125 MHz, DMSO-*d*₆, ppm): 173.4, 167.5, 157.5, 150.5, 137.9, 123.7, 119.0, 112.8, 23.4, and 20.9. FTIR (ν_{max}, cm^{−1}): 2024–1660 (ν_{Ar}), 1600 (ν_{C=N}), 1430 (ν_{C-H}), and 1143 (ν_{C-O}). HRMS (ESI⁺): *m/z*, calculated for C₂₈H₂₇N₄O₂Zn [M + H]⁺ 515.1425; found 515.1414.

ZnL22. Yield (84%). ¹H NMR (250 MHz, DMSO-*d*₆, ppm): 9.46 (s, 2H), 7.81, 7.34, 7.03, 6.98 (Ar-H, 10H), 2.66 (s, 3H) and 2.28 (s, 3H). ¹³C NMR (300 MHz, CDCl₃, ppm): 164.7, 161.9, 156.8, 155.0, 149.7, 136.9, 128.5, 126.2, 119.9, 113.8, 29.7 and 25.6. FTIR (ν_{max}, cm^{−1}): 2015–1650 (ν_{Ar}), 1600 (ν_{C=N}), 1437 (ν_{C-H}), 1146 (ν_{C-O}), and 565 (ν_{C-Br}). HRMS (ESI⁺): *m/z*, calculated for C₂₈H₂₅BrN₄O₂Zn [M + H]⁺ 670.9636; found 670.9624.

ZnL13. Yield (61%). ¹H NMR (250 MHz, DMSO-*d*₆, ppm): 9.34 (s, 1H), 8.15, 7.65, 7.55, 7.40, 7.17, 6.87, 6.66 (Ar-H, 18H), 2.28 (s, 3H), 2.17 (s, 3H), and 1.97 (s, 3H). ¹³C NMR (125 MHz, DMSO-*d*₆, ppm): 173.5, 167.5, 157.4, 150.9, 137.9, 129.7, 127.5, 123.0, 120.3, 118.9, 114.7, 111.0, 23.4, 21.29, and 21.12. FTIR (ν_{max}, cm^{−1}): 2016–1640 (ν_{Ar}), 1600 (ν_{C=N}), 1438 (ν_{C-H}), 1135 (ν_{C-O}), and 1083 (ν_{C-N}). HRMS (ESI⁺): *m/z*, calculated for C₃₄H₂₉N₄O₃S₂Zn [M + H]⁺ 669.0973; found 669.0962.

ZnL23. Yield (77%). ¹H NMR (250 MHz, DMSO-*d*₆, ppm): 9.36 (s, 1H), 8.14, 7.68, 7.53, 7.40, 7.16, 6.93, 6.65 (Ar-H, 17H), 2.37 (s, 3H), 2.28 (s, 3H), and 1.93 (s, 3H). ¹³C NMR (125 MHz, DMSO-*d*₆, ppm): 173.7, 168.2, 155.8, 150.6, 129.6, 127.1, 122.8, 120.5, 114.9, 23.6, and 21.3. FTIR (ν_{max}, cm^{−1}): 2016–1650 (ν_{Ar}), 1600 (ν_{C=N}), 1437 (ν_{C-H}), 1134 (ν_{C-O}), 1083 (ν_{C-N}), and 563 (ν_{C-Br}). HRMS (ESI⁺): *m/z*, calculated for C₃₄H₂₈BrN₄O₃S₂Zn [M + H]⁺ 747.0078; found 747.0064.

2.3. Optical properties

The electronic absorption spectra were acquired with a Hewlett-Packard 8452A diode array spectrophotometer. Their steady-state fluorescence spectra were recorded with a Cary Eclipse Varian spectrofluorometer using the absorption maxima as the excitation wavelength and λ_{exc} = 375 nm to excite PFO in the PFO:zinc(II) complex. Fluorescence decays of the PFO:Zn(II) composites were recorded by time-correlated single-photon counting (TCSPC) using an Edinburg Analytical Instruments FL 900 spectrofluorometer with an MCP-PMT (Hamamatsu R3809U-50) with a pulsed laser operating at λ_{exc} = 370.8 nm (EPL-375, FWHM 10 nm, and pulse width 77.0 ps), and λ_{exc} = 404.2 nm (EPL-405, FWHM 10 nm, and pulse width 46.3 ps). The instrument response function (IRF) was recorded using a Ludox[®] scatter. At least 10 000 counts in the peak channel were accumulated for the lifetime determination. The emission decays were fitted as exponential functions using the software FAST as expressed in eqn (1):

$$I(t) = \sum_{i=1}^n B_i e^{-t/\tau_i} \quad (1)$$



where $I_{(i)}$ is the signal intensity sum of the electronically excited species weighted by B_i (preexponential factor) and with a life-time constant τ_i .

The confocal fluorescence microscopy (fluorescence microscopy) images of the PFO:Zn(II) films were obtained with a Leica SP5 confocal laser scanning microscope, using an HCX PL APO CS 10 \times 0.4 DRY objective.

2.4. Solution-processed OLEDs

OLEDs were fabricated on a glass substrate with a patterned indium tin oxide (ITO) ($20 \Omega \square^{-1}$) from Ossila B.V. company. The ITO was cleaned with Hellmanex[®] 2% (30 min), hot deionized water (15 min), acetone (5 min), and 2-propanol (5 min), *via* ultrasonic bath, followed by washing with hot deionized water, drying with $N_2(g)$ jet, and treating with UV-ozone cleaner (10 min). Poly(3,4-ethylene-dioxythiophene)-poly(styrene sulfonate) (PEDOT:PSS) was purchased from Clevis, H. C. Starck, and the thin film (30 nm) was deposited by dynamic spin-coating at 5000 rpm for 30 s, followed by thermal annealing at 120 $^{\circ}C$ for 15 min. PVK ($M_w = 1100000 \text{ g mol}^{-1}$) was purchased from Sigma-Aldrich. The solution was processed in dry tetrahydrofuran at 5 mg mL^{-1} and the thin film (80 nm) was obtained by spin-coating at 3000 rpm for 30 s and annealing at 80 $^{\circ}C$ for 10 min. Poly(9,9-di-*n*-octylfluorenyl-2,7-diyl) was supplied by American Dye Source, Inc. (ADS129BE, $M_w = 25000\text{--}150000 \text{ g mol}^{-1}$). The solution was processed in dry tetrahydrofuran at a final concentration of 10 mg mL^{-1} and spin-coated at 2000 rpm for 30 s. Active layers' thin films based on PFO:Zn(II)-homo and heteroligand compounds were obtained from the same PFO's procedure, from the host-guest system with 1% mol mol^{-1} of each Zn(II) coordination compound. All THF solutions were filtered with a Ossila's PTFE hydrophobic 0.10 μm . 1,3,5-Tris(3-pyridyl-3-phenyl)benzene (TmPyPB) was purchased from Sigma-Aldrich and deposited by thermal evaporation (50 nm) at a ratio of 0.1 \AA s^{-1} under a pressure of $\sim 10^{-6}$ mbar, and cathode calcium (10 nm) and aluminum (90 nm) were evaporated at 0.1 and 1.0 \AA s^{-1} respectively. The final device structure was ITO|PEDOT:PSS|PVK|PFO:Zn(II) compound (1% mol mol^{-1})|TmPyPB|Ca|Al.

The optoelectrical properties were analyzed by current-voltage-brightness (I - V - L) measurements using a 2400 Keithley Source Meter and a Konica Minolta LS-100 Luminance Meter, coupled with a close-up lens no. 110, $\varnothing = 40.5 \text{ mm}$, 10–20 cm. The EL spectra were acquired using a USB2000+ Ocean Optics diode array spectrometer. The CIE 1931 chromaticity coordinates as we as all OLEDs' efficiency calculations were done in a home-made software, assuming devices Lambertian electro-emission profile.

2.5. Computational details

Single-crystal X-ray structures of ZnL11, ZnL22 and ZnL23 were used to calculate the excited state. The ZnL23 structure was optimized in the ground state by density functional theory (DFT) after which its final geometry was determined. The predictions were performed using the PBE0³² functional and aug-cc-pVDZ^{33,34} basis set. The vertical excitation was obtained

via the same level of calculation using the time-dependent DFT (TD-DFT) method at the vacuum level. All calculations were performed with ORCA version 5.0.3,^{35,36} and visualized in Gabedit 2.5.1³⁷ and Avogadro 1.2³⁸ software.

3. Results and discussion

3.1. Structural properties

Fig. 2 shows the ZnL11, ZnL22 and ZnL13 refined structures obtained by single-crystal X-ray diffraction. The ZnL23 structure was optimized by DFT. The relevant structural data are summarized in Table 1. For all these compounds, dihedral torsion occurs due to the interaction of zinc with oxygen and nitrogen, which results in the formation of a pseudo five coordinate complex. Some similar zinc(II) compounds present tetrahedral structures with a pyridine moiety that interacts with the zinc atom distorting the tetrahedral arrangement with a torsion angle in the same range (between 70 $^{\circ}$ and 77 $^{\circ}$).³⁹

3.2. Optical properties

This study aims to evaluate the effect of substitution between two different ligands at a zinc(II) site. For this, the photophysics characterization was performed, focusing on the main properties of the zinc compounds. First, UV-vis spectra were obtained in the solid-state (drop-cast film deposition) and thin film (dispersion produced by the spin-coating technique). Considering that PFO is active in the visible spectra, the zinc complexes were dispersed in a spectroscopically inert "host" as PMMA, and the results are shown in Table 2.

According to our theoretical results, the L3 ligand had a greater bandgap energy than L1 and L2. In fact, the absorption spectra of ZnL11 and ZnL22 showed a small redshift in relation to those of ZnL13 and ZnL23 in the PMMA film (Fig. 3). Due to the small intensity of absorption, at 410–420 nm, the excitation spectra concerning the PL maxima were recorded. The maxima band of excitation is close to the first weak band of the

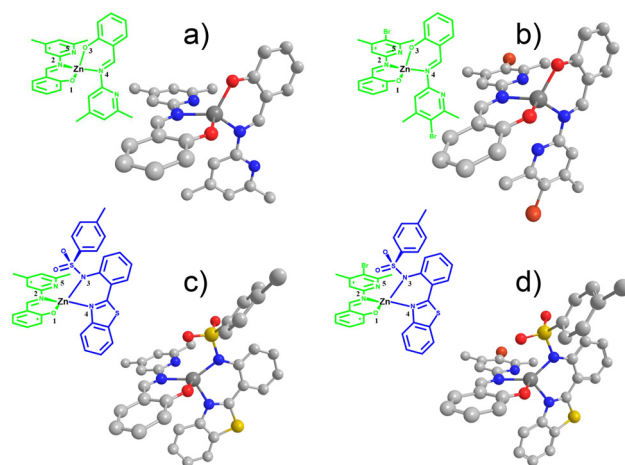


Fig. 2 The structures obtained by X-ray diffraction of ZnL11 (a), ZnL22 (b), ZnL13 (c) and ZnL23 (omitted hydrogen atom). (d) Optimized structures using PBE0/aug-cc-pVDZ at the vacuum level.



Table 1 Relevant distances and dihedral torsion angles between the ligands and zinc obtained via X-ray diffraction and DFT optimization using PBE0/aug-cc-pVDZ at the vacuum level. The distances are presented in angstroms (Å) and the torsion is presented in degrees (°)

Zinc(II) complex	Zn-1	Zn-2	Zn-3	Zn-4	Zn-5	Dihedral (1-2-3-4)
ZnL11	1.933	2.006	1.960	2.004	2.699	70.1
ZnL13	1.959	1.999	2.057	1.982	2.590	76.5
ZnL22	1.937	1.991	1.937	1.991	2.809	76.4
ZnL23^a	1.971	2.016	1.989	2.060	2.559	72.6

^a Obtained by a theoretical approach.

Table 2 Summary of optical data of **ZnL11**, **ZnL13**, **ZnL22** and **ZnL23** in PMMA films at 1% and in the solid state. λ_{abs} and λ_{em} (nm) are the absorption and emission maxima excited at 425 nm, respectively; SS is the Stokes shift (cm^{-1}), and τ is the lifetime (ns)

Zinc(II) complex	Condition	λ_{abs} (nm)	λ_{em} (nm)	SS (cm^{-1})	τ (ns)
ZnL11	PMMA	412	483	3568	2.4
	Solid-state	428	529	4461	4.1
ZnL13	PMMA	409	473	3308	2.7
	Solid-state	442	533	3863	5.9
ZnL22	PMMA	417	480	3147	2.1
	Solid-state	443	530	3705	1.8
ZnL23	PMMA	413	477	3249	3.0
	Solid-state	446	529	3518	2.8

absorption spectra. An explanation is that more excitation energy promotes the transition of electrons to a higher state than during emission and internal conversion occurs *via* a nonradiative pathway. In terms of fluorescence, **ZnL11** and **ZnL13** show a blue-shifted emission profile compared to the

ZnL22 and **ZnL23** ones. The photophysical differences between these molecules are linked to the presence of the bromine atom, which is a heavy atom, and may cause an energy shift in conjugated organic molecules.²⁹

The theoretical predictions showed that the first transition is a $\pi-\pi^*$ transition that occurs between the ligands. Due to the tetrahedral angle compared to the other one, the oscillator strength for the highest occupied molecular orbital (HOMO) \rightarrow lowest unoccupied molecular orbital (LUMO) for the first transition is small. Although the transitions were calculated at the vacuum level, the band positions were compared with the PMMA film and presented a small difference in energy (~ 0.1 eV). Additionally, the electronic density of the HOMO-1, HOMO, LUMO, and LUMO+1 of all the complexes is shown in Fig. S15 (ESI[†]). As mentioned, L3 has a greater bandgap energy than L1 and L2 which contributes to the first orbital levels and increases their energy gap regarding the **ZnL11** and **ZnL22** series (Table 3). In the electronic transitions, based on the X-ray structure, **ZnL11** has one of the N atoms directed to the zinc metal and another out of the molecule. This fact decreases the symmetry and causes a small energy splitting between both ligands in the complex. However, the first strong transition ($H_{-1} \rightarrow L, f = 0.29$) occurs in the same ligand. In parallel, **ZnL22** shows the same ligands position in the molecule, increasing the symmetry, and causing an $H \rightarrow L$ transition distributed throughout the molecule. For heteroligand Zn(II) complexes, the first transition occurs between $H \rightarrow L$ and $H_{-1} \rightarrow L$ for **ZnL13** and **ZnL23**, respectively. These transitions are mainly from L1 to L3 in **ZnL13** and from L3 to L2 in **ZnL23**. Along with these transitions, the difference between the dipole moment in the ground- and the excited state of the heteroligand is higher than the homoligand Zn(II) complex. It is worth mentioning that the difference between **ZnL22** and **ZnL23** is 16.3 Debye. Thus, **ZnL13** and **ZnL23** complexes showed a charge transfer character between the different ligands characterized by higher dipole moments regarding **ZnL11** and **ZnL22**.

In terms of solid-state absorption, all the complexes exhibited a spectral redshift in terms of the absorption in the PMMA film. The decrease in energy is partly due to aggregation in the structures. Fig. 3(c) shows the emission spectra in the solid-state of all the complexes. The emission spectra are similar, with green emission at approximately 530 nm, and redshift when compared to that of the PMMA film. Solid-state investigations are important for understanding OLED dynamics and

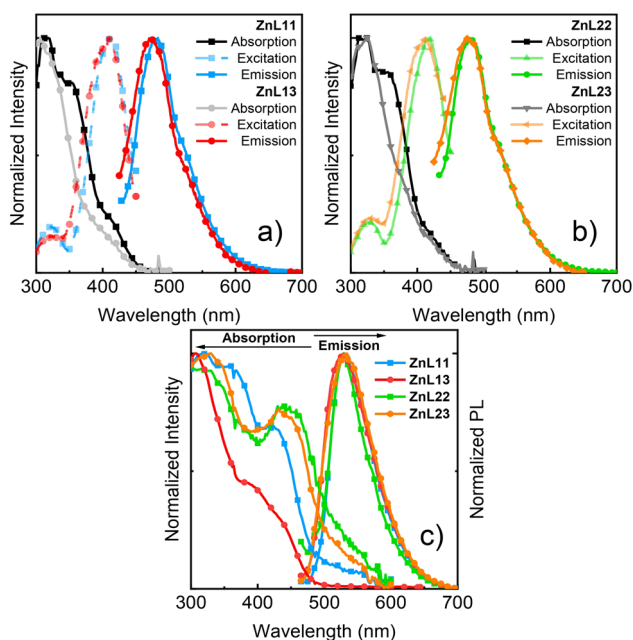


Fig. 3 Normalized absorption, excitation, and emission spectra of **ZnL11** and **ZnL13** (a) and **ZnL22** and **ZnL23** (b) in the 1% PMMA film at room temperature. The emission was excited by the maximum excitation wavelength. (c) Absorption and emission spectra in the solid-state of all complexes.

Table 3 Relevant data from the theoretical calculations of PBE0/aug-cc-pVDZ at the vacuum level. λ_{calc} is the calculated electronic transition, f is the oscillator strength and their assignment, $\Delta\mu$ is the difference of dipole moments between excited- and ground states, the value of the energies of the HOMO and LUMO, and the energy gap

Zinc(II) complex	λ_{calc} (nm)	f	Assignment	$ \Delta\mu $ (Debye)	HOMO (eV)	LUMO (eV)	E_{gap} (eV)
ZnL11	446	<0.001	$H \rightarrow L$ 99%	14.1	-5.7	-2.9	2.8
ZnL13	392	0.06	$H \rightarrow L$ 77%	15.4	-5.6	-2.4	3.2
ZnL22	417	0.03	$H \rightarrow L$ 82%	3.3	-5.7	-2.7	3.0
ZnL23	393	0.01	$H_{-1} \rightarrow L$ 89%	19.5	-5.9	-2.7	3.2

further possible emission from aggregation sites. The fluorescence lifetimes were longer for **ZnL11** and **ZnL13** than those for **ZnL22** and **ZnL23**. The heavy-atom effect causes a decay pathway by intersystem crossing and the lifetime is decreased.⁴⁰

3.3. Förster resonance energy transfer

The first approach characterized the photophysical properties of zinc(II) complexes in the inert host and solid state allows us to differentiate photophysical aspects in terms of steady-state and time-resolved emission features. Nevertheless, for solution-processed OLED devices, due to the low solubility of Zn(II) coordination compounds, these small molecules cannot be used alone to prepare good thin films. Therefore, it is necessary to employ a host–guest system mentioned before. The host must be a semiconducting material that can be easily wettable-processed, with the complementary energy levels of the zinc(II) complexes, *i.e.* the host must have large energy levels than the guest molecules, at same time transport charge-carriers.

The HOMO and LUMO energy levels below the vacuum of zinc(II) compounds were determined by DFT/TDDFT calculations, considering a theoretical error of about 0.1 eV. Besides this, their energy gaps are between the polyfluorene (PFO) HOMO and LUMO energetic levels (band gap).¹⁵ PFO is a well-known host in the literature and has an emission band at 430–440 nm. To investigate the efficiency of the energy transfer between the paired host/guest complex, the Förster resonance energy transfer (FRET) mechanism was used. The FRET condition involves an overlap between the emission band of the donor (host) and the absorption band of the acceptor (guest) within a short distance (1–10 nm) of the chromophores (Fig. S16, ESI†). Therefore, we prepared thin films by spin coating onto a glass substrate with pure PFO and PFO/zinc(II)-complexes at various solution ratios of 0.1, 0.5, 1.0, and 2.5% in the mol zinc(II) complex in PFO. Fig. 4(a) shows the emission of PFO with different ratios of **ZnL11**, exciting in the PFO chromophore (370 nm). The neat PFO thin film has a maximum emission band centered at 438 nm, related to the pure electronic emission (0-0) of the in-plane organized β -phase, with its well-defined vibronic progression at 460 (0-1) and 485 (0-2) nm. Besides this, a small contribution of the glassy PFO's α -phase pure electronic emission (0-0) around 425 nm was observed.⁴¹ In general, these emission bands decrease the intensity with the insertion of **ZnL11** and generates a broadband at 500 nm attributed to the zinc(II) complex emission.¹⁹ The emission shifts for the PMMA film are due to the chemical environment changes. The efficiency of FRET was measured by monitoring the lifetime of the PFO host. Fig. 4(b) and (c) show the emission decay and the decrease in the short term with the addition of **ZnL11** and **ZnL22** at ratios of 0.1, 0.5, 1.0, and 2.5%, respectively. The FRET efficiency (E_{FRET})⁴² was calculated using eqn (2), which determines the host emission quenching of the host in the presence of the guest.

$$E_{\text{FRET}} (\%) = \left(1 - \frac{\tau_{\text{acceptor}}}{\tau_{\text{donor}}}\right) \times 100 \quad (2)$$

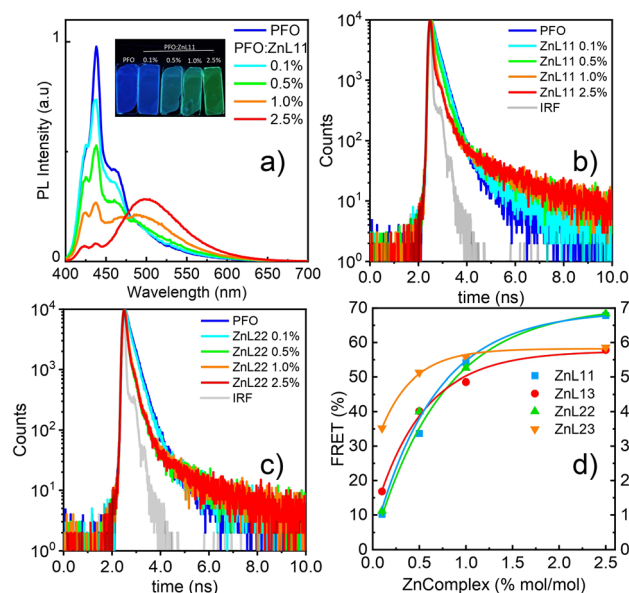


Fig. 4 (a) Fluorescence spectra of the thin films ($\lambda_{\text{exc}} = 375$ nm). (b) and (c) Emission decays of **ZnL11** and **ZnL22** in different host–guest ratios ($\lambda_{\text{exc}} = 370$ nm; $\lambda_{\text{em}} = 440$ nm). (d) FRET efficiency (%) for all complexes. Data from **ZnL22**, **ZnL13** and **ZnL23** are summarized in the ESI† (see Fig. S17 and S18).

where τ_{donor} is the lifetime of the pure PFO decay, and τ_{acceptor} is the lifetime of the mixture of PFO/Zn(II).

Fig. 4(d) shows the plot of the equation data concerning the zinc(II) complex ratio. The proportion of 1% between the zinc(II) complexes in PFO is lower than that between the other complex, with a good FRET efficiency of 30–45%. The maximum efficiency occurred at 2.5% at 58–68%; however, there was a small difference in the efficiency at a 1.0% ratio. Table S1 (ESI†) summarizes the principal FRET data.

A critical factor in solution-processed OLEDs is the deposited film morphology. Typically, a loss of ordered molecular conformation leads to a high formation of intrinsic defects that are responsible for unbalanced electrical charge densities (*via* effect in electrical mobilities) and/or losses by recombination. These effects are not only related to the emitter itself, but also the host plays an important role.^{43,44} In this sense, an investigation based on confocal microscopy has shown the solubility of the zinc(II) complexes in the PFO host. The images were obtained using an argon laser at 458 nm (Fig. 5), where Zn(II) coordination compounds are preferentially excited. It is possible to observe the formation of aggregated structures, following an increase in the concentration of Zn(II)-homo and hetero-ligand compounds. As a general trend, Zn(II) coordination compounds based on L1 (**ZnL11** and **ZnL13**) are more soluble than those based on L2 (**ZnL22** and **ZnL23**), which form aggregates even at low concentrations (0.5%), leading to great changes in the morphology with increasing concentrations. The FRET efficiency is intricately linked with this observation, and, beyond a 1.0% ratio, all zinc(II) complexes saturate the host, being unfavorable to make a good film for a solution-processed OLED.

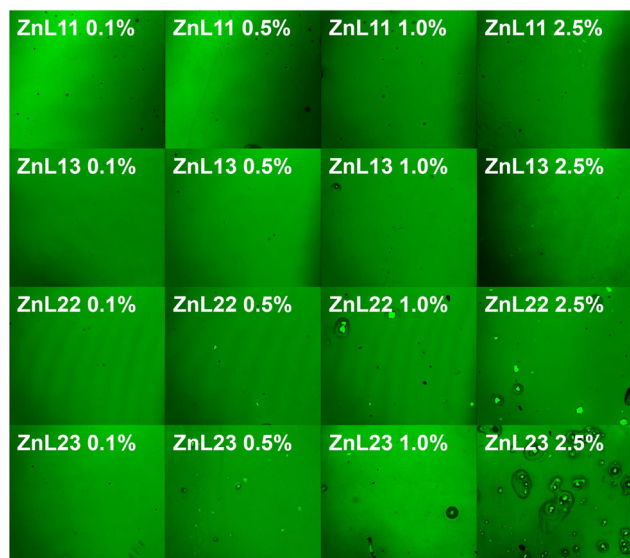


Fig. 5 Confocal microscopy images of PFO:Zn(II) coordination compounds (λ_{exc} = at 458 nm).

3.4. Solution-processed OLEDs' optoelectronic properties

Considering the photophysical and morphological characterization of PFO-based zinc-homo (ZnL11 and ZnL22) and heteroligand (ZnL13 and ZnL23) complex composites, including energy transfer studies, we successfully employed them as emissive layers (EMLs) in solution-processed OLEDs in

a proof of concept. Considering the FRET (Fig. 4(d)) efficiency and confocal microscopy images (Fig. 5), we chose a 1% (mol mol⁻¹) concentration of each zinc(II) coordination compound in the PFO semiconducting polymer matrix because 1% concentration is the limit threshold for FRET efficiency and does not strongly disturb the thin-film morphology of the composite.

The solution-processed OLEDs were assembled following the straightforward architecture previously described (scheme in Fig. 6(d)). PEDOT:PSS is a well-known metal-like conducting polymer blend that was used to optimize the ITO working function to -5.2 eV, while simultaneously acting as a hole injection layer.⁴⁵ High-molecular weight PVK is a hole transporting polymer ($\mu_{\text{h}} \approx 1 \times 10^{-6} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$) that has an E_{HOMO} of -6.1 eV and an E_{LUMO} of -1.2 eV,⁴⁶ which enables PVK to be a good buffer layer, coupled with hole transport and electron blocking characteristics, confining the electrons on the interface with the emissive layer. TmPyPB was used as both an electron transport layer and a hole blocking layer (E_{LUMO} of -2.7 eV and an E_{HOMO} of -6.8 eV). The electron mobility of TmPyPB ($\mu_{\text{e}} \approx 1 \times 10^{-3} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$)⁹ has the same magnitude as the PFO hole mobility,⁴⁷ while the PFO electron mobility is very small; therefore, the combination of PVK and TmPyPB among the EML will guarantee the electrical balance of the proposed device. Finally, Ca/Al was thermally evaporated to optimize the cathode working function to -2.9 eV. Table 4 summarizes the optoelectronic characteristics of all the solution-processed OLEDs.

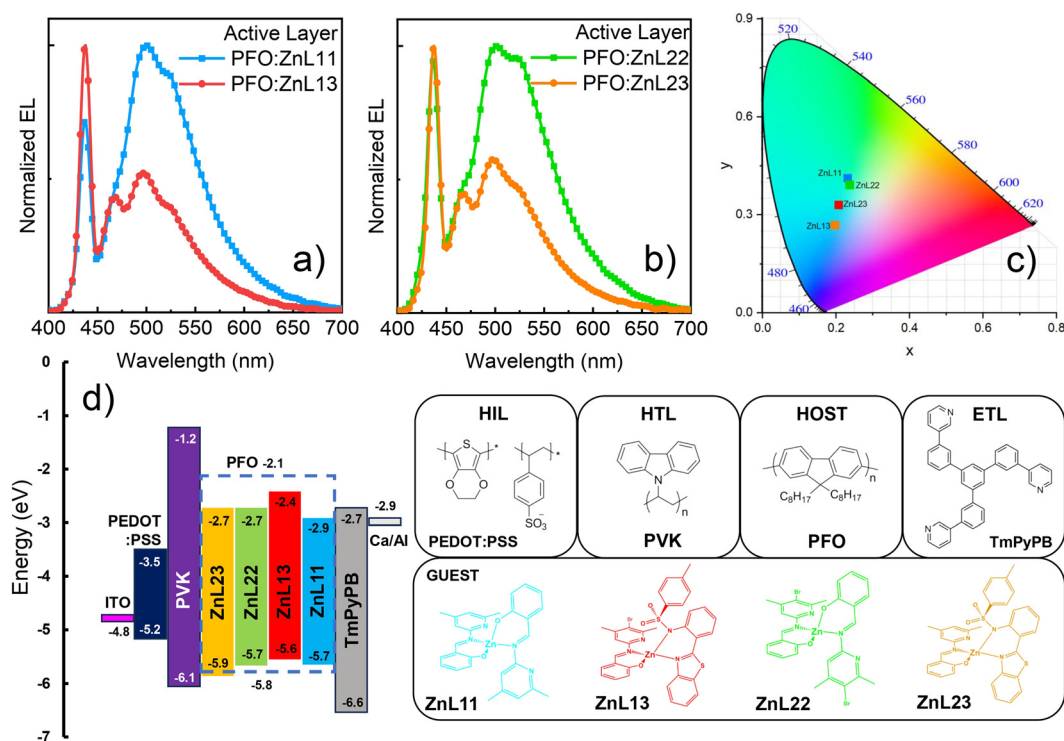


Fig. 6 Electroluminescence spectra of ZnL11 (blue curve) and ZnL13 (red curve) (a) and ZnL22 (green curve) and ZnL23 (orange curve) (b) in the PFO matrix (1% mol mol⁻¹), and the OLED chromaticity coordinate diagram CIE 1931 (c). Diagram level energy of each layer of the device and their chemical structures (d).

Table 4 Summary of the optoelectronic properties of PFO-based solution-processed zinc(II)-complex OLEDs

Optoelectronic properties	ZnL11	ZnL13	ZnL22	ZnL23
V_{on} (V)	12.5	9.5	9.0	8.5
L_{max} (cd m^{-2})	533.1	351.7	668.2	536.8
EQE (%)	1.40	1.38	1.21	1.84
EQE _{est} (%)	1.66	1.91	1.64	1.50
EQE (100 cd m^{-2})	1.25	1.30	1.06	1.55
R_{off} (%)	11	6	12	16
η_{C} (cd A^{-1})	2.08	1.58	1.81	2.11
η_{P} (lm W^{-1})	0.38	0.30	0.37	0.49
λ_{EL} (nm)PFO; zinc(II) complex	437; 501	437; 498	437; 502	437; 498
V_{TFL} (V)	9.60	10.96	9.91	10.67
N_{T} (cm^{-3})	3.18×10^{17}	3.63×10^{17}	3.28×10^{17}	3.54×10^{17}
E_{T} (meV)	189	215	265	257
μ_{eff} ($\text{cm}^2 \text{V}^{-1} \text{s}^{-1}$)	2.74×10^{-10}	8.46×10^{-10}	5.86×10^{-10}	2.82×10^{-10}
μ_{SCLC} ($\text{cm}^2 \text{V}^{-1} \text{s}^{-1}$)	1.97×10^{-7}	1.63×10^{-6}	3.76×10^{-7}	2.97×10^{-7}
θ_0	1.39×10^{-3}	5.21×10^{-2}	1.56×10^{-3}	9.51×10^{-4}
n_0 (cm^{-3})	4.43×10^{14}	1.90×10^{15}	5.13×10^{14}	3.37×10^{14}
μ_{TCLC} ($\text{cm}^2 \text{V}^{-1} \text{s}^{-1}$)	2.88×10^{-7}	2.57×10^{-7}	2.21×10^{-7}	7.92×10^{-7}

In general, the OLEDs' electroluminescence (EL) spectra exhibit a combination of zinc(II) complex compounds and PFO matrix electroemission (Fig. 6(a) and (b)). A similar behavior was observed in the steady-state PL measurements. However, **ZnL11** and **ZnL22** compounds ($\lambda_{\text{EL}} = 501$ and 523 nm, and $\lambda_{\text{EL}} = 502$ and 525 nm, respectively) have shown more intense EL bands in contrast to the zinc(II)-heteroligand compounds (**ZnL13** – $\lambda_{\text{EL}} = 496$ nm and 524 nm; **ZnL23** – $\lambda_{\text{EL}} = 498$ nm and 525 nm), when compared to the PFO EL profile ($\lambda_{\text{EL}}^{(0-0)} = 437$ nm and $\lambda_{\text{EL}}^{(0-1)} = 468$ nm). These features lead to solution-processed OLEDs based on the zinc(II)-homoligands exhibiting greenish-blue emission, while zinc(II)-heteroligands to exhibit cyan-blue emission, according to the CIE1931 color diagram (Fig. 6(c)). Additionally, it is interesting to note that the OLED EL colors are very close to the center of the CIE diagram, which can enable the application of these devices in lighting technology (some specific lighting/decorative and signage applications), besides the usual display applications.

The observed behavior should be originated from several different factors, including, for instance, different materials solubility, energy level diagram compatibility, spectral overlap between PFO emission and zinc(II) complex absorption, complex PLQYs, device charge-carrier transport and recombination efficiency, electrical carrier injection balance and intrinsic defects, in which some of them can alter the recombination region in the organic layers of the device. From photophysical, FRET and confocal microscopy studies, no direct correlation could be observed when the EL features were contrasted, with the exception of the energy level alignment of each zinc(II) complex compound with the PFO matrix (Fig. 6(d)). To better understand the solution-processed PFO-based composite framework, we focused on the optoelectronic properties of the devices, in terms of their figures of merit, as well as charge-carrier transport characteristics under space-charge conditions (Fig. 7).

Fig. 7 presents the overall OLEDs' figures of merit based on the zinc(II)-homo and heteroligand complexes in the PFO-matrix as EMLs, particularly current density and brightness *versus* voltage curves (Fig. 7(a)), external quantum, current and

power efficiencies (EQE, η_{C} and η_{P} , respectively) *versus* brightness curves (Fig. 7(b)), and DC charge-carrier transport characteristics (Fig. 7(c)).

In general, the **ZnL11** and **ZnL22** compounds presented high brightness and driving voltage values in contrast to those of the **ZnL13** and **ZnL23** complexes, according to the sequence **ZnL11** > **ZnL22** > **ZnL13** > **ZnL23**. On the other hand, the heteroligand compounds exhibited better performance parameters in terms of all the figures of merit, with **ZnL23** > **ZnL13** \approx **ZnL11** > **ZnL22**. From our data, zinc(II) complexes presents the following figures of merit: **ZnL23** – $V_{\text{on}} = 8.5$ V@ $L_{\text{max}} = 536.8 \text{ cd m}^{-2}$ @EQE_{max} = 1.84%@ $\eta_{\text{Cmax}} = 2.11 \text{ cd A}^{-1}$ @ $\eta_{\text{Pmax}} = 0.49 \text{ lm W}^{-1}$;

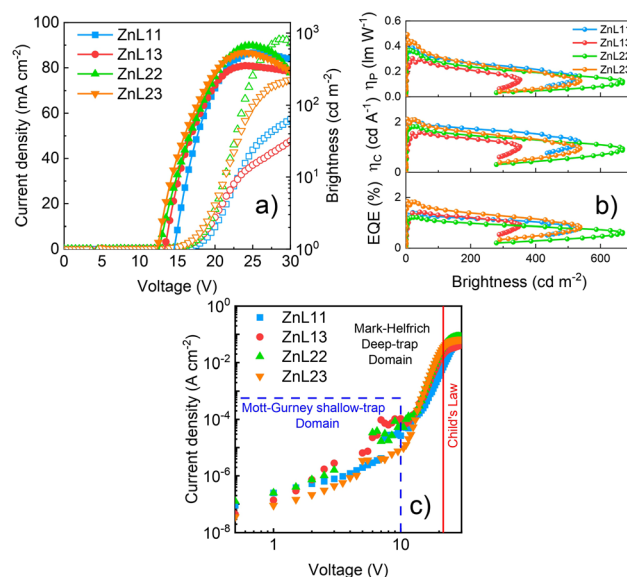


Fig. 7 Current density and brightness *versus* voltage curves (a), figures of merit *versus* brightness (roll-off) curves (b), and DC charge-carrier transport characteristics, according to $J \times V$ (log-log) curves, and current density *versus* voltage curve and the shallow-trap and deep-trap domain, and Child's Law (c) of the PFO-based zinc(II)-homo and heteroligand compound OLEDs.



ZnL13 – $V_{\text{on}} = 9.5 \text{ V}$, $L_{\text{max}} = 351.7 \text{ cd m}^{-2}$, $\text{EQE}_{\text{max}} = 1.38\%$, $\eta_{\text{Cmax}} = 1.58 \text{ cd A}^{-1}$, $\eta_{\text{Pmax}} = 0.30 \text{ lm W}^{-1}$; **ZnL11** – $V_{\text{on}} = 12.5 \text{ V}$, $L_{\text{max}} = 533.1 \text{ cd m}^{-2}$, $\text{EQE}_{\text{max}} = 1.40\%$, $\eta_{\text{Cmax}} = 2.08 \text{ cd A}^{-1}$, $\eta_{\text{Pmax}} = 0.38 \text{ lm W}^{-1}$; and **ZnL22** – $V_{\text{on}} = 9.0 \text{ V}$, $L_{\text{max}} = 668.2 \text{ cd m}^{-2}$, $\text{EQE}_{\text{max}} = 1.21\%$, $\eta_{\text{Cmax}} = 1.81 \text{ cd A}^{-1}$, $\eta_{\text{Pmax}} = 0.37 \text{ lm W}^{-1}$. Additionally, it is important to note that the obtained experimental EQE values of these solution-processed OLEDs are very close to the estimated ones, considering an approximation of the emissive layer PLQY. As far as we know, these are the best results that have been achieved for zinc(II) salicylidene coordination compounds as emissive layers in electroluminescent diodes.^{19,26,48–51}

Another important highlight that should be noted is that the OLEDs have a very low roll-off (R_{off}) of the EQE (Fig. 6(b)) as the brightness increases until $L = 100 \text{ cd m}^{-2}$, with **ZnL13** – $R_{\text{off}} = 6\% < \text{ZnL11}$ at $R_{\text{off}} = 11\% < \text{ZnL22}$ at $R_{\text{off}} = 12\% < \text{ZnL23}$ at $R_{\text{off}} = 16\%$. These behaviors could be directly linked to the fact that the **ZnL13** and **ZnL23** complexes can easily undergo charge transfer, as predicted by our DFT and TD-DFT simulations, which can facilitate the charge carriers' transport toward the emissive layer; in contrast to zinc(II)-homoligand compounds, although, as previously mentioned, the molecular conformation in the active layer can induce intrinsic defects acting as traps for the electrical carriers. Despite, and at least for **ZnL23**, confocal microscopy images do not predict such a hypothesis, and the nanodomains of defects can exist.

To support such interpretations, we employed space-charge models in the DC current density *versus* voltages curves in shallow- (Mott–Gurney's law), deep- (Mark–Helfrich's law) and free-trap domains (Child's law) (Fig. 7(c); fit details on the ESI† Fig. S19–S22). It is well stated that space-charge models can be applied to obtain the charge-carrier mobilities when only one type of charge drifts along the device structure (electron- or hole-only diode).⁵² Despite this, an estimate of the global electrical mobility could be achieved by applying these models in an OLED, although it has ambipolar electrical injection, transport and recombination.¹⁵ At low driving voltages, during the shallow-trap filling process, the current density depends on a quadratic power with an applied voltage, also known as space-charge limited current (SCLC) behavior, as described by the Mott–Gurney equation^{53–55}

$$J_{\text{SCLC}} = \frac{9}{8} \varepsilon \varepsilon_0 \mu_{\text{eff}} \frac{V^2}{d^3} \quad (3)$$

where J_{SCLC} is the current density at the SCLC domain, ε is the semiconducting layer dielectric constant, ε_0 is the free-space electrical permittivity, μ_{eff} is the effective electrical mobility at the SCLC shallow-trap region, V is the applied voltage, and d is the semiconductive layer thickness. During this process, all obtained OLEDs present very low effective electrical charge-carrier in the magnitude of $10^{-10} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$, which indicates that the devices may have a considerable density of trap-states, coupled with a small density of free-carriers, simultaneously with a charge-carrier transport governed by the minor carrier (in our case, electrons). SCLC effective electrical mobilities in the descending order are: $\mu_{\text{eff}}^{(\text{ZnL13})} > \mu_{\text{eff}}^{(\text{ZnL22})} > \mu_{\text{eff}}^{(\text{ZnL23})} \approx \mu_{\text{eff}}^{(\text{ZnL11})}$. Applying the SCLC model at the trap-free domain,

Mott–Gurney's law becomes Child's law and trap-free electrical mobility can be rewritten as follows:⁵⁵

$$\mu_{\text{eff}} = \mu_{\text{SCLC}} \theta_0 \quad (4)$$

where the trap-density parameter (θ_0) relates the density of free carriers (n_0) to the total density of trap states (N_T) and it is described as follows:

$$\theta_0 = \frac{n_0}{n_0 + N_T} \quad (5)$$

After the trap-filling process, the electrical mobilities of all the OLEDs increase by approximately 10^3 orders of magnitude, $\mu_{\text{SCLC}}^{(\text{ZnL22})} = 3.76 \times 10^{-7} > \mu_{\text{SCLC}}^{(\text{ZnL23})} = 2.97 \times 10^{-7} > \mu_{\text{SCLC}}^{(\text{ZnL11})} = 1.97 \times 10^{-7} > \mu_{\text{SCLC}}^{(\text{ZnL13})} = 1.63 \times 10^{-7} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$. Looking only for those values, in comparison with the shallow trap-filling electrical mobility ones, no absolute conclusive correlation with device figures of merit can be clearly observed. However, correlating OLED figures of merit with the θ_0 parameter, it is clear that the density of trap-states and free-carriers impact, in our case, the electroemissive device efficiency loss (R_{off}), following the same trend (see Table 4). To obtain more insights about this, we determine the total density of trap-states (N_T) of each kind of OLED by applying the trap-filled-limit voltage (V_{TFL}) relationship:⁵⁵

$$V_{\text{TFL}} = \frac{q N_T d^2}{2 \varepsilon \varepsilon_0} \quad (6)$$

where q represents the elementary charge. V_{TFL} corresponds to the point where Mott–Gurney's law intersects the deep-trap filling process behavior in $J \times V$ data (also known as trapped-charge limited current – TCLC), during the device's operation. This behavior can be described by following the Mark–Helfrich equation:^{55,56}

$$J_{\text{TCLC}} = q^{1-l} \mu_{\text{TCLC}} n_0 \left(\frac{2l+1}{l+1} \right)^{l+1} \left(\frac{l}{(l+1)N_T} \right)^l \frac{V^{l+1}}{d^{2l+1}} \quad (7)$$

where l is the temperature (T)-dependent trap-energy (E_T) parameter, which can be expressed as $\left(l = \frac{E_T}{k_B T} \right)$, and μ_{TCLC} is the electrical mobility during the deep-trap filling. Considering these parameters and combining eqn (5) and (6), we can also estimate n_0 , and finally, determine μ_{TCLC} values for each kind of device.

Taking into account the obtained values for n_0 and μ_{TCLC} (Table 4), we can observe some correlations with the OLEDs' figures of merit: (i) n_0 values follow the same trend with devices roll-off, as observed for θ_0 values; and (ii) the electrical mobility during the deep-trap filling process of each OLED has shown a correlation with the devices figures of merit in terms of EQE, η_c and η_p . In other words, OLEDs with better optoelectronic properties exhibit high μ_{TCLC} values, while those with lower efficiency losses with brightness increments have high density of free-carriers. Although expected, our data seem to follow a well trend.

4. Conclusions

In summary, our study reports the synthesis and evaluation of four zinc(II) complexes with two newly synthesized and



characterized ligands, aiming to modulate their photophysical and optoelectronic properties for applications in solution-processed OLEDs. Our approach involved the strategic combination of different ligands forming Zn(II) coordination compounds in a composite with PFO in 1% minimizing the morphological changes in the thin film. Theoretical predictions indicated that the L3 ligand possessed a higher bandgap energy compared to L1 and L2, resulting in a blue emission when complexed with zinc(II), thereby creating a wider energy gap relative to the green emission of **ZnL11** and **ZnL22**. Consequently, while there were no significant alterations observed in the photophysical properties, slight modifications were noted in the optoelectronic characteristics of the complexes. Notably, our study revealed an increase in external quantum efficiency (EQE) associated with enhancements in electrical mobility and trap-density parameters. This underscores the effectiveness of our novel approach in utilizing zinc(II) complexes with different ligands to fine tune the optoelectronic properties, thus offering a new perspective on their application in OLEDs. In essence, our findings present a promising framework for the rational design of zinc(II) complexes, wherein the insertion of distinct ligands can effectively tailor optoelectronic properties while maintaining morphological integrity. This approach opens new possibilities for the development of solution-processed OLEDs with improved performance and versatility, contributing to the advancement of lighting technologies and beyond.

Author contributions

E. S. M.: conceptualization, formal analysis, investigation, writing – original draft, and writing – review and editing; L. G. T. A. D.: conceptualization, investigation, writing – original draft, and writing – review and editing; F. S. R.: supervision, writing – original draft, and writing – review and editing; J. C. G.: supervision, conceptualization, formal analysis, investigation, writing – original draft, and writing – review and editing. L. F. R. P.: funding acquisition, project administration, resources, software assembly, and writing – review and editing. T. D. Z. A.: funding acquisition, project administration, resources, and writing – review and editing.

Data availability

The data supporting this article have been included as part of the ESI.†

Conflicts of interest

The authors declare no conflicts of interest.

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