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Rapid room temperature synthesis of red iridium(III) complexes containing a four-membered Ir–S–C–S chelating ring for highly efficient OLEDs with EQE over 30%†

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Three red cyclometalated iridium(III) complexes (4tfmpq)₂Ir(dipdte), (4tfmpq)₂Ir(dpdtc) and (4tfmpq)₂Ir(Czdtc) (4tfmpq = 4-(4-(trifluoromethyl)phenyl)quinazoline, dipdte = *N,N*-diisopropyl dithiocarbamate, dpdtc = *N,N*-diphenyl dithiocarbamate, and Czdtc = *N*-carbazoyl dithiocarbamate) containing the unique four-membered Ir–S–C–S backbone ring were synthesized in five minutes at room temperature with good yields, and the Gibbs free energy calculation results indicate that all reactions are exothermic and thermodynamically favorable processes. The emission colors (λ_{peak} = 641–611 nm), photoluminescence quantum efficiencies (Φ_{p} = 58.3–93.0%) and bipolar properties can be effectively regulated by introducing different electron-donating substituents into the dithiocarbamate ancillary ligands. Employing these emitters, organic light emitting diodes (OLEDs) with double emissive layers exhibit excellent performances with a maximum brightness over 60 000 cd m^{−2}, a maximum current efficiency of 40.68 cd A^{−1}, a maximum external quantum efficiency (EQE_{max}) of 30.54%, and an EQE of 26.79% at the practical luminance of 1000 cd m^{−2}. These results demonstrate that Ir(III) complexes with sulfur-containing ligands can be rapidly synthesized at room temperature, which is key to the production of metal luminescent materials for large-scale application in highly efficient OLEDs.

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Introduction

Highly efficient phosphorescent cyclometalated complexes have been widely employed to fabricate organic light-emitting diodes (OLEDs) because of their unique optoelectronic properties

including relatively short triplet excited-state lifetime, high phosphorescence quantum yields and excellent color tunability.¹ Especially, iridium(III) complexes are the most important emitters, and the demand for phosphorescent materials during OLED industrialization is particularly large.² In order to speed up the process of OLED industrialization, cost reduction is extremely important. But for the production of most Ir(III) complexes, the last reaction process is always to reflux the [(C[^]N)₂Ir(μ -Cl)]₂ chloride-bridged dimer with cyclometalated or ancillary ligands at high temperature for a long time, which would inevitably increase the manpower and cost.³ It is important, but difficult to find suitable ligands which can form phosphorescent complexes at room temperature efficiently to reduce the cost of OLED industrialization significantly.⁴

For the previously reported Ir(III) complexes, ancillary ligands such as acetylacetone (acac) or 2-pyridinecarboxylic acid (pic) with the central iridium atom mainly formed six or five-membered ring structures. But Ir(III) complexes containing four-membered ring structures make the angle formed by the two coordinated atoms of the ancillary ligand and the iridium atom reduced, suggesting that four-membered metallocycles would possess greater coordination strain energy.⁵ However, four-membered ring structures of Ir(III) complexes with sulfur

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† Electronic supplementary information (ESI) available: Details of materials, measurements, X-ray crystallography, electrochemical tests and theoretical calculations. Procedures of OLED fabrication and measurements. The crystallographic data, electronic cloud density distributions, TG curves, cyclic voltammograms and lifetime curves of Ir(III) complexes. The flexible scanning of the Ir–S bond length. The calculated free energy changes ΔG of formation of the two S–Ir coordination bonds. The isodensity surface plots and HOMO/LUMO orbital levels. Current efficiency versus luminance of devices with different doped concentrations. ¹H NMR and ¹⁹F NMR spectra. CCDC 1832334 and 1832357. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c8sc05605f

‡ Lu and Su have the same contributions to this paper.

atoms in the ancillary ligands are rarely studied in OLEDs,⁶ and there is no new progress in the research of Ir(III) complexes containing the four-membered ring based on the Ir–S–C–S backbone. Moreover, these ancillary ligand derivatives have strong selectivity to the cyclometalated ligand for efficient Ir(III) complexes. Furthermore, in comparison with those well-developed green Ir(III) complexes, efficient red emitters are limited and the performances of red OLEDs are still not satisfactory owing to the energy gap law.⁷ For example, Suh *et al.* reported an Ir(mphmq)₂(tmd) based high-performance device with a maximum EQE of 24.3% and the CIE coordinates of (0.65, 0.35).^{7f} Lately, Yang *et al.* reported devices using (ptq)₂Ir(acac) with the CIE coordinates of (0.61, 0.36) and a maximum EQE of 22.9%.^{7g} Cho *et al.* reported a thiophene-phenylquinoline-based Ir(III) complex (Th-PQ)₃Ir and the corresponding OLEDs achieved a maximum EQE of 21.3% with an EL emission peak at 612 nm adopting solution-processed technology.^{7h} Recently, Chi *et al.* reported a bis-tridentate Ir(III) complex of [Ir(mimb)(L3)] and the corresponding devices achieved the highest results of a maximum EQE of 27.4% with CIE coordinates of (0.63, 0.38).⁷ⁱ

With this consideration, three sulfur atom containing dithiocarbamate derivatives constructed with different electron-donating groups (dipdte = *N,N*-diisopropyl dithiocarbamate, dpdte = *N,N*-diphenyl dithiocarbamate, and Czdtc = *N*-carbazolyl dithiocarbamate) as ancillary ligands were developed for Ir(III) complexes (4tfmpq)₂Ir(dipdte), (4tfmpq)₂Ir(dpdte) and (4tfmpq)₂Ir(Czdtc) using the electron-deficient 4-(4-(trifluoromethyl)phenyl)quinazoline (4tfmpq) moiety as the cyclometalated ligand (Scheme 1) in this paper. Diphenylamine and carbazole are very classical hole transport units in OLED hole transport and host materials. The application of diphenylamine

and carbazole in phosphorescent materials is more conducive to the balance of the distribution of holes and electrons, and is crucial for improving device efficiency and reducing roll-off. Moreover, the stable dithiolate compounds can reduce the work function of the emitter and the turn-on voltage in OLEDs.⁸ In addition, due to the nitrogen frame of the 4tfmpq ligand, these complexes would have bipolar properties, which are beneficial for their device performances.⁹

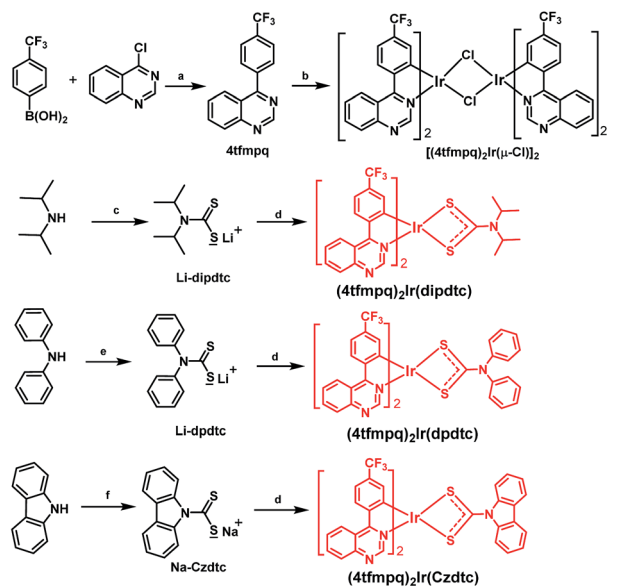
Results and discussion

Preparation and characterization of complexes

As shown in Scheme 1, three Ir(III) complexes were synthesized by the reaction of the [(C[^]N)₂Ir(μ-Cl)]₂ chloride-bridged dimer with dithiocarbamate derivatives. The four-membered heterocycle containing dithiocarbamate derivatives in (4tfmpq)₂Ir(dipdte), (4tfmpq)₂Ir(dpdte) and (4tfmpq)₂Ir(Czdtc) were accessed by employing electron-rich units with bulkier backbone substituents to favor *S,S*-chelation of the electron-deficient Ir(III) center.^{5,6,10} Thus, the isopropyl, phenyl and carbazolyl moieties tend to push the lone pairs of electrons on the nitrogen atoms toward the metal center, which could greatly favor the chelating bonding mode.^{5,6,11} All complexes can be synthesized at room temperature in 5 min with high yields above 70% and good thermal stability (with decomposition temperature above 370 °C, Table 1 and Fig. S1†) owing to the strong coordination capability of the sulfur atom with the iridium atom, which is also helpful to boost their sublimation yields (>90%). Because most Ir(III) complexes were prepared at high temperature for a long time, the sulfur atom containing ligands developed in this study can reduce the cost of materials and OLED industrialization significantly.

To further explore the reason for the rapid synthesis of these Ir(III) complexes, density functional theory (DFT) calculations at the B3LYP/Lan12DZ^{12,13} level with the solvent effect treated using a SMD model¹⁴ were conducted. The calculated Gibbs free energy changes, Δ*G*, indicate that all three reactions are exothermic and thermodynamically favorable processes (from −59.4 to −94.9 kJ mol^{−1}, Fig. 1). Assuming the formation of the two Ir–S bonds is stepwise, the kinetics of the complex formation by flexibly scanning the Ir–S bond length for the reaction of (4tfmpq)₂Ir(dipdte) were examined. The obtained potential energy surfaces show that as the S and Ir approach each other, the energy of the system gradually decreases until the equilibrium S–Ir bond length is reached. Further reduction of the S–Ir distance leads to a sharp energy increase (Fig. S2†). Hence, the formation of the S–Ir coordination bond does not involve the transition state of the conventional sense, and it is not necessary to overcome the corresponding energy barrier. The calculated free energy changes Δ*G* (−42.4 kJ mol^{−1}, Fig. S3†) further demonstrate that the formation of the first Ir–S bond is thermodynamically favorable. The formation of the second Ir–S bond is similar to the first one, both kinetically and thermodynamically (−60.6 kJ mol^{−1}, Fig. S3†).

Molecular structures by the ORTEP drawing of the Ir(III) complexes (4tfmpq)₂Ir(dipdte) and (4tfmpq)₂Ir(Czdtc) are shown in Fig. 2, and the crystallographic data and the selected



- (a) Pd(PPh₃)₄, Na₂CO₃, THF–H₂O, 70°C, 12h; (d) EtOCH₂CH₂OH, rt, 5 min;
 (b) IrCl₃, EtOCH₂CH₂OH, 115°C, 12h; (e) *n*-BuLi, CS₂, 0°C, 12h;
 (c) LiOH, CS₂, benzene–H₂O, rt, 2h; (f) NaH, CS₂, 40°C, 3h;

Scheme 1 The synthetic routes to the ligands and complexes.



Table 1 Photophysical data of (4tfmpq)₂Ir(dipdte), (4tfmpq)₂Ir(dpdtc) and (4tfmpq)₂Ir(Czdtc) in CH₂Cl₂ solution

Complex	<i>T</i> _d ^a (°C)	Absorption ^b (λ nm)	Emission ^b (λ _{max} nm)	τ _{298 K} ^b (μs)	Φ _p ^c (%)	<i>E</i> _T ^d (eV)	HOMO/LUMO ^e (eV)
(4tfmpq) ₂ Ir(dipdte)	384	263/285/453/592	641	1.41	58.3	2.04	−5.44/−3.19
(4tfmpq) ₂ Ir(dpdtc)	376	261/324/443/543	628	1.55	84.3	2.07	−5.50/−3.20
(4tfmpq) ₂ Ir(Czdtc)	371	282/357/532/570	611	1.50	93.0	2.18	−5.61/−3.22

^a Decomposition temperature. ^b Measured in degassed CH₂Cl₂. ^c Φ: emission quantum yields were calculated with the *fac*-Ir(ppy)₃ standard in degassed CH₂Cl₂ solution (Φ_p = 0.4). ^d *E*_T (triplet energy): estimated from the highest-energy peaks of the 77 K phosphorescence spectra in CH₂Cl₂ (10^{−5} M). ^e HOMO (eV) = −(*E*_{ox} − *E*_{1/2,Fc}) − 4.8, LUMO (eV) = HOMO + *E*_{bandgap}.

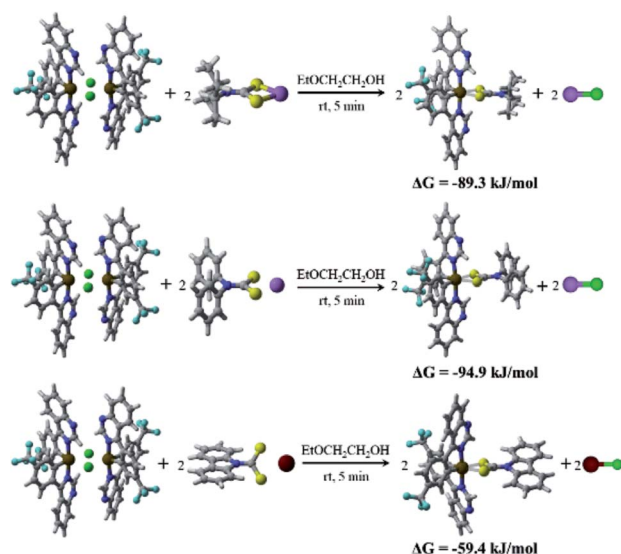


Fig. 1 The calculated Gibbs free energy change Δ*G* of the three reactions of (4tfmpq)₂Ir(dipdte), (4tfmpq)₂Ir(dpdtc) and (4tfmpq)₂Ir(Czdtc) complexes.

bond lengths/angles are summarized in Tables S1 and S2 (ESI[†]), respectively. Both complexes have a distorted octahedral coordination geometry around the iridium center surrounded by three chelating ligands with *cis*-C–C and *trans*-N–N dispositions in which the N–Ir–N angles for these complexes are all almost 175°. The four-membered Ir–S–C–S backbones could lead to significantly more acute S–Ir–S bite angles (71.4° and 72.0°) than seen for the larger five-

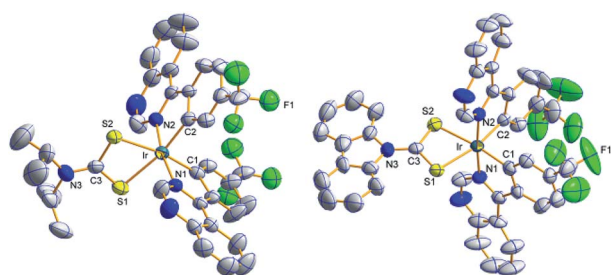


Fig. 2 ORTEP diagram of (4tfmpq)₂Ir(dipdte) (CCDC no. 1832357) and (4tfmpq)₂Ir(Czdtc) (CCDC no. 1832334) with the atom-numbering schemes. Hydrogen atoms are omitted for clarity. Ellipsoids are drawn at the 50% probability level.

membered or six-membered heterocycles such as those based on pic and acac derivatives.^{5,15,16} In (4tfmpq)₂Ir(Czdtc), the bond length of C3–S1 (1.710 Å) which is similar to that of C3–S2 (1.702 Å) also illustrates that −1 charge of dithiocarbamate is scattered over both sulfur atoms. The two coordinated S atoms of the ancillary ligand reside in the equatorial plane *trans* to the metalated C (4tfmpq) atoms. From the above analysis, this compact and tensile bonding mode is well understood.

Electrochemical properties and theoretical calculations

The highest occupied molecular orbital (HOMO) and lowest unoccupied molecular orbital (LUMO) energy levels of the dopants are important for the structure design of OLEDs. To determine the HOMO/LUMO energy levels, the electrochemical behaviors of the three complexes were measured by cyclic voltammetry in CH₂Cl₂ solution at room temperature, relative to an internal ferrocenium/ferrocene reference (Fc⁺/Fc) (Fig. S4,† and Table 1). As revealed previously by electrochemistry and theoretical calculations of cyclometalated Ir(III) complexes, it is generally considered that the reduction process occurs mainly on the heterocyclic portion of the cyclometalated C⁺N ligands, whereas oxidation processes largely involve the Ir–aryl center and the ancillary ligand.¹⁷ Based on the above analysis, these complexes have almost equal LUMO energy levels (−3.19 to −3.22 eV) due to the same quinazoline-based frame, whereas their HOMO levels systematically change with the variation of the ancillary ligands from −5.44 eV of (4tfmpq)₂Ir(dipdte) containing diisopropylamine to −5.50 and −5.61 eV of (4tfmpq)₂Ir(dpdtc) and (4tfmpq)₂Ir(Czdtc) complexes, respectively, by introducing the diphenylamine and carbazole groups into the ancillary ligands. These results suggest that the diphenylamine and carbazole groups in the ancillary ligands would stabilize the HOMO level and positively shift the oxidation potentials.

To obtain further electronic structure information, the HOMO/LUMO distributions of these Ir(III) molecules were also obtained with DFT calculations. As shown in Fig. S5,† the HOMO orbitals of the three Ir(III) complexes are mostly located on the 4tfmpq (43.55–47.90%) together with the d orbitals of the iridium atom (39.42–47.66%) with a small portion of the dithiocarbamate derivatives (8.66–12.67%). And a larger scale distribution of electron clouds on the iridium atom also indicates an efficient MLCT of the phosphorescent complexes, crucial to high PL quantum efficiency. With the decrease of



the electron-donating ability of the substituents from diisopropylamine and diphenylamine to carbazole, the electron clouds distributed over the dithiocarbamate derivatives decrease gradually from 12.67% and 8.79% to 8.66%, which verifies the gradual change of the HOMO level of the corresponding complexes (Table S3[†]). The LUMOs are mostly distributed over the π^* orbitals of the cyclometalated ligand (93.33–94.35%) and to a small extent on Ir d orbitals (3.45–4.13%) and ancillary ligands (2.20–2.34%). The calculated LUMOs remain almost unchanged, yet the HOMOs vary significantly, which correlate well with the electrochemical results (Table S3[†]).

Photophysical properties

Fig. 3 shows the ultraviolet-visible absorption and photoluminescence spectra of $(4\text{tfmpq})_2\text{Ir}(\text{dipdte})$, $(4\text{tfmpq})_2\text{Ir}(\text{dpdte})$ and $(4\text{tfmpq})_2\text{Ir}(\text{Czdtc})$ complexes at room temperature in CH_2Cl_2 , and their photophysical data are listed in Table 1. The intense absorption bands below 400 nm arise from the spin-allowed intraligand $\pi \rightarrow \pi^*$ transitions, and the relatively weak bands in the visible region (400–630 nm) can be assigned to the mixed $^1\text{MLCT}$ and $^3\text{MLCT}$ (metal-to-ligand charge-transfer) states, or the LLCT (ligand-to-ligand charge-transfer) transition through strong spin-orbit coupling of the iridium atom.¹⁸ Compared with the complex $(4\text{tfmpq})_2\text{Ir}(\text{dipdte})$ using the *N,N*-diisopropyl dithiocarbamate ancillary ligand ($\lambda_{\text{peak}} = 641$ nm, Fig. 3(b)), complexes $(4\text{tfmpq})_2\text{Ir}(\text{dpdte})$ containing *N,N*-diphenyl dithiocarbamate and $(4\text{tfmpq})_2\text{Ir}(\text{Czdtc})$ with *N*-carbazolyl dithiocarbamate exhibit 13 and 30 nm hypsochromic shifts, coinciding with the variation of calculated energy gaps.

The PL quantum yields of the three Ir(III) complexes were measured as 58.3%, 84.3% and 93.0%, respectively, which may be attributed to the gradually increased rigidity of the corresponding Ir(III) complexes by gradually increasing the rigidity of the electron-donating substituents in the dithiocarbamate ancillary ligand from diisopropylamine and diphenylamine to carbazole. When the rigidity of the ligands is enhanced gradually, the non-radiative transitions caused by the vibration and rotation of a large number of C–H and C–C bonds are much reduced, which is also a way of energy consumption for the triplet states of the Ir(III) complexes. In addition, the lifetimes of the three complexes are in the microsecond range (1.41 μs for $(4\text{tfmpq})_2\text{Ir}(\text{dipdte})$, 1.55 μs for $(4\text{tfmpq})_2\text{Ir}(\text{dpdte})$ and 1.50 μs

for $(4\text{tfmpq})_2\text{Ir}(\text{Czdtc})$, respectively) (Fig. S6[†]), which is indicative of the phosphorescence origin for the excited states in each case.

OLED performance

To illustrate their electroluminescence (EL) properties, typical OLEDs using these complexes as dopants were fabricated. Firstly, single-emissive-layer devices with ITO/HAT-CN (dipyr-azino[2,3-f:2',3'-h]quinoxaline-2,3,6,7,10,11-hexacarbonitrile, 6 nm)/HAT-CN (0.2 wt%):TAPC (di-[4-(*N,N*-ditolylamino) phenyl] cyclohexane, 50 nm)/(4tfmpq)₂Ir(dipdte) or (4tfmpq)₂Ir(dpdtc) or (4tfmpq)₂Ir(Czdtc) (3 wt%):2,6DCzPPy (2,6-bis(3-(carbazol-9-yl)phenyl)pyridine, 10 nm)/Tm3PyP26PyB (1,3,5-tris(6-(3-(pyridin-3-yl)phenyl)pyridin-2-yl) benzene, 60 nm)/LiF (1 nm)/Al (100 nm) are named as S1–S3, respectively. As shown in Scheme 2, the materials of HAT-CN and LiF served as hole- and electron-injecting interface modified materials, respectively. The bipolar material 2,6 DCzPPy was chosen as the host material. The TAPC material with a high HOMO level (−5.5 eV) acts as the hole-transport medium, and the layer of TAPC is doped with 0.2 wt% HAT-CN to further enhance the hole-transporting ability, while Tm3PyP26PyB with a low LUMO level (−2.7 eV) is used as the electron-accepting and electron-transport material. The HOMO and LUMO levels of Ir(III) complexes were all embedded between those of 2,6 DCzPPy (HOMO = 6.10 eV and LUMO = 2.60 eV). Thus, efficient energy transfer from the host to Ir(III) complexes occurred in the emitting layer (EML). Furthermore, holes and electrons can be injected to the HOMO and LUMO levels of the host material or trapped by the dopants directly owing to the difference of the HOMO/LUMO levels between the Ir(III) complexes and the host materials. And in the bipolar host 2,6 DCzPPy the excitons can be easily formed and distributed more balanced, which is helpful to reduce the turn-on voltage. All the devices were optimized and they display the best EL performances all at 3 wt% doping concentration, and the details of optimizing the doping concentration of Ir(III) emitters are included in the ESI.[†] The electroluminescence (EL) spectra, current density (*J*)–voltage (*V*)–luminance (*L*), current efficiency (η_c) and power efficiency (η_p) versus luminance, and EQE as a function of luminance characteristics of the devices

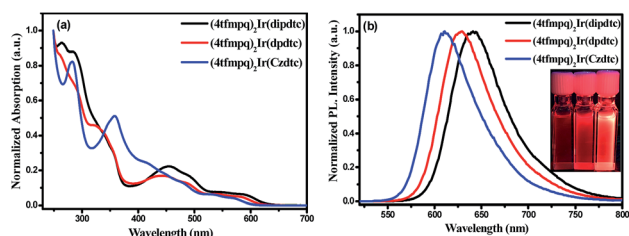
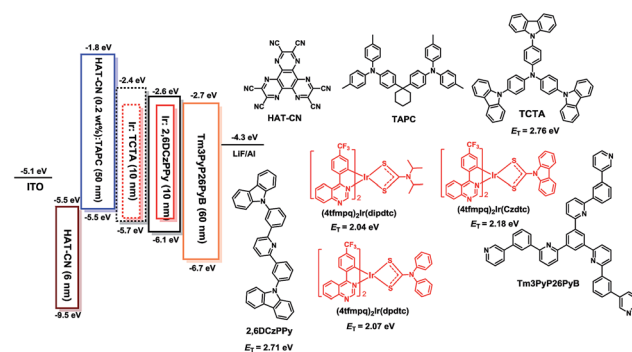


Fig. 3 The UV-vis absorption (a) and emission (b) spectra of the three complexes $(4\text{tfmpq})_2\text{Ir}(\text{dipdte})$, $(4\text{tfmpq})_2\text{Ir}(\text{dpdte})$ and $(4\text{tfmpq})_2\text{Ir}(\text{Czdtc})$ in degassed dichloromethane (5×10^{-5} M) at room temperature.



Scheme 2 Energy level diagram of the HOMO and LUMO levels of materials investigated and their chemical molecular structures.



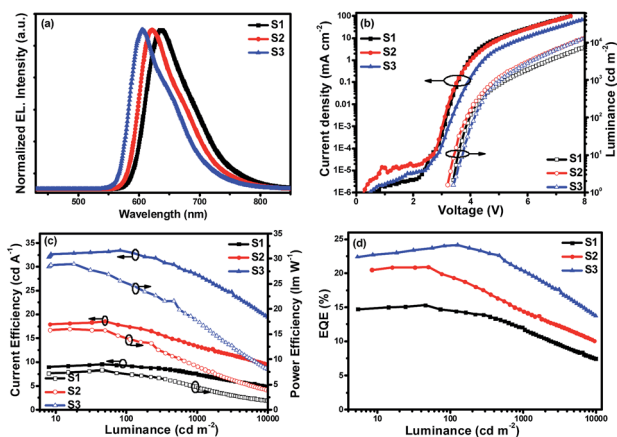


Fig. 4 Characteristics of single-emitting-layer devices: (a) EL spectra, (b) J - V - L curves, (c) η_c - L - η_p curves and (d) EQE- L curves.

are displayed in Fig. 4, and the parameters of these devices are summarized in Table 2.

Fig. 4(a) shows the normalized EL spectra of the three devices measured at the current of 1 mA with peaks at about 636, 621 and 605 nm for devices S1–S3, which are very close to the PL spectra of the three Ir(III) complexes, indicating that the EL emission of the devices originates from the triplet excited states of the phosphors in which all the Commission Internationale de l'Eclairage (CIE) color coordinates fall in the red region. Due to the same cyclometalated ligand and similar molecular structures of the ancillary ligands, the device performances of these materials mainly depend on their PL efficiencies and bipolar properties. With the changes of the substituents from diisopropylamine and diphenylamine to carbazole, the resulting complexes from (4tfmpq)₂Ir(dipdte) and (4tfmpq)₂Ir(dpdtc) to (4tfmpq)₂Ir(Czdtc) exhibit more notable bipolar properties, better carrier balances and enhanced photoluminescence quantum efficiency, which would lead to the gradually enhanced OLED performances.

Respectively, the device S1 (Φ_p of (4tfmpq)₂Ir(dipdte) is 58.3%) shows the lowest EL performances with a maximum luminance ($L_{\max}$) above 26 000 cd m^{-2} , a peak current efficiency ($\eta_{c,\max}$) of 9.81 cd A^{-1} , a peak power efficiency ($\eta_{p,\max}$) of 7.86 lm W^{-1} and a maximum external quantum efficiency ($\text{EQE}_{\max}$) of 15.30%. Relatively, the device S2 (Φ_p of (4tfmpq)₂Ir(dpdtc) is 84.3%) displays better performances with a $L_{\max}$, $\eta_{c,\max}$ and

$\text{EQE}_{\max}$ of 39 870 cd m^{-2} , 18.56 cd A^{-1} and 20.91%, respectively. Due to the highest Φ_p of (4tfmpq)₂Ir(Czdtc) (93.0%), the device S3 exhibits the best characteristics with a $L_{\max}$, $\eta_{c,\max}$ and $\text{EQE}_{\max}$ of 56 880 cd m^{-2} , 32.92 cd A^{-1} and 24.18%, respectively.

In order to further improve the EL performances of these complexes, another hole transport material TCTA (4,4',4''-tris(carbazol-9-yl)triphenylamine) was introduced as the second host for these emitters and also as a "hole ladder" layer due to its suitable HOMO level (−5.70 eV) between TAPC and 2,6DCzPPy (Scheme 2). The OLEDs with double emissive layers with the configuration of ITO/HAT-CN (6 nm)/HAT-CN (0.2 wt%):TAPC (50 nm)/Ir(III) complexes (x wt%):TCTA (10 nm)/Ir(III) complexes (x wt%):2,6DCzPPy (10 nm)/Tm3PyP26PyB (60 nm)/LiF (1 nm)/Al (100 nm) using the dopants of (4tfmpq)₂Ir(dipdte), (4tfmpq)₂Ir(dpdtc) and (4tfmpq)₂Ir(Czdtc) are named as D1–D3, respectively (Scheme 2). The corresponding EL characteristics of the devices are shown in Fig. 5, and the detailed results are also summarized in Table 2. The stepwise changed HOMO energy levels of TAPC (−5.5 eV), TCTA (−5.7 eV) and 2,6DCzPPy (−6.1 eV) are beneficial for the hole injection and transport. Similarly, they are also beneficial for the injection and transport of electrons owing to the gradually changed LUMO energy levels of Tm3PyP26PyB (−2.7 eV), 2,6DCzPPy (−2.6 eV) and TCTA (−2.4 eV). Thus, holes and electrons will be distributed in more balanced emissive layers and the exciton recombination zone is expected to be broadened. Furthermore,

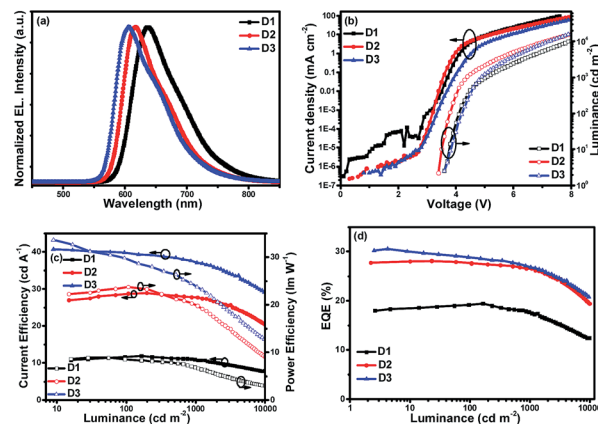
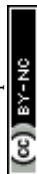


Fig. 5 Characteristics of double emitting-layer devices: (a) EL spectra, (b) J - V - L , (c) η_c - L - η_p and (d) EQE- L curves.

Table 2 The key EL data of devices S1–S3 and D1–D3

Device	$V_{\text{turn-on}}^a$ (V)	$L_{\max}$ (cd m^{-2})	$\eta_{c,\max}$ (cd A^{-1})	$\eta_{\text{ext},\max}$ (%)	$\eta_{p,\max}$ (lm W^{-1})	η_c^b (cd A^{-1})	η_{ext}^b (%)	CIE ^c (x, y)
S1	3.4	26 490	9.51	15.30	7.86	7.38	11.69	(0.68, 0.31)
S2	3.2	39 870	18.56	20.91	16.00	13.27	14.36	(0.66, 0.33)
S3	3.4	56 880	32.92	24.18	28.94	28.14	20.33	(0.63, 0.36)
D1	3.6	33 540	11.89	19.46	8.87	10.76	17.47	(0.68, 0.31)
D2	3.4	46 920	28.95	28.10	20.97	27.74	26.53	(0.65, 0.33)
D3	3.6	60 950	40.68	30.54	33.63	37.10	26.79	(0.63, 0.36)

^a Applied voltage recorded at a luminance of 1 cd m^{-2} . ^b Recorded at 1000 cd m^{-2} . ^c Measured at 1000 cd m^{-2} .



the Tm3PyP26PyB HOMO energy level is 0.6 eV lower than that of 2,6DCzPPy and the TAPC LUMO energy level is 0.6 eV higher than that of TCTA, which will result in excitons (hole–electron pairs) being well conformed within emissive layers and the triplet exciton quenching being avoided effectively. From Fig. 5 and Table 2, it can be observed that all the devices with double emissive layers show better performances than S1–S3.

Respectively, D1 based on (4tfmpq)₂Ir(dipdte) at a doping concentration of 3 wt% displayed good performances with a $L_{\max}$ above 33 000 cd m^{−2}, $\eta_{c,\max}$ over 11.89 cd A^{−1}, $\eta_{p,\max}$ over 8.87 lm W^{−1} and EQE_{max} of 19.46%. The device D2 based on (4tfmpq)₂Ir(dpdtc) at a doping concentration of 1 wt% displayed higher characteristics with a $L_{\max}$ of 46 920 cd m^{−2}, an $\eta_{c,\max}$ of 28.95 cd A^{−1}, an $\eta_{p,\max}$ of 20.97 lm W^{−1} and an EQE_{max} of 28.10% with CIE coordinates of (0.65, 0.33), which are quite close to the standard red demanded by the National Television System Committee (NTSC). Device D3 based on (4tfmpq)₂Ir(Czdtc) with the *N*-carbazolyl dithiocarbamate ancillary ligand at 3 wt% doped concentration showed the best EL performances with a high brightness of 60 950 cd m^{−2} and the $\eta_{c,\max}$, EQE_{max} and $\eta_{p,\max}$ are up to 40.68 cd A^{−1}, 30.54% and 33.63 lm W^{−1}, respectively. The CIE coordinates of (0.63, 0.36) also fall in the red region. The EQEs of 30.54% and the maximum brightness of 60 950 cd m^{−2} are among the highest results ever reported for red Ir(III) complexes.⁷ Furthermore, the device efficiency roll-off is not serious. As for device D3, the η_c and EQE can still be obtained as 37.10 cd A^{−1} and 26.79%, respectively, when the luminance reaches 1000 cd m^{−2}.

Summarized from the investigated results, the superior device properties may result from the following facts. Firstly, dithiocarbamate derivatives with different electron-donating moieties and the nitrogen frame of 4-(4-(trifluoromethyl)phenyl)quinazoline were applied as the ancillary ligands and cyclometalated ligand for stable Ir(III) complexes with bipolar properties, which may contribute to a wider recombination area and a more balanced distribution for the excitons. Secondly, the coordination capability of the sulfur atom with the iridium atom is strong, and the stable dithiolate compounds can reduce the work function of the emitter and the threshold electric field.⁸ The different dithiocarbamate derivatives can lower the LUMO energy levels of the complexes, and the electron transport properties can benefit from lowering the LUMO levels of the dopants. Finally, double light-emitting layers were adopted, which showed improved and balanced charge-injection/transporting properties compared to single layer-emitting devices.¹⁹

Conclusions

In conclusion, three cyclometalated Ir(III) complexes containing the unique four-membered Ir–S–C–S backbone ring were rapidly prepared at room temperature with good yields above 70% and high PLQYs up to 93.0% in CH₂Cl₂. The Gibbs free energy calculation results confirm that all reactions are exothermic and thermodynamically favorable processes. Through changing the ancillary ligands with different electron-donating groups, the rigidity and bipolar properties of these

Ir(III) complexes can be effectively tuned, resulting in regularly changed emission wavelength and OLED performances. The double-emissive-layer devices using the complex with *N*-carbazolyl dithiocarbamate as the ancillary ligand displays exceedingly good performances with a maximum brightness over 60 000 cd m^{−2} and an EQE_{max} of 30.54%. Even at the brightness of 1000 cd m^{−2}, the EQE can still be obtained as 26.79%, respectively. This study provides a useful route to synthesize Ir(III) complexes with the four-membered ring Ir–S–C–S structure rapidly, which is key to the industrial production of phosphorescent materials for high-performance OLEDs.

Experimental section

Synthesis of the salts of dithiocarbamate derivatives

Lithium *N,N*-diisopropyl dithiocarbamate (Li-dipdte). Diisopropyl amine (40 mmol) was added to a stirring slurry of LiOH (2.0 g, 48 mmol) in water (3 mL) and CS₂ (3.6 g, 48 mmol) in benzene (10 mL). An exothermic reaction ensued, followed by formation of yellow to orange precipitates. After 1 h, petroleum ether was added and the reaction mixture was filtered. The solids were washed well with petroleum ether and dried *in vacuo* to give a white to pale yellow solid, which was used in the next procedure without further purification.

Lithium *N,N*-diphenyl dithiocarbamate (Li-dpdte). A solution of diphenylamine (500 mg, 2.96 mmol) was dissolved in degassed THF (6 mL), then cooled to 0 °C and treated with *n*-BuLi (30 mL, 5.92 mmol). After stirring for 30 min at 0 °C, CS₂ (2 mL, 29.6 mmol) was added. The ice bath was removed and stirred for 12 h at room temperature. The crude mixture was recrystallized from THF in Et₂O, to afford lithium *N,N*-diphenyl dithiocarbamate as an off-white solid, which was used in the next procedure without further purification.

Sodium *N*-carbazolyl dithiocarbamate (Na-Czdtc). Carbazole powder (3.34 g, 0.02 mol) dissolved in 30 mL of THF was added dropwise into 0.72 g of NaH (0.03 mol) suspended in 5 mL of THF in an ice-water bath under a N₂ atmosphere. Afterwards, the mixture was stirred for another 15 min at room temperature. Then, 1.33 mL of CS₂ (0.022 mol) was added dropwise into the system. The system was subsequently immersed in an oil bath at 40 °C and reacted for 3 h. During the reaction time, the color of the system gradually turned dull-red. The system of sodium thionocarboxylate in THF was quenched by adding a small amount of water in an ice-water bath. After the solvent was removed from the reaction mixture, 100 mL of water was added into the residue, and the aqueous phase was filtered. The filtrate was washed with diethyl ether three times to obtain pure sodium thionocarboxylate aqueous solution, which was used in the next procedure without further purification.

Synthesis of iridium(III) complexes

IrCl₃ (0.64 g, 2.14 mmol) and 2.4 equivalents of cyclometalated ligand (5.14 mmol) were added in a 2-ethoxyethanol and water mixture. Then, the solution was heated for 16 h at 110 °C. After the addition of water, the precipitated red powder of [(C[^]N)₂Ir(μ-Cl)]₂ chloride-bridged dimer was filtered and



reacted with dithiocarbamate derivatives without further purification for 5 minutes at room temperature in EtOCH₂CH₂OH. Then the solution was concentrated and the resulting residue was purified by silica gel column chromatography (CH₂Cl₂/petroleum ether 1 : 2 (v/v)) and vacuum sublimation gave red crystals.

(4tfmpq)₂Ir(dipdte). Yield: 77%. ¹H NMR (400 MHz, CDCl₃) δ 10.43 (s, 2H), 8.85 (d, *J* = 8.6 Hz, 2H), 8.43 (d, *J* = 8.4 Hz, 2H), 8.29 (d, *J* = 8.0 Hz, 2H), 8.04–7.96 (m, 2H), 7.88–7.83 (m, 2H), 7.18 (dd, *J* = 8.4, 1.3 Hz, 2H), 6.74 (s, 2H), 1.37 (d, *J* = 6.8 Hz, 8H), 1.28–1.22 (m, 4H). ¹⁹F NMR (377 MHz, CDCl₃) δ –63.19 (s). HR-MS, *m/z*: calcd for C₃₇H₃₀N₅F₆S₂Ir, 915.1476 [M]; found 916.1557 [M + H]⁺. Anal. calcd for C₃₇H₃₀N₅F₆S₂Ir: C, 48.57; H, 3.30; N, 7.65. Found: C, 48.67; H, 3.32; N, 7.75%.

(4tfmpq)₂Ir(dpdtc). Yield: 75%. ¹H NMR (400 MHz, CDCl₃) δ 10.31 (d, *J* = 2.4 Hz, 2H), 8.73 (d, *J* = 8.5 Hz, 2H), 8.30 (d, *J* = 8.4 Hz, 2H), 8.24 (d, *J* = 8.2 Hz, 2H), 7.93 (t, *J* = 7.7 Hz, 2H), 7.76 (dd, *J* = 8.4, 7.2 Hz, 2H), 7.42–7.32 (m, 6H), 7.32–7.24 (m, 2H), 7.23–7.17 (m, 2H), 7.07 (dd, *J* = 8.4, 1.2 Hz, 2H), 6.66 (s, 2H). ¹⁹F NMR (377 MHz, CDCl₃) δ –63.25 (s). HR-MS, *m/z*: calcd for C₄₃H₂₆N₅F₆S₂Ir, 983.1163 [M]; found 984.1247 [M + H]⁺. Anal. calcd for C₄₃H₂₆N₅F₆S₂Ir: C, 52.54; H, 2.67; N, 7.12. Found: C, 52.74; H, 2.47; N, 7.32%.

(4tfmpq)₂Ir(Czdte). Yield: 70%. ¹H NMR (400 MHz, CDCl₃) δ 10.33 (s, 2H), 9.35–9.18 (m, 2H), 8.84 (d, *J* = 8.5 Hz, 2H), 8.47 (d, *J* = 8.4 Hz, 2H), 8.24 (d, *J* = 8.2 Hz, 2H), 8.03–7.95 (m, 2H), 7.95–7.89 (m, 2H), 7.85 (dd, *J* = 11.5, 4.1 Hz, 2H), 7.40–7.30 (m, 4H), 7.30–7.22 (m, 2H), 6.82 (s, 2H). ¹⁹F NMR (377 MHz, CDCl₃) δ –63.18 (s). HR-MS, *m/z*: calcd for C₄₃H₂₄N₅F₆S₂Ir, 981.1007 [M]; found 982.1090 [M + H]⁺. Anal. calcd for C₄₃H₂₄N₅F₆S₂Ir: C, 52.65; H, 2.47; N, 7.14. Found: C, 52.47; H, 2.61; N, 6.99%.

Conflicts of interest

There are no conflicts to declare.

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