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Bay-substituted octaazaperopyrenedioxides as solid-state emitters for strong light-matter coupling†

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Molecules based on the octaazaperopyrenedioxide (OAPPDO) motif represent a new class of organic emitters with the ability to form amorphous films when processed from solution while retaining their emissive properties. Here, we report the synthesis of bay-functionalized OAPPDO derivatives with thioether-substituents, which are structurally characterized by their twisted π -core. The combination of this structural motif with suitable substituents at the chromophore facilitates the deposition of neat and smooth films from solution, in which the molecules remain electronically decoupled, thus drastically reducing non-radiative decay pathways. An analysis of absorption and emission spectra of solutions and spincoated thin films is provided along with details of their structural morphology and thermodynamics. The amorphous thin films display spectroscopic characteristics similar to those of the isolated molecules in solution and exhibit photoluminescence quantum yields up to 17%. The high oscillator strength of smooth films of these OAPPDO derivatives enables their integration in Fabry–Pérot microcavities with high quality factors and the observation of strong light-matter coupling with clear exciton–polariton modes and large coupling strength.

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1. Introduction

Strong interactions between dye molecules in crystalline or amorphous thin films are beneficial for charge transport but pose a significant challenge for the development of highly emissive thin films for applications such as organic light-emitting diodes (OLEDs),^{1,2} luminescent solar concentrators,^{3,4} or laser gain media.^{5,6} Various nonradiative decay pathways reduce the photoluminescence quantum yield (PLQY) of densely packed and electronically coupled molecules, such as concentration quenching, singlet fission, or the formation of non-emissive aggregates.^{7,8} Methods to electronically decouple luminescent molecules, and thus maintain their emissive

properties in the solid state, include dilution within a polymer or small molecule matrix,^{9,10} integration in nanoporous frameworks,^{11,12} or the protection of the chromophore core with bulky substituents.^{13–15} Unfortunately, these strategies often lead to a lower density of emitters per volume or poor film formation.

One large area of application for molecular solid-state emitters is exciton–polaritonics, *i.e.*, strong light-matter coupling in optical microcavities. Excitonic materials enclosed in a resonant Fabry–Pérot (or similar) cavity can reach the so-called strong-coupling regime when the exchange of energy between the cavity photons and excitons is faster than the decay of either component.^{16–18} New mixed light-matter states are formed (lower and upper polariton) that enable the precise control of light absorption and emission in these microcavities while keeping the emitters the same. Optoelectronic devices, such as OLEDs and photodiodes can be tuned in the strong coupling regime through adjusting the coupling strength (*via* the concentration of emitters) and detuning of the cavity (*via* cavity thickness).^{19–22} An example of such tuning is the angle-independent emission wavelength of ultrastrongly coupled OLEDs.^{23,24} Furthermore, due to the high exciton binding energy of their excitons, organic emitters can facilitate exciton–polariton

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condensation and polariton lasing under suitable conditions even at room temperature.^{25–30} To reach the strong or ultrastrong coupling regime, the density and oscillator strength of the molecules in the cavity must be high enough. Furthermore, the lifetime of the excitons must be sufficiently long and the quality factor of the produced cavity should be as high as possible. The latter is often limited by the roughness of the emitter film. To reach polariton lasing the photoluminescence quantum yield of the emitter also has to be high. Although there have been several examples of polariton lasing, the field is still limited by the availability of organic emitters with high oscillator strength, which retain their absorption and emission properties in solution-processed neat and smooth thin films.

To address this limitation, we apply an alternative strategy for the formation of smooth, amorphous and aggregate-free thin films with a high density of emissive small molecules. To suppress aggregation of π -conjugated molecules we make use of non-planar π -systems instead of bulky sidegroups. Based on initial studies with *peri*-substituted tetraazaperylenes,³¹ we have recently begun to investigate the suitability of octaazaperopyrenedioxides (OAPPDOs)^{32,33} (see Fig. 1) as emitters at high concentrations in solution as well as in condensed thin films and their application as fluorophores in two-photon micro-printed 3D emissive structures.³⁴

Formally, the OAPPDO compounds may be viewed as tetraazaperylenes that are decorated with *N*-substituted urea units in both opposite *peri*-positions. In particular, we found that the facile modification of these fluorophores in the *bay*-positions enables efficient tuning of their luminescent properties. Moreover, the *bay*-substituents of the tetraazaperylene core induce a twist of the polycyclic aromatic core which may reduce the degree of π - π interaction. This twist therefore potentially favours retention of the photophysical properties of isolated chromophores in condensed amorphous or crystalline thin films.

Here, we investigate the impact of introducing sulfur-based groups in substituents in the *bay* positions of OAPPDOs (see Fig. 1) on their photophysical properties. This structural

modification barely alters their absorption and photoluminescence properties, both in solution and in solution-processed thin films. Moreover, we found that these compounds have the tendency to form smooth and amorphous neat films with good PLQY. We further demonstrate the ability of thin films of OAPPDO derivatives to enter the strong coupling regime in suitable Fabry–Pérot microcavities with nearly angle-independent emission, hence expanding the library of potential organic molecules for polaritonic devices.

2. Results and discussion

2.1. Synthesis of OAPPDO thioethers and sulfones and their crystal structures

The previously described³³ *bay*-chlorinated OAPPDO **1** enabled the facile preparation of thioether-functionalized derivatives **2a–f** via nucleophilic substitution in high yields (see Scheme 1). The corresponding thiolates were prepared *in situ* or in the case of the perfluoro derivative **2e** the copper thiolate was reacted with **1**. Both are common approaches for the sulfur-functionalization of perylene dyes.³⁵ Thioethers **2a**, **b** and **f** were oxidized to the corresponding sulfones **3a–c** with excess *meta*-chloroperbenzoic acid (*m*-CPBA) in good yields (for a detailed description of the synthesis, see ESI†). To investigate the packing of the thioether-functionalized OAPPDOs in the solid state, single crystals of **2a–c**, **2e** and **3a** were grown by slow evaporation of concentrated solutions of each compound and crystallographic structure analysis was performed (see Fig. 2 and Tables S1–S3, ESI†). The tetraazaperylene core of all OAPPDO fluorophores was found to be highly twisted (twist angles for **2a**: 28.99(4)° and 30.43(5)° for two independent molecules in the cell, **2b**: 30.83(4)°, **2c**: 29.92(5)°, **2e**: 30.32(8)°, **3a**: 31.16(4)°) due to the large van der Waals radii of the sulfur atoms of the thioether or sulfone functionalities and the resulting steric repulsion in the *bay* area. Consistent with the general structural characteristics of rylene, the individual

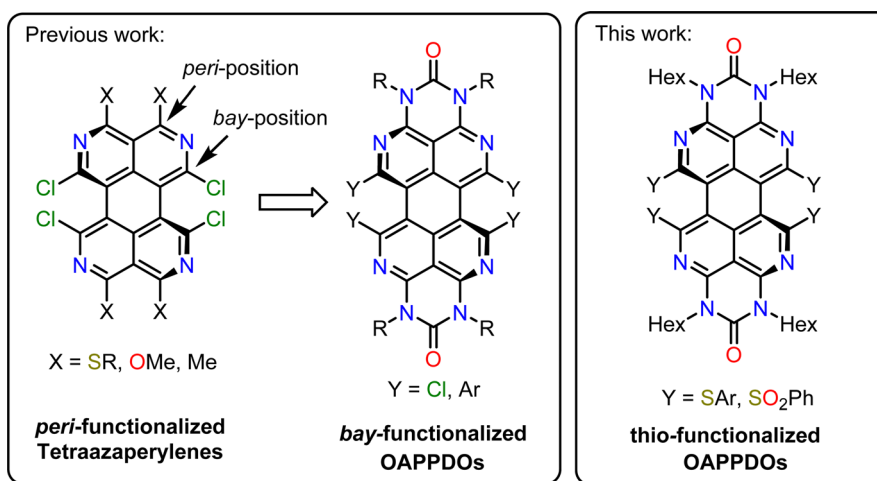
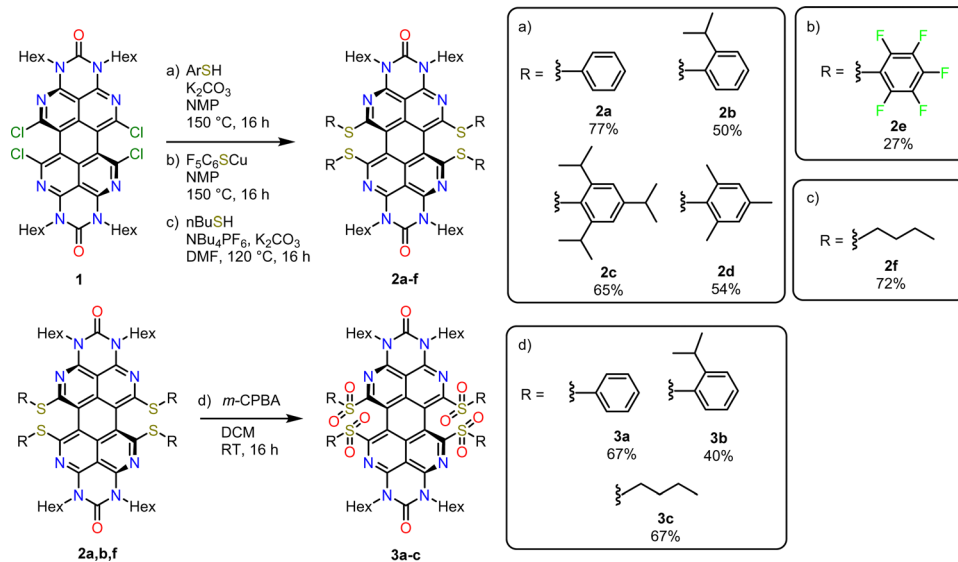


Fig. 1 Previously investigated fluorophores based on a tetraazaperylene core and their functionalization with sulfur-based groups at *bay*-positions (this work).





Scheme 1 Synthesis of thioether- and sulfone-functionalized OAPPDOs. Reaction conditions: (a) ArSH, K_2CO_3 , NMP, 150 °C, 16 h; (b) F_5C_6SCu , NMP, 150 °C, 16 h; (c) *n*-BuSH, NBu_4PF_6 , K_2CO_3 , DMF, 120 °C, 16 h; (d) *m*-CPBA, DCM, room temperature, 16 h.

naphthyridine moieties within the tetraazaperylene core were found to be “fused” through long C–C bonds of around 1.46 Å. The principal structural data are in the typical range of previously reported *bay*-functionalized tetraazaperylene dyes.^{31–33}

In all compounds, the steric demand of the *bay*- and *N*-hexyl-substituents at the periphery of the fluorophore suppresses significant π – π -stacking of the perylene cores. Additionally, the variation of the *bay*-substituents leads to distinctly different patterns of molecular packing. In compounds **2a**, **2b** and **2e** the molecules are stacked to give a herringbone (**2a**) and sandwich herringbone (**2b** and **2e**) packing motif with distances of approximately 10.7 Å (**2a**), 3.4 Å (**2b**) and 3.3 Å (**2e**) between the polycyclic aromatic cores. The latter two suggest weak π – π -interactions at the *peri* region of the perylene units while the central pericyclic cores appear to be ‘decoupled’. In contrast to that, compounds **2c** and **3a** exhibit both an offset stacked packing pattern with inter-layer distances of approximately 4.5 Å and 7.8 Å, respectively. Such inhibition of π – π -interactions by twisting the π -core system has been studied for numerous perylenediimide (PDI) derivatives^{36–40} and can lead to dramatic changes in the photophysical properties of thin PDI films. For example, successive brominations in the *bay* position of a PDI derivative and concomitant opening of the twist angle of the perylenic core drastically increased the yield of triplet formation.⁴¹ Moreover, a recent study on singlet fission in amorphous PDI thin films has shown that the twisted polycyclic aromatic cores suppressed the formation of competing excimer states.⁴² The following sections will therefore first focus on the optical properties of these new OAPPDO dyes in solution and, subsequently, in neat thin films.

2.2. Absorption and emission properties in solution

All OAPPDO derivatives are highly soluble in polar organic solvents, while the thioether-functionalized derivatives also

display good solubility in nonpolar solvents such as *n*-hexane. The absorption spectra of the compounds exhibit intense bands in the visible spectral region with large extinction coefficients (ϵ) and a characteristic vibronic progression (see Fig. 3 and Table 1 and Fig. S1, S2 (ESI[†]) for spectra of the other derivatives), indicating coupling between the electronic transitions and the vibrational modes of the tetraazaperylene cores.

The absorption maxima of the thioether-functionalized OAPPDOs **2a–f** range from 521 to 533 nm with molar absorption coefficients of around $22\,400\ M^{-1}\ cm^{-1}$. Although all derivatives are non-planar, the vibronic structure for the $S_0 \rightarrow S_1$ (ground state to first excited state) transition remained similar for all OAPPDOs, indicating the pronounced rigidity of the tetraazaperylene core, consistent with previous reports for PDI derivatives.⁴³ Time-dependent density-functional theory (TD-DFT) calculations (basis set: B3LYP-GD3(BJ)/Def2-TZVPP)^{44–49} of **2a**, **2e**, **2f**, **3a** and **3c** confirmed that the maxima around 500 nm correspond to the first singlet transition ($S_0 \rightarrow S_1$) (see Fig. S3–S7 and Tables S4–S8, ESI[†]). Additionally an intense band between 400 nm and 450 nm is observed and can be attributed to a $S_0 \rightarrow S_2$ transition, which is enhanced due to the core twist of the OAPPDOs.³⁷

All the OAPPDO derivatives are fluorescent and their photoluminescence (PL) spectra are approximately the mirror image of their respective $S_0 \rightarrow S_1$ transition (see Fig. 3 and Fig. S1, S2 for PL spectra of the other derivatives, ESI[†]). Notably, this type of mirror image emission pattern was found to be even more pronounced for the sulfone-derivatized compounds **3a–c**, indicating a highly rigid tetraazaperylene core. This increased core rigidity for the sulfone-functionalized OAPPDOs is also reflected in the emission Stokes shifts (see Table 1), which are between $1400\ cm^{-1}$ and $1700\ cm^{-1}$ for *bay*-sulfido derivatives **2a–f** but only 736 – $891\ cm^{-1}$ for the sulfone-derivatives **3a–c**. The observed photophysics reflect the key structural modification



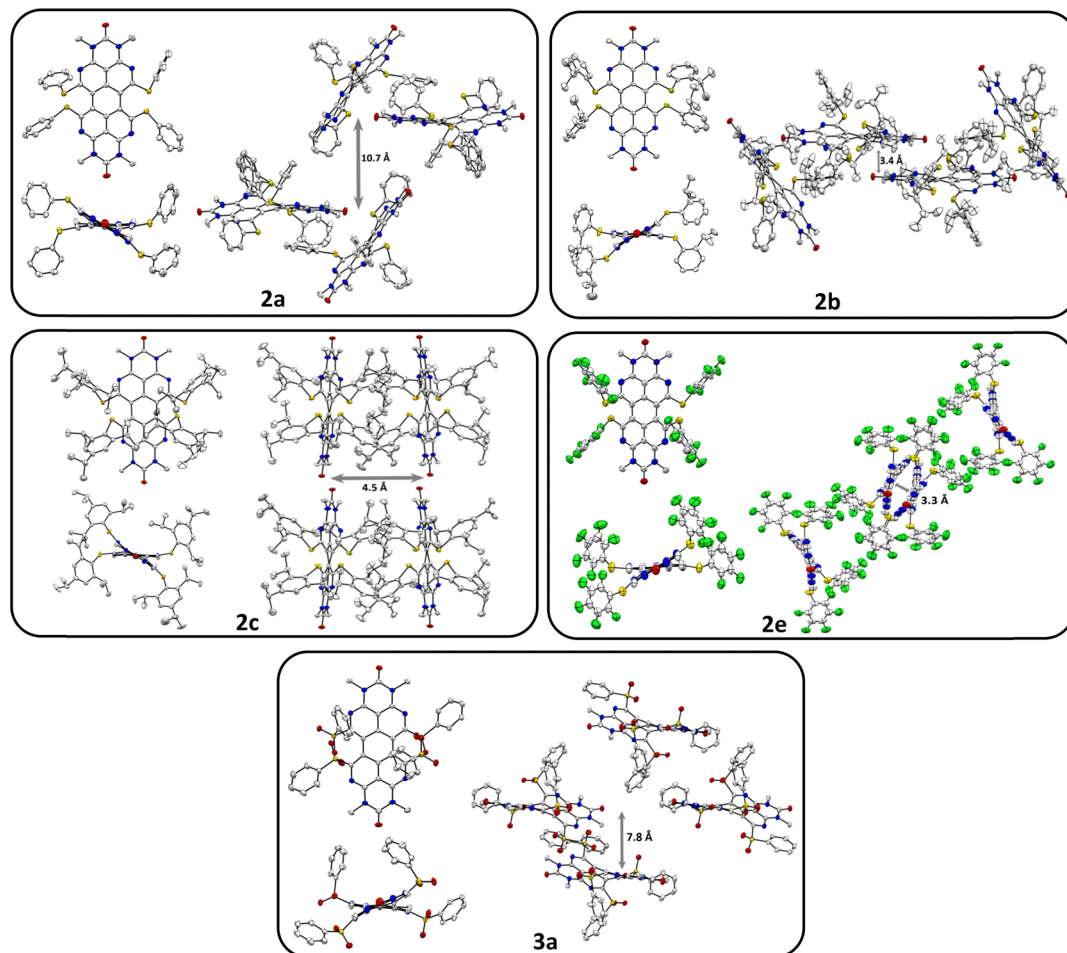


Fig. 2 Solid state structures and molecular packing motifs of thioether-functionalized OAPPDOs **2a** (view along [111]), **2b** (view along *a* axis), **2c** (view along *c* axis), **2e** (view along *a* axis), and sulfone-functionalized OAPPDO **3a** (view along *a* axis). *N*-hexyl substituents, hydrogen atoms and disorder are omitted for clarity. Thermal displacement ellipsoids at 50% probability level.

upon oxidation of the sulfide-functions in the *bay*-positions, namely the transformation of the twofold substituted sulfur atoms in the thioethers to the fourfold coordinated S-atoms in the sulfones. This significant increase in steric bulk in the immediate proximity of the *bay*-area rigidifies the chromophore with regard to the (activated) libration around the long molecular axis. Concomitantly, the increased oxidation state of the sulfur leads to a slight contraction and thus stabilization of the C–S bonds. The latter is a well-established pattern in organo-element chemistry.

PLQYs of up to 84% in solution were found for the compounds listed in Table 1. Notably, the oxidation of the *bay*-thioether units to the corresponding sulfones did not change the absorption behaviour significantly and only led to a small bathochromic shift of the maxima of less than 10 nm. However, the PLQY increased by about 10% upon oxidation. The fluorescence lifetimes (τ) of the OAPPDOs in dichloromethane (DCM) solution are listed in Table 1. All compounds showed a simple mono-exponential decay with lifetimes values between 5.7 ns and 8.0 ns, as typically expected for good fluorescent dyes. The corresponding rate constants for radiative (k_r) and non-radiative (k_{nr}) decay were also

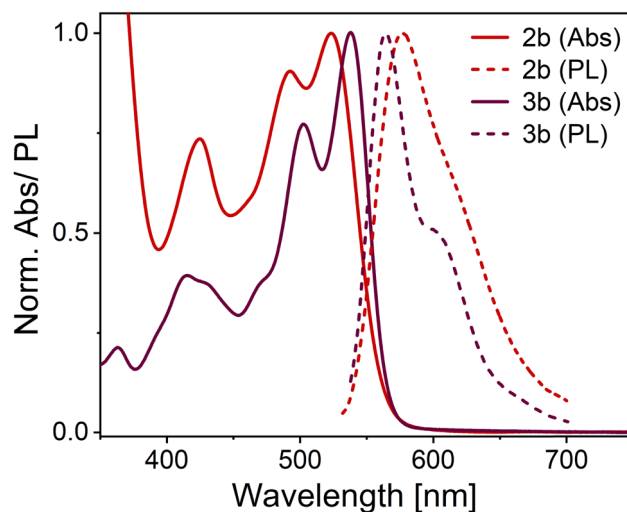


Fig. 3 Normalized absorption (Abs) and emission (PL) spectra of compounds **2b** and **3b** in dichloromethane (concentration ca. 10^{-5} M).

estimated from the measured PLQY and fluorescence lifetime. However, they were all fairly similar, except for slightly higher k_r



Table 1 Photophysical data of OAPPDOs **2a–f** and **3a–c** (in dichloromethane, concentration ca. 10^{-5} M). Maximum absorption and emission wavelengths ($\lambda_{\text{abs,max}}$, $\lambda_{\text{em,max}}$), extinction coefficient (ϵ), PL quantum yield (PLQY), Stokes shift and PL lifetime (τ) as measured by TCSPC (see ESI), the estimated rate constants for radiative (k_r) and non-radiative (k_{nr}) decay

	$\lambda_{\text{abs,max}}$ [nm] ($\log_{10} \epsilon$)	$\lambda_{\text{em,max}}$ [nm] (PLQY) ^a (%)	Stokes shift [cm^{-1}]	τ [ns]	k_r [10^7 s^{-1}]	k_{nr} [10^7 s^{-1}]
2a	525 (4.33)	577 (68)	1716	7.4	9.2	4.3
2b	532 (4.37)	576 (67)	1408	6.9	9.7	4.8
2c	533 (4.47)	575 (73)	1370	6.4	11.4	4.2
2d	527 (4.52)	569 (55)	1401	5.9	9.3	7.6
2e	521 (4.36)	563 (77)	1431	6.1	12.6	3.8
2f	530 (4.31)	573 (70)	1416	8.0	8.8	3.7
3a	536 (4.42)	558 (82)	736	6.1	13.4	2.9
3b	537 (4.57)	553 (84)	891	6.6	12.7	2.4
3c	528 (4.42)	551 (81)	791	5.7	14.2	3.3

^a Excitation wavelength: $\lambda_{\text{abs,max}} - 5$ nm. PLQY measured with an Ulbricht sphere (absorbance < 0.1).

and lower k_{nr} for the sulfones compared to the thioethers, which is in agreement with their higher PLQYs.

2.3. Thin films of OAPPDO derivatives

Neat thin films of compounds **2b–e** and **3a** were prepared by spin-coating from *o*-xylene or dichloromethane solution (for parameters see Table S9, ESI[†]). The compounds and solvents were mainly chosen to maximize the concentration of dissolved molecules while also ensuring smooth film formation during spincoating. Dichloromethane was required for the sulfone compounds due to their higher polarity. The thicknesses of the spin-coated neat films ranged from 120 nm to 200 nm. All films were annealed at 80 °C for two hours to remove residual solvents and to avoid thermal cracking during the atomic layer deposition of aluminium oxide (AlO_x) for encapsulation. All five OAPPDO compounds studied in this context (**2b–e**, **3a**) formed homogeneous and smooth thin films that displayed no visible signs of crystallization or birefringence in cross-polarized optical micrographs (see Fig. S8–S12, ESI[†]). All thin films were found to be amorphous at room temperature. Atomic force microscopy (AFM) images of the films of compounds **2d** and **3a** (see Fig. 4) also showed no signs of crystallization on the nanometer scale and were extremely smooth. The same was observed for the other OAPPDO derivatives (see Fig. S13, ESI[†]). The root mean square (RMS) roughness determined from AFM images was less than 0.5 nm (see Table 2), highlighting the amorphous nature and the excellent smoothness of these films, which is necessary for the fabrication of high quality optical cavities.

To further confirm the amorphous nature of the compounds in the solid state, their thermal behaviour was studied. Thermogravimetric analysis (TGA) and differential scanning calorimetry (DSC) of all compounds were performed under suitable heating/cooling conditions (see Fig. S14–S31, ESI[†]). Of all derivatives, only compound **2d** displayed exothermic crystallization, setting in at 175 °C (max. at 189 °C) with an enthalpy of 16.1 J g⁻¹, followed by an endothermic melting process beginning at 211 °C (see Fig. 5). Interestingly, compounds **2f**, **3b** and **3c** feature one or two glass transitions at 55 °C, 54 °C (and 119 °C) and 45 °C, respectively. Compounds **2a** and **2b** only showed an endothermic melting process starting at 147 °C and 138 °C, respectively, while compounds **2c–d** did not show any observable thermodynamic processes between 0 °C and 275 °C. In summary, except for compound **2d**, the investigated OAPPDO derivatives generated no crystalline phases under the conditions of film processing and annealing.

2.4. Absorption and PL of OAPPDO thin films

The absorption and photoluminescence spectra of neat thin films of the five OAPPDO derivatives, **2a–2e** and **3a** were recorded. As shown in Fig. 6, all of these films display a well-defined vibronic progression of the $S_0 \rightarrow S_1$ transition between 500 nm and 550 nm. The absorption spectra are very similar in terms of shape and spectral position to the absorption spectra in solution (see Fig. S32, ESI[†]). The principal absorption band in the visible exhibits only a minor shift compared to the spectrum in solution (from 0 nm to 10 nm), which can be attributed solely to the change in the dielectric environment.

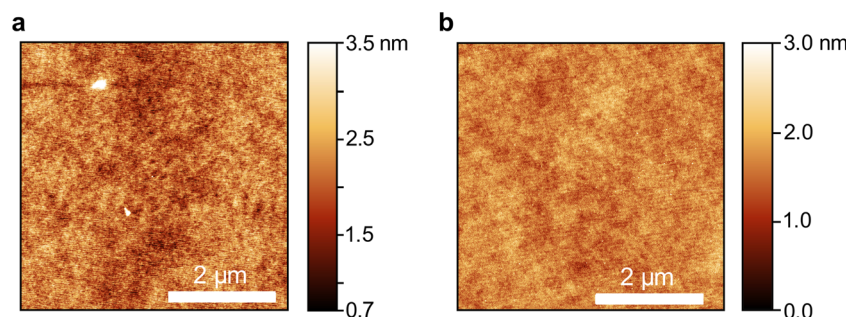


Fig. 4 Atomic force microscopy images of a neat film of (a) compound **2d** and (b) compound **3a**. The scale bar is 2 μm .



Table 2 Summary of root mean square (RMS) roughness, Stokes shifts, PLQYs, and fluorescence lifetimes of spincoated thin films of compounds **2b–e** and **3a**. Reported τ_{avg} are amplitude-averaged lifetimes. The rate constants for radiative (k_r) and non-radiative (k_{nr}) decay are calculated from the amplitude-averaged lifetimes and PLQY

	2b	2c	2d	2e	3a
RMS [nm]	0.40	0.40	0.40	0.40	0.30
Stokes shift [cm^{-1}]	1230	985	930	1495	850
PLQY (%)	2.2	4.5	2.8	17	3.8
τ_1 [ns]	0.23	0.28	0.56	4.20	0.14
τ_2 [ns]	1.88	2.46	3.08	1.00	3.93
τ_3 [ns]	—	0.66	0.20	—	0.70
τ_{avg} [ns]	0.36	0.37	0.37	2.04	0.23
k_r [10^7 s^{-1}]	6.1	12.2	7.6	8.3	16.5
k_{nr} [10^7 s^{-1}]	271.7	258.1	262.7	40.7	418.3

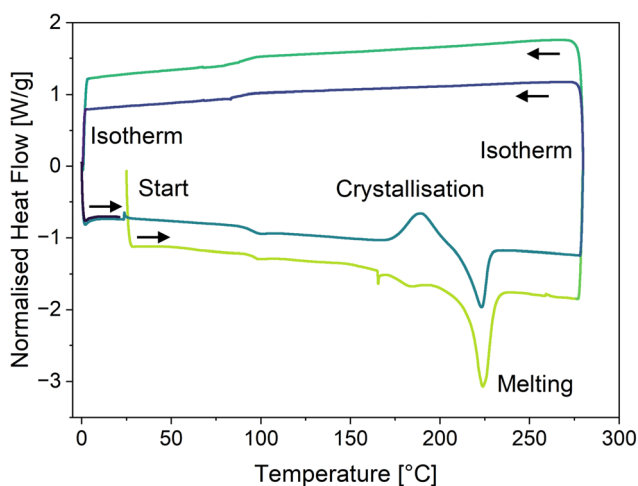


Fig. 5 Differential scanning calorimetry thermogram of compound **2d** showing an exothermic crystallisation process at ca. 180 °C and an endothermic melting process at ca. 220 °C (positive heat flow: exotherm, negative heat flow: endotherm). The first cycle is indicated in green and the second cycle in blue.

There is no sign of aggregate formation or excitonic coupling, excluding any formation of J- or H-aggregates.⁵⁰ The absence of such spectral changes further corroborates the extremely weak π - π and Coulombic interactions between OAPPDO molecules in the solid state as already indicated in the molecular packing (see Fig. 2).

The PL spectra of thin films of compounds **2b–e** display an emission band around 550–600 nm with Stokes shifts of about 1230 cm^{-1} , 985 cm^{-1} , 930 cm^{-1} and 1495 cm^{-1} , respectively (see Table 2) which, except for compound **2e**, is even smaller than the values observed in solution. Interestingly, the solid-state PL of compound **3a** retains its well-resolved vibronic structure (see Fig. 7), which is a rare feature for small organic molecules. It was previously only observed for highly twisted⁴¹ or shielded⁴³ PDIs. It further corroborates the lack of electronic coupling between the molecules. The well-resolved absorption and emission peaks in the neat film make compound **3a** a prime candidate for strong light-matter coupling.

The PLQYs (see Table 2) measured for thin films of compounds **2b–d** and **3a** were significantly reduced (to ca. 3%)

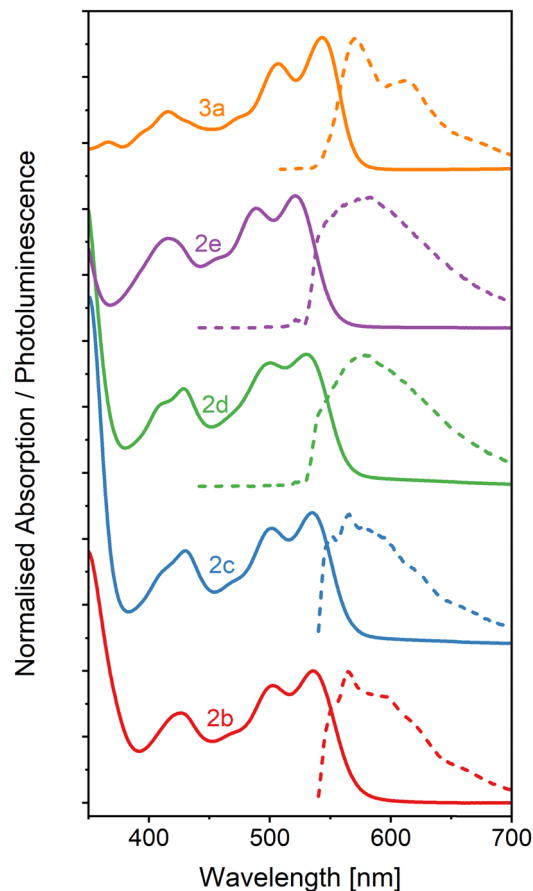


Fig. 6 Absorption (solid lines) and PL (dashed lines) spectra of neat OAPPDO thin films. All films were excited at 532 nm.

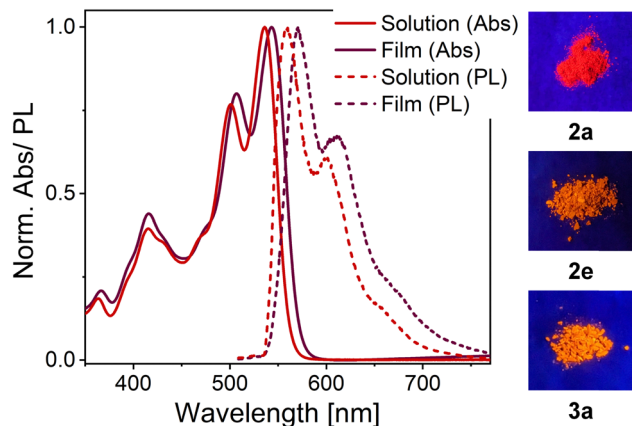


Fig. 7 Absorption (Abs, solid lines) and emission (PL, dashed lines) of compound **3a** in dichloromethane (red) and in neat thin film (dark red). The clear vibronic progression is retained for both absorption and emission spectra of the neat film (left). Photographs of OAPPDOs **2a**, **2e** and **3a** as powders under UV-illumination ($\lambda_{\text{exc}} = 365 \text{ nm}$, right).

compared to their PLQY in solution (for excitation wavelengths see Tables S10 and S11, ESI†). The films also showed multi-exponential PL decays that could be characterized by three lifetimes (τ_1 – τ_3) or one amplitude-averaged lifetime (τ_{avg}) of a



few hundred picoseconds (see Table 2 and Fig. S33–S35 (ESI[†]), for excitation wavelengths and film formation parameters see Table S12, ESI[†]). This fast and multiexponential decay indicates the presence of efficient non-radiative decay channels (quenching) despite the negligible electronic coupling of the molecules. Interestingly, the PLQY of compound **2e** remained fairly high (17%) even for a neat film. This value is comparable to PLQYs reported for neat PDI films.^{9,15,51} The corresponding rate constants for radiative and non-radiative decay were estimated from the measured PLQYs and amplitude-averaged lifetimes (see Table 2). Despite the increased uncertainty of the measured values for thin films compared to solutions, the rate constants for radiative decay remained within the same range as for the solutions. However, the non-radiative decay increased by almost two orders of magnitude for all films except for compound **2e**, for which it only increased by a factor of ten. The amplitude-averaged fluorescence lifetime for thin films of compound **2e** stayed in the nanosecond range and thus similar to that in solution. All of these properties indicate that the chromophores of compound **2e** interact even less than the other OAPPDO compounds. This might be a result of partial shielding of the polycyclic aromatic cores by perfluorophenyl groups on the thioether units in the *bay*-positions. Fluorinated groups are also known to efficiently limit water and oxygen diffusion in molecular thin films, which may also reduce the related quenching.

Furthermore, the photostability of the thin films was investigated under UV light exposure (see Experimental section) in air. Films made of compounds **2b–e** were photobleached within seconds, probably due to the weak photostability of the thioether C–S bonds. However, compound **3a**, with the oxidized sulfone units, showed much higher photostability, with only a 10% reduction of the PL intensity after UV exposure (see Table S13, ESI[†]). It is noteworthy, that encapsulation of the OAPPDO thin films with 10 nm of aluminium oxide resulted in much higher photostability (*i.e.*, less the 5% PL reduction after the same exposure), suggesting that the photobleaching is largely caused by reactive oxygen species. High PLQY but also high photostability of the neat films are basic requirements for any possible observation of polariton lasing in strongly coupled microcavities.

2.5. Strong light-matter coupling in Fabry–Pérot cavities

Given the high density of emitters and the excellent smoothness of neat OAPPDO films, these new fluorophores should be suitable for achieving strong light-matter coupling in cavities. To this end, the films were integrated in Fabry–Pérot cavities (see Fig. 8a). This was accomplished by spincoating neat OAPPDO layers (100–200 nm thickness) on a highly reflective silver mirror (100 nm Ag) and thermal evaporation of 30 nm of silver as a semi-transparent top mirror.

Fig. 8b displays the angle resolved reflectivity and photoluminescence spectra of the highly photostable compound **3a** embedded in a Fabry–Pérot cavity. They clearly show that the bare excitonic transitions (E_{X_1} , E_{X_2} of the vibronic progression of the first singlet transition) and the cavity mode (E_c) are replaced by an upper (UP), middle (MP) and lower (LP) polariton branch

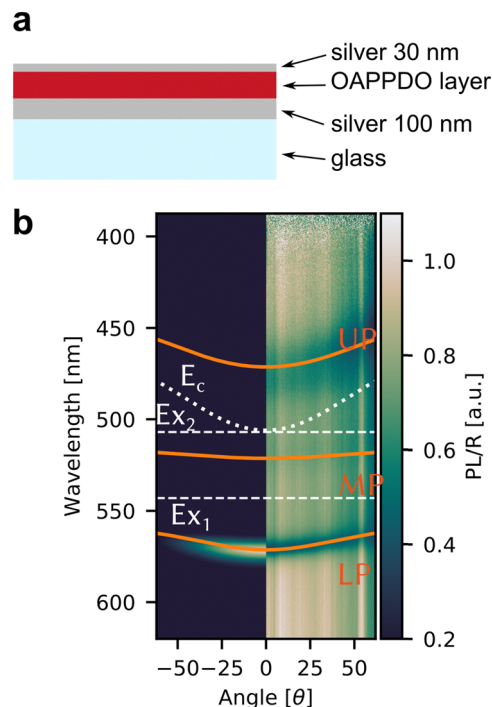


Fig. 8 (a) Schematic cross section of a semi-transparent Fabry–Pérot cavity with a neat OAPPDO layer between two silver mirrors. (b) Angle-resolved PL (left) and reflectivity (right) spectra of a Fabry–Pérot cavity containing compound **3a**. The solid orange lines are the fitted upper (UP), middle (MP) and lower (LP) polaritonic branches, the grey dotted line is the bare cavity mode (E_c). The horizontal grey dashed lines indicate the excitonic transitions (E_{X_1} , E_{X_2}) of compound **3a**.

resulting from hybridization as shown previously for molecules with prominent vibronic progression.¹⁷ The avoided crossing of the polariton modes and the excitonic transitions is a strong indication that the cavity is indeed in the strong coupling regime. PL only occurs from the LP mode and hence fitting of the three polariton modes was performed based on the reflectance data. The same observations were made for cavities with compounds **2b–e** (see Fig. S36, ESI[†]). The linewidth of the polariton modes enabled the extraction of the quality factors of the cavities, which range between 20 and 64. These values are very good for simple silver mirrors and are a direct result of the excellent smoothness of the OAPPDO thin films.

From the angle-resolved reflectivity spectrum of the cavity (right panel of Fig. 8b), the Rabi splitting $\hbar\Omega_R$ can be extracted by fitting the UP, MP and LP modes with a coupled harmonic oscillator model, taking into account the two excitonic transitions and the cavity mode (see Experimental section). The Rabi splitting for a cavity with compound **3a** reached 304 meV for the first excitonic transition and 290 meV for the second excitonic transition. The data for cavities with all other OAPPDO derivatives are summarised in Table 3. The extracted values for $\hbar\Omega_R$ are significantly larger than the linewidths of both the bare excitonic transitions and the bare cavity mode, hence confirming again that the cavities are in the strong coupling regime.

A comparison of the Rabi splittings (304 meV and 290 meV) for compound **3a** with its excitonic transition energies (2.17 eV



Table 3 Summary of the Rabi splittings $\hbar\Omega_R$, bare cavity energy E_c and the quality-factor Q for Fabry–Pérot cavities containing compounds **2b–e** and **3a**

Compound	2b	2c	2d	2e	3a
$\hbar\Omega_1$ [meV]	270	240	242	280	304
$\hbar\Omega_2$ [meV]	190	220	238	200	290
E_c [eV]	2.45	2.33	2.30	2.40	2.45
Q	48	37	48	21	64

and 2.03 eV) indicates that the cavities are approaching the so-called ultrastrong coupling regime, *i.e.*, the coupling strength becomes comparable (>10 – 20%) to the molecular transition energy.⁵² It is difficult to reach the ultrastrong coupling regime in a cavity with diluted emitters in a matrix, hence, it is usually overserved for neat films of emitters with high oscillator strength.^{53,54} Such strong coupling also leads to a very flat dispersion of the lower polariton modes and hence nearly angle-independent emission wavelengths, which is very useful for light-emitting devices.²⁴ These observations emphasize again the excellent film quality and high molecular density of neat OAPPDO films as well as their strong light absorbing and emitting properties.

3. Conclusion

In summary, we have presented a new class of amorphous solid-state emitters, which are derivatives of octaazaperopyrenedioxide (OAPPDO), an *N*-heteropolycyclic dye with a non-planar azaperylene-derived π -system at its core. The introduction of sulfur-based groups at the *bay*-positions of the tetraazaperylene core leads to a torsional twist in the π -core of about 30° that results in highly soluble compounds that exhibit high molar extinction coefficients and photoluminescence quantum yields in solution. Their tailored molecular properties allow the OAPPDOs to be solution-processed into high density neat films with monomer-like absorption and emission properties and photoluminescence quantum yields up to 17%. Those thin films are amorphous at room temperature, highly homogeneous and exhibit exceptional surface smoothness. Moreover, OAPPDO films are photostable against UV irradiation when encapsulated while the sulfone derivatives were found to be photostable even without any encapsulation. Finally, we demonstrated strong light-matter coupling of thin films of OAPPDOs in simple metal-clad microcavities. We believe that this new class of amorphous solid-state emitters provides promising candidates for optoelectronic and photonic applications where amorphous and highly emissive layers are required.

4. Experimental section

4.1. Synthesis

The detailed synthetic protocol for all compounds is described in the ESI.†

4.2. Thermal characterization of materials

TGA (thermal gravimetric analysis) measurements were performed with a Mettler Toledo TGA 2 device. DSC (differential scanning calorimetry) was performed using a TA Instruments Discovery DSC 250 calorimeter (**2b**, **2c**, **2d**, **2e**) or a Mettler Toledo DSC821 (**2a**, **2f**, **3a**, **3b**, **3c**).

4.3. Thin film and preparation

Cleaned, alkali-free glass substrates (AF32eco, Schott, $20 \times 25 \times 0.5$ mm³) and EagleXG glass substrates were used for all thin film measurements. Compounds **2b–e** were dissolved in *o*-xylene and compound **3a** in DCM, respectively. Neat films were prepared by spincoating directly from solution on cleaned glass substrates (for parameters see Table S9, ESI†). Atomic force microscopy images were recorded with a Bruker Dimension Icon atomic force microscope (AFM) in ScanAsyst[®] mode. The root mean square surface roughness was extracted using Gwyddion 2.59 software. For encapsulation, a 10 nm AlO_x layer was deposited on top of the neat OAPPDO layer by atomic layer deposition (ALD) (Ultratech, Savannah S100, precursor trimethyl-aluminium, Strem Chemicals, Inc.) at low temperature (80 °C) to avoid thermal degradation of the molecules. For Fabry–Pérot cavities, a reflective bottom silver mirror (100 nm) was deposited by thermal evaporation onto a cleaned glass substrate. The OAPPDO layer was spincoated on top and a 30 nm silver mirror completed the cavity.

4.4. Optical characterization

Absorption spectra were acquired with an Agilent Cary 6000i or Cary 5000i UV-Vis-NIR absorption spectrometer. Photoluminescence spectra in solution were recorded with a Varian Cary Eclipse fluorescence spectrophotometer and photoluminescence quantum yields (PLQY) were measured on a JASCO spectrofluorometer FP-8500 equipped with an ILF-835j100 mm integrating sphere. PL spectra of thin films were obtained by excitation with a 532 nm laser diode (OBIS, cw) and collection of the emitted light with a 60× objective (Olympus, NA = 0.9). Scattered laser light was blocked with a 550 long pass filter. The emission was imaged onto the entrance slit of a Princeton Instruments IsoPlane SCT 320 spectrometer and spectra were acquired with a PIXIS:400 BR camera (Princeton Instruments).

For PLQY values of thin films, the emitter materials were spin-coated onto EagleXG glass substrates as described above. They were subsequently characterized in a PLQY measurement system (Hamamatsu Photonics) equipped with an integrating sphere, xenon lamp (L9799-01), monochromator (A10080-01), and multi-channel analyser (C10027). The samples were excited at different wavelengths (see Table S10, ESI†). All data were analyzed using the Hamamatsu U6039-06 software. All PLQY measurements were performed in dry nitrogen.

All optical images of thin films were acquired using an Olympus BX51 microscope. The optical constants and thickness of the thin films (on EagleXG glass) were determined from variable angle spectroscopic ellipsometry (M2000, J.A. Woollam),



transmission spectra, and subsequent modeling (CompleteEase software, J.A. Woollam) using a general oscillator model (see Fig. S37, ESI†).

4.5. Angle-resolved measurements

Angle-resolved reflectivity spectra of the Fabry–Pérot cavities were measured in a home-built Fourier microscopy setup^{55,56} using a halogen light source (Ocean Optics, HL-2000-HP) focused onto the sample *via* a 60× objective (Olympus, MPLAPON60X) with a numerical aperture of 0.9. For photoluminescence measurements, a laser diode (OBIS, CW 532 nm) was used as an excitation source, and the scattered laser light was blocked by a long-pass filter (550 nm cut-off). The reflected/emitted light was imaged from the back focal plane of the objective onto the entrance slit of the spectrometer (Princeton Instruments IsoPlane SCT 320). A linear polarizer was placed in front of the spectrometer to select TE polarization. Angle-resolved spectra were recorded with a Si CCD camera (Princeton Instruments, PIXIS:400).

4.6. Fluorescence lifetime measurements

For fluorescence lifetime measurements by time-correlated single-photon counting (TCSPC), dilute solutions and thin films of compounds **2b** and **2e** were excited with the spectrally filtered output of a picosecond-pulsed super-continuum laser source. A dichroic long-pass filter blocked scattered laser light (550 nm cut-off). The collected emission was focused onto a silicon avalanche photodiode (Micro Photon Devices) and a time-correlated single-photon counting module (PicoHarp 300, PicoQuant) recorded the PL decay traces, which were fitted as a mono-exponential or multi-exponential decay. Time-resolved PL decays of dilute solutions and thin films of compounds **2c–d** and **3a** were measured with a time-correlated single photon counting setup (FluoTime 250, TimeHarp 260, PMA Hybrid detector, PicoQuant) at room temperature at an excitation wavelength of 373 nm. The employed excitation wavelengths and solvents are given in Tables S11 and S12 (ESI†).

4.7. Photostability measurements

The photostability of thin films of all compounds was estimated using the following procedure. A fluorescence image was recorded with an Olympus BX51 fluorescence microscope under UV excitation using a 10× objective. Samples were then exposed to the same UV light using a 20× objective for 5 minutes. Finally, a fluorescence image was recorded using the same parameters as the first image. The resulting photostability is reported as the decrease of fluorescence intensity before and after UV exposure or as ‘bleached’ when no more fluorescence could be recorded from the exposed area.

4.8. Coupled harmonic oscillator (CHO) fitting procedure

The reflectivity dispersion plots were fitted using a 3 × 3 coupled harmonic oscillator (CHO) model to extract couplings

strengths. Solving the following Hamiltonian leads to the polariton energy as eigenvalues:

$$H = \begin{pmatrix} E_c(\theta) & V_1 & V_2 \\ V_1 & E_1 & 0 \\ V_2 & 0 & E_2 \end{pmatrix}$$

where E_1 and E_2 are the exciton energies, V_1 and V_2 are the coupling strengths for the corresponding transitions. The cavity energy E_c as a function of angle θ is given by:

$$E_c(\theta) = \frac{E_c(0)}{\sqrt{1 - \left(\frac{\sin(\theta)}{n_{\text{eff}}}\right)^2}}$$

with n_{eff} the effective refractive index.

The quality factors (Q) of the cavities were determined using:

$$Q = \frac{\lambda}{\Delta\lambda}$$

where λ is the wavelength of the lower polariton mode at $\theta = 0^\circ$, and $\Delta\lambda$ is the full width at half maximum.

Author contributions

M. H. performed optical experiments if not otherwise noted and analyzed the data. R. E. and P. J. carried out all syntheses and chemical characterizations. T. W. proofread the manuscript. J. B. collected and analyzed the crystallographic data. S. S. and S. L. F. performed and analyzed the TCSPC measurements of compound **2a–b**, **2f** and **3b–c**. A. M., F. L. R. and F. T. C. carried out the ellipsometry measurements, PLQY and lifetime measurements of thin films of compounds **2c–d** and **3a** under the supervision of M. C. G.; J. Z. and L. H. G. conceived and supervised the project. J. Z., L. H. G., R. E., and M. H. co-wrote the manuscript with input from all co-authors.

Conflicts of interest

There are no conflicts to declare.

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References

- H. Uoyama, K. Goushi, K. Shizu, H. Nomura and C. Adachi, *Nature*, 2012, **492**, 234–238.
- S. Reineke, F. Lindner, G. Schwartz, N. Seidler, K. Walzer, B. Lüssem and K. Leo, *Nature*, 2009, **459**, 234–238.
- J. Roncali, *Adv. Energy Mater.*, 2020, **10**, 2001907.
- B. Zhang, P. Zhao, L. J. Wilson, J. Subbiah, H. Yang, P. Mulvaney, D. J. Jones, K. P. Ghiggino and W. W. H. Wong, *ACS Energy Lett.*, 2019, **4**, 1839–1844.
- A. J. C. Kuehne and M. C. Gather, *Chem. Rev.*, 2016, **116**, 12823–12864.
- Y. Jiang, Y.-Y. Liu, X. Liu, H. Lin, K. Gao, W.-Y. Lai and W. Huang, *Chem. Soc. Rev.*, 2020, **49**, 5885–5944.
- A. Huber, J. Dubbert, T. D. Scherz and J. Voskuhl, *Chem. – Eur. J.*, 2023, **29**, e202202481.
- J. Gierschner, J. Shi, B. Milián-Medina, D. Roca-Sanjuán, S. Varghese and S. Park, *Adv. Opt. Mater.*, 2021, **9**, 2002251.
- B. Zhang, H. Soleimaninejad, D. J. Jones, J. M. White, K. P. Ghiggino, T. A. Smith and W. W. H. Wong, *Chem. Mater.*, 2017, **29**, 8395–8403.
- M. G. Ramírez, S. Pla, P. G. Boj, J. M. Villalvilla, J. A. Quintana, M. A. Díaz-García, F. Fernández-Lázaro and Á. Sastre-Santos, *Adv. Opt. Mater.*, 2013, **1**, 933–938.
- Y. Wen, T. Sheng, X. Zhu, C. Zhuo, S. Su, H. Li, S. Hu, Q.-L. Zhu and X. Wu, *Adv. Mater.*, 2017, **29**, 1700778.
- J. Yu, Y. Cui, H. Xu, Y. Yang, Z. Wang, B. Chen and G. Qian, *Nat. Commun.*, 2013, **4**, 2719.
- M. J. Hollamby and T. Nakanishi, *J. Mater. Chem. C*, 2013, **1**, 6178–6183.
- D. Schmidt, M. Stolte, J. Süß, A. Liess, V. Stepanenko and F. Würthner, *Angew. Chem., Int. Ed.*, 2019, **58**, 13385–13389.
- M. Stolte, T. Schembri, J. Süß, D. Schmidt, A.-M. Krause, M. O. Vysotsky and F. Würthner, *Chem. Mater.*, 2020, **32**, 6222–6236.
- D. G. Lidzey, D. D. C. Bradley, M. S. Skolnick, T. Virgili, S. Walker and D. M. Whittaker, *Nature*, 1998, **395**, 53–55.
- R. J. Holmes and S. R. Forrest, *Phys. Rev. Lett.*, 2004, **93**, 186404.
- A. Graf, L. Tropsch, J. Zaumseil and M. C. Gather, in *Handbook of Organic Materials for Electronic and Photonic Devices*, ed. O. Ostroverkhova, Woodhead Publishing, 2nd edn, 2019, ch. 9, pp. 281–307.
- J. R. Tischler, M. S. Bradley, V. Bulović, J. H. Song and A. Nurmikko, *Phys. Rev. Lett.*, 2005, **95**, 036401.
- B. Siegmund, A. Mischok, J. Benduhn, O. Zeika, S. Ullbrich, F. Nehm, M. Böhm, D. Spoltore, H. Fröb, C. Körner, K. Leo and K. Vandewal, *Nat. Commun.*, 2017, **8**, 15421.
- A. Mischok, J. Lüttgens, F. Berger, S. Hillebrandt, F. Tenopala-Carmona, S. Kwon, C. Murawski, B. Siegmund, J. Zaumseil and M. C. Gather, *J. Chem. Phys.*, 2020, **153**, 201104.
- E. Eizner, J. Brodeur, F. Barachati, A. Sridharan and S. Kéna-Cohen, *ACS Photonics*, 2018, **5**, 2921–2927.
- S. Gambino, M. Mazzeo, A. Genco, O. Di Stefano, S. Savasta, S. Patané, D. Ballardini, F. Mangione, G. Lerario, D. Sanvitto and G. Gigli, *ACS Photonics*, 2014, **1**, 1042–1048.
- A. Mischok, S. Hillebrandt, S. Kwon and M. C. Gather, *Nat. Photonics*, 2023, **17**, 393–400.
- S. Kéna-Cohen and S. R. Forrest, *Nat. Photonics*, 2010, **4**, 371–375.
- J. D. Plumhof, T. Stöferle, L. Mai, U. Scherf and R. F. Mahrt, *Nat. Mater.*, 2014, **13**, 247–252.
- T. Cookson, K. Georgiou, A. Zasedatelev, R. T. Grant, T. Virgili, M. Cavazzini, F. Galeotti, C. Clark, N. G. Berloff, D. G. Lidzey and P. G. Lagoudakis, *Adv. Opt. Mater.*, 2017, **5**, 1700203.
- C. P. Dietrich, A. Steude, L. Tropsch, M. Schubert, N. M. Kronenberg, K. Ostermann, S. Höfling and M. C. Gather, *Sci. Adv.*, 2016, **2**, e1600666.
- S. K. Rajendran, M. Wei, H. Ohadi, A. Ruseckas, G. A. Turnbull and I. D. W. Samuel, *Adv. Opt. Mater.*, 2019, **7**, 1801791.
- T. Ishii, K. Miyata, M. Mamada, F. Bencheikh, F. Mathevet, K. Onda, S. Kéna-Cohen and C. Adachi, *Adv. Opt. Mater.*, 2022, **10**, 2102034.
- T. Wesp, P. Valsalan, A. Kochan, M. Hertzog, H. Wadepohl, J. Zaumseil and L. H. Gade, *Chem. – Eur. J.*, 2022, **28**, e202202661.
- T. Wesp, T. Bruckhoff, J. Petry, H. Wadepohl and L. H. Gade, *Chem. – Eur. J.*, 2022, **28**, e202200129.
- T. Wesp, T. Bruckhoff, H. Wadepohl and L. H. Gade, *Chem. – Eur. J.*, 2022, **28**, e202201706.
- R. Eichelmann, J. Monti, L.-Y. Hsu, F. Kröger, J. Ballmann, E. Blasco and L. H. Gade, *Mol. Syst. Des. Eng.*, 2023, **8**, 1470–1476.
- Y. Zhou, B. Xue, C. Wu, S. Chen, H. Liu, T. Jiu, Z. Li and Y. Zhao, *Chem. Commun.*, 2019, **55**, 13570–13573.
- Z. Chen, M. G. Debije, T. Debaerdemaeker, P. Osswald and F. Würthner, *ChemPhysChem*, 2004, **5**, 137–140.
- Z. Chen, U. Baumeister, C. Tschierske and F. Würthner, *Chem. – Eur. J.*, 2007, **13**, 450–465.
- T. E. Kaiser, H. Wang, V. Stepanenko and F. Würthner, *Angew. Chem., Int. Ed.*, 2007, **46**, 5541–5544.
- T. E. Kaiser, V. Stepanenko and F. Würthner, *J. Am. Chem. Soc.*, 2009, **131**, 6719–6732.
- F. Würthner, C. R. Saha-Möller, B. Fimmel, S. Ogi, P. Leowanawat and D. Schmidt, *Chem. Rev.*, 2016, **116**, 962–1052.
- K. Nagarajan, A. R. Mallia, V. S. Reddy and M. Hariharan, *J. Phys. Chem. C*, 2016, **120**, 8443–8450.
- K. M. Felter, R. K. Dubey and F. C. Grozema, *J. Chem. Phys.*, 2019, **151**, 094301.
- Á. J. Jiménez, M.-J. Lin, C. Burschka, J. Becker, V. Settels, B. Engels and F. Würthner, *Chem. Sci.*, 2014, **5**, 608–619.
- C. Lee, W. Yang and R. G. Parr, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1988, **37**, 785–789.
- A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648–5652.
- F. Weigend and R. Ahlrichs, *Phys. Chem. Chem. Phys.*, 2005, **7**, 3297–3305.
- F. Weigend, *Phys. Chem. Chem. Phys.*, 2006, **8**, 1057–1065.
- S. Grimme, J. Antony, S. Ehrlich and H. Krieg, *J. Chem. Phys.*, 2010, **132**, 154104.



- 49 S. Grimme, S. Ehrlich and L. Goerigk, *J. Comput. Chem.*, 2011, **32**, 1456–1465.
- 50 N. J. Hestand and F. C. Spano, *Chem. Rev.*, 2018, **118**, 7069–7163.
- 51 M.-J. Lin, Á. J. Jiménez, C. Burschka and F. Würthner, *Chem. Commun.*, 2012, **48**, 12050–12052.
- 52 P. Forn-Díaz, L. Lamata, E. Rico, J. Kono and E. Solano, *Rev. Mod. Phys.*, 2019, **91**, 025005.
- 53 S. Kéna-Cohen, S. A. Maier and D. D. C. Bradley, *Adv. Opt. Mater.*, 2013, **1**, 827–833.
- 54 M. Held, A. Graf, Y. Zakharko, P. Chao, L. Tropsch, M. C. Gather and J. Zaumseil, *Adv. Opt. Mater.*, 2018, **6**, 1700962.
- 55 A. Graf, L. Tropsch, Y. Zakharko, J. Zaumseil and M. C. Gather, *Nat. Commun.*, 2016, **7**, 13078.
- 56 J. M. Lüttgens, F. J. Berger and J. Zaumseil, *ACS Photonics*, 2021, **8**, 182–193.

