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# Dissecting non-radiative decay in donor-functionalized radicals with a mode-resolved model

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Donor-functionalized radicals based on the poly-chlorinated trityl moiety are an emerging class of efficient emitters for next-generation optoelectronics. Their performance is governed by the non-radiative decay of a charge-transfer (CT) state, making its understanding crucial for molecular design. We present a mode-resolved model that links the decay to structural motifs through molecular vibrations. The decisive role of exact exchange in density functional modeling is highlighted, alongside solvation and anharmonic effects. Using representative radical emitters, we show that suppression of non-radiative decay in systems with a perpendicular donor–acceptor arrangement arises from reduced coupling of the promoting mode to the CT exciton and a diminished anharmonic contribution. To elucidate this mechanism, we perform a sensitivity analysis of the contributions of individual modes to the Franck–Condon weighted density of states, including anharmonic effects.

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## 1 Introduction

Organic molecular emitters are widely applied in optoelectronic devices, while ongoing research continues to optimize their photophysical properties through molecular design.<sup>1–3</sup> Conventionally, these emitters have a closed-shell ground state and suffer radiative losses due to optically dark triplet states.<sup>4</sup> Donor-functionalized radical emitters based on the tris(2,4,6-trichlorophenyl)methyl (TTM) moiety circumvent this issue by having a doublet ground state. In these radical emitters, radiative losses are primarily associated with the non-radiative decay of the intramolecular charge-transfer (CT) state, which arises from electron transfer between the donor and radical (acceptor) moieties.<sup>5–7</sup>

Gaining a theoretical understanding of non-radiative decay is crucial for guiding molecular design. The energy-gap law,<sup>8,9</sup> supported by both theory and experiment, dictates that the rate constant  $k_{\text{nr}}$  for non-radiative decay falls off exponentially as the energy gap  $\Delta E$  increases,

$$k_{\text{nr}} = \frac{C^2 \sqrt{2\pi}}{\hbar \sqrt{\hbar \omega_{\text{e}} \Delta E}} \exp \left\{ -\frac{\Delta E}{\hbar \omega_{\text{e}}} \left[ \ln \left( \frac{\Delta E}{\sum_{\mu} A_{\mu}} \right) - 1 \right] \right\}. \quad (1)$$

The transition is driven by high-frequency modes with index  $\mu$ , which contribute to the reorganization energy by  $A_{\mu}$  and are

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combined into an effective mode with averaged angular frequency  $\omega_{\text{e}}$ . The parameter  $C$  describes the coupling between the electronic states involved in the transition.

Donor-functionalized TTM radicals do not conform to the energy-gap law, as they emit in the deep-red/near-infrared region yet exhibit relatively slow non-radiative decay rates.<sup>10,11</sup> In addition, chemical modifications of either the donor or the acceptor unit can lead to a further marked reduction of non-radiative decay without substantially altering the energy gap.<sup>11,12</sup> This behavior has been attributed to suppressed couplings of the promoting high-frequency vibrations to the CT exciton, either due to the localization of vibrations on the donor or acceptor moiety, or as a consequence of orbital effects arising from molecular symmetry.<sup>11,13</sup>

Quantitative modeling of non-radiative decay in donor-functionalized radicals has so far relied on the Marcus–Levich–Jortner (MLJ) model,<sup>14–18</sup> combining a classical description of low-frequency modes with a harmonic quantum-mechanical treatment of an effective (e) mode representing all high-frequency vibrations,

$$k_{\text{nr}}^{(\text{MLJ})} = \frac{C^2 \sqrt{2\pi}}{\hbar \sigma} \sum_{\nu} F_{\nu, \text{e}} \exp \left[ \frac{1}{2} \left( \frac{\nu \hbar \omega_{\text{e}} + A_{\text{c}} - \Delta E}{\sigma} \right)^2 \right]. \quad (2)$$

In the above expression, the sum runs over the vibrational quantum number  $\nu$  of the effective mode with harmonic Franck–Condon factor  $F_{\nu, \text{e}}$ . The width parameter  $\sigma$  is related to the classical reorganization energy  $A_{\text{c}}$  and the temperature  $T$  via  $\sigma = \sqrt{2A_{\text{c}}k_{\text{B}}T}$ . Due to the size of the radical emitters, only



density functional theory (DFT) is practically feasible, and the CT character of these systems requires the use of range-separated hybrid functionals with additional long-range exact exchange (eX).<sup>19–22</sup>

Because the effective mode in the MLJ model does not directly correspond to a particular molecular vibration, it is challenging to relate non-radiative decay to molecular design. Here, we present a mode-resolved quantitative model that explicitly connects non-radiative decay to the molecular structure. We begin by addressing key theoretical considerations, covering the effects of long-range eX, solvation, and anharmonicity. Using the systems under investigation, we show that a quantitative and physically sound description of non-radiative decay is possible. The decay mechanism can be related to particular molecular vibrations, allowing the observed trends in rate constants  $k_{\text{nr}}$  to be interpreted in an intuitive manner.

## 2 Theoretical framework

In the present work, we investigate the non-radiative decay in donor-functionalized radical emitters in solution. The intramolecular charge-transfer (CT) character of the decaying state represents a crucial aspect for theoretical modeling. In addition to discussing the theoretical details of the rate constant  $k_{\text{nr}}$  for non-radiative decay, we also provide remarks on the description of CT states within the (time-dependent) formalism of density functional theory ((TD)-DFT). Furthermore, we briefly address the treatment of solvation effects using state-specific continuum solvation models and include a short note on the rate constant  $k_{\text{r}}$  for radiative decay.

### 2.1 Non-radiative decay

Our treatment of non-radiative decay by internal conversion is inspired by the MLJ model which combines a classical treatment of low-frequency modes with a (harmonic) quantum mechanical treatment of high-frequency vibrations in terms of an effective mode.<sup>14–18</sup> Analogous to the MLJ model, we essentially adopt the classical description from Marcus theory.<sup>14,15</sup> For the quantum-mechanical description, however, we abandon the one-effective mode model and explicitly consider the (harmonic) contributions from all quantum-mechanical modes. Moreover, we introduce anharmonic corrections to contributions of selected high-frequency modes based on the Morse potential.<sup>23</sup> We note that our approach can be regarded as a reduced linear vibronic coupling model. In this section, we present the complete expression for the rate constant  $k_{\text{nr}}$  (including classical and quantum-mechanical contributions) and focus on the mode-resolved quantum-mechanical treatment of high-frequency vibrations.

We note that the quantum-mechanical mode contributions are distinguished as either harmonic or anharmonic. In Section S1.1 of the SI, we provide a detailed derivation of the rate constant considering exclusively harmonic (h) contributions, along with remarks on the complete rate constant, which also includes classical (c) and anharmonic (a) contributions.<sup>14–18,24–39</sup> The final

and complete expression for the rate constant can be written as

$$k_{\text{nr}}^{(\text{cha})} = \frac{\pi}{\hbar} |\mathbf{C}|^2 \langle \rho_{\text{cha}} \rangle_{(\Delta E)}. \quad (3)$$

It should be emphasized that this expression relies on the Condon approximation.<sup>30–32</sup> In the above expression, the first leading contribution is the squared norm of the first-order non-adiabatic coupling vector  $\mathbf{C}$ , which is composed of the normal mode projections  $C_{\mu}$ ,

$$C_{\mu} = \hbar \omega_{\mu} \left\langle \Psi_{\text{i}} \left| \frac{\partial}{\partial Q_{\mu}} \right| \Psi_{\text{f}} \right\rangle_{\mathbf{Q}=\mathbf{0}}. \quad (4)$$

The (dimensionless) normal coordinate and (harmonic) angular frequency of mode  $\mu$  are denoted as  $Q_{\mu}$  and  $\omega_{\mu}$ , respectively, and the wavefunctions  $\Psi_{\text{i}}$  and  $\Psi_{\text{f}}$  refer to the initial (i) and final (f) adiabatic electronic state. We remark that  $\mathbf{C}$  is evaluated at the displacement vector  $\mathbf{Q} = \mathbf{0}$ , corresponding to the equilibrium geometry of the emitting state.

The second leading contribution in eqn (3) is the total Franck–Condon weighted density of states  $\rho_{\text{cha}}$  (FCWD)<sup>31,32,39</sup>—including classical, harmonic and anharmonic mode contributions—averaged around the energy gap  $\Delta E$  of the non-radiative transition. Classifying the normal modes as classical, harmonic, or anharmonic, the total FCWD  $\rho_{\text{cha}}$  can be represented as a convolution of the FCWDs corresponding to the respective mode classes,

$$\rho_{\text{cha}} = \rho_{\text{c}} \circ \rho_{\text{h}} \circ \rho_{\text{a}} = \rho_{\text{c}} \circ \rho_{\text{q}}. \quad (5)$$

In the second equality of the previous expression, we summarized the harmonic and anharmonic FCWDs as the overall quantum-mechanical contribution  $\rho_{\text{q}}$ .

In the following, we outline the concepts relevant to a mode-resolved description of non-radiative decay, with the harmonic FCWD  $\rho_{\text{h}}$  as an illustrative example. As shown in Section S1.1 of the SI,  $\rho_{\text{h}}$  is given as a function of the energy gap  $\Delta E$  by

$$\rho_{\text{h}}(\Delta E) = \sum_{\nu_{\mu}} \prod_{\mu} F_{\mu, \nu_{\mu}}^{(\text{h})} \delta \left( \Delta E - \sum_{\mu} \varepsilon_{\mu, \nu_{\mu}}^{(\text{h})} \right). \quad (6)$$

The first summation in the above expression runs over all possible combinations of quantum numbers  $\nu_{\mu}$ —corresponding to vibrational states within the final electronic state—associated with the vibrational energies  $\varepsilon_{\mu, \nu_{\mu}}^{(\text{h})} = \nu_{\mu} \hbar \omega_{\mu}$ . Assuming non-radiative decay from the vibrational ground state of the emitting state, the harmonic Franck–Condon (FC) factors  $F_{\mu, \nu_{\mu}}^{(\text{h})}$  are given by Poisson distributions of the quantum numbers  $\nu_{\mu}$ ,<sup>36</sup>

$$F_{\mu, \nu_{\mu}} = \frac{S_{\mu}^{\nu_{\mu}}}{\nu_{\mu}!} e^{-S_{\mu}}. \quad (7)$$

It should be emphasized that the above expression implies the displaced harmonic oscillator model (see also Section S1.1 of the SI). The Huang–Rhys (HR) factor  $S_{\mu}$  can be interpreted as a direct measure for the coupling between a mode  $\mu$  and the exciton.<sup>35</sup> Denoting the (dimensionless) displacement of the



equilibrium position of mode  $\mu$  between initial and final vibrational state as  $\Delta_\mu$  and the corresponding contribution to the total reorganization energy  $\Lambda$  as  $\Lambda_\mu$ , the HR factor  $S_\mu$  can be expressed as

$$S_\mu = \frac{A_\mu}{\hbar\omega_\mu} = \frac{1}{2}A_\mu^2. \quad (8)$$

Within the displaced harmonic oscillator model, the mode displacements  $\Delta_\mu$  can be related to the projections  $G_\mu$  of the initial state electronic gradient  $\mathbf{G}$  evaluated at the equilibrium geometry of the final electronic state,

$$\Delta_\mu = \sqrt{\omega_\mu} \left( \frac{G_\mu}{\omega_\mu^2} \right), \quad (9)$$

see also Sections S1.1 and S2.2 of the SI for additional details. The excitonic coupling of a mode  $\mu$ ,  $S_\mu$ , can thus be associated with the extent to which the corresponding vibrational motion perturbs the energy of the initial (excited) electronic state.

According to eqn (6), a mode  $\mu$  contributes substantially to  $\rho_h$  if the FC factors  $F_{\mu,\nu_\mu}^{(h)}$  are large at vibrational energies  $\varepsilon_{\mu,\nu_\mu}^{(h)}$  near the gap  $\Delta E$ . The required vibrational quantum number  $\nu_\mu$  is relatively small for a high-frequency mode. Hence, a high-frequency mode with significant excitonic coupling  $S_\mu$  accelerates non-radiative decay due to a large vibronic transition probability  $F_{\mu,\nu_\mu}^{(h)}$  at the gap  $\Delta E$  of the transition.

To conclude our examination of quantum mechanical contributions, we consider the inclusion of anharmonic corrections in terms of displaced Morse oscillators. The concepts introduced above can also be extended to anharmonic oscillators, which are essential for the description of (highly) excited vibrational states. A convenient description of anharmonic vibrations can be achieved by using the Morse potential,<sup>23,28,40,41</sup>

$$V_\mu^{(a)} = \tilde{D}_{e,\mu}(1 - e^{-a_\mu Q_\mu})^2, \quad (10)$$

with the well-depth  $\tilde{D}_{e,\mu}$  and the stiffness  $a_\mu$ .

The anharmonic FCWD,  $\rho_a$ , can be obtained from eqn (7) by substituting  $F_{\mu,\nu_\mu}^{(h)}$  and  $\varepsilon_{\mu,\nu_\mu}^{(h)}$  with the corresponding expressions  $F_{\mu,\nu_\mu}^{(a)}$  and  $\varepsilon_{\mu,\nu_\mu}^{(a)}$  derived for displaced Morse oscillators. The presumably small displacements  $\Delta_\mu$  between the equilibrium positions of the Morse oscillators are thereby adopted from the displaced harmonic oscillator model (*cf.* eqn (9)). While the harmonic FC factors  $F_{\mu,\nu_\mu}^{(h)}$  in eqn (7) are invariant with respect to the direction of  $\Delta_\mu$  due to the symmetry of the harmonic potentials, the sign of  $\Delta_\mu$  becomes crucial in the anharmonic case.

In Section S1.2 of the SI, we provide a detailed account on relevant expressions for displaced Morse oscillators and discuss possible effects of the anharmonic contributions.<sup>23,28,40–45</sup> The displacements  $\Delta_\mu$  are defined as positive when the average change in bond lengths, relative to the initial state and determined by projection onto internal coordinates, is positive. For a high-frequency mode, the vibronic transition terminates near the inner turning point of the (final) Morse potential if  $\Delta_\mu > 0$ ,

in which case an acceleration of non-radiative decay due to anharmonic contributions can be anticipated (see Section S1.2).

The well-depth for all anharmonic modes is chosen as  $\tilde{D}_{e,\mu} = 30\,000\text{ cm}^{-1}$  as suggested by previous works.<sup>28,40,41</sup> Section S1.3 of the SI provides a brief discussion of this choice and indicates its justification through a stability analysis of the total FCWD  $\rho_{\text{cha}}$ .<sup>28,40,41,46</sup>

## 2.2 Low-frequency modes

For vibrational modes with sufficiently low frequency, the spacings between adjacent vibrational energy levels are small relative to the thermal energy  $k_B T$  at relevant temperatures  $T$  (room temperature), which motivates a classical description. In analogy to the MLJ model, our distinction between classical (c) and quantum-mechanical (q) modes corresponds to a division of the total reorganization energy  $\Lambda$  for displaced harmonic oscillators,<sup>14–18</sup>

$$\Lambda = \sum_\mu \hbar\omega_\mu S_\mu = \Lambda_c + \Lambda_q. \quad (11)$$

The above summation runs over all normal modes  $\mu$  and the reorganization energies  $\Lambda_c$  and  $\Lambda_q$  collect the contributions from classical and quantum-mechanical modes, respectively.

We adopt the expression for the classical FCWD  $\rho_c$  from Marcus theory, which corresponds to a Gaussian line shape function centered at the classical reorganization energy  $\Lambda_c$ .<sup>14,15</sup>

$$\rho_c(\Delta E) = \frac{1}{\sigma_c \sqrt{2\pi}} \exp \left[ -\frac{1}{2} \left( \frac{\Delta E - \Lambda_c}{\sigma_c} \right)^2 \right]. \quad (12)$$

The Gaussian width  $\sigma_c = \sqrt{2\Lambda_c k_B T}$  accounts for a thermal broadening of  $\rho_c$  and can be associated with a thermalization of the initial state with respect to the vibrational states of the classical modes.

## 2.3 Density functional theory

As indicated earlier, the CT character of the decaying state is an essential aspect of the theoretical description. The difficulties in describing CT states with density functionals is generally related to the many-electron self-interaction error (ME-ESIE), which implies an over-stabilization of fractionally charged subsystems due to semi-local exchange.<sup>47–51</sup> Although the admixture of eX into the exchange–correlation functional can presumably cure the ME-ESIE, global hybrid functionals (constant amount of eX) reportedly fail in describing CT states.<sup>52–55</sup>

A general remedy can be found in the (three-parameter) Coulomb-attenuating method (CAM) approach,<sup>56,57</sup> which introduces a distant-dependent admixture of eX and a range-separation of the exchange interactions between electrons.<sup>58</sup>

The Coulomb operator,  $\frac{1}{r}$ , is divided into a short-range and a long-range part,

$$\frac{1}{r} = \frac{1 - \alpha - \beta \text{erf}(\gamma r)}{r} + \frac{\alpha + \beta \text{erf}(\gamma r)}{r}. \quad (13)$$

Exchange contributions from the first term on the r.h.s. are evaluated using a semi-local exchange functional, while



contributions from the second term are obtained by means of eX. The CAM parameters  $\alpha$  and  $\beta$  measure the (additional) amount of eX in the short-range and the long-range limit, respectively. We note in passing that the value of  $\alpha$  is typically chosen in accordance with global hybrids ( $\alpha \approx 20\%$ ). The range-separation parameter  $\gamma$  corresponds to the inverse of the distance distinguishing the short-range from the long-range regime and allows adjustment of the balance between eX and semi-local exchange.<sup>59</sup>

We recall that the CT states are treated within TD-DFT, and analysis of the corresponding formalism provides a detail essential to our study.<sup>60</sup> Under the assumption of zero overlap between the donating orbital  $\psi_d$  and the accepting orbital  $\psi_a$  of the CT excitation, the only non-vanishing element of the electron-hole interaction matrices is given by<sup>61</sup>

$$A_{da} = \varepsilon_a - \varepsilon_d - \iint d\mathbf{r}_1 d\mathbf{r}_2 C_{eX}(r) \psi_d(\mathbf{r}_1) \frac{1}{r} \psi_a(\mathbf{r}_2). \quad (14)$$

The orbital energies of  $\psi_d$  and  $\psi_a$  are denoted by  $\varepsilon_d$  and  $\varepsilon_a$ , and  $C_{eX}(r)$  is the coefficient for eX, which generally depends on the distance  $r = |\mathbf{r}_1 - \mathbf{r}_2|$  between the position vectors  $\mathbf{r}_1$  and  $\mathbf{r}_2$ . It follows that eX weights the electrostatic attraction between the hole and the particle of the CT excitation, as expressed by the Coulomb integral on the r.h.s. of eqn (14). Discarding the Coulomb integral removes the attraction between hole and particle, allowing large charge separations without energetic penalty, consistent with the tendency toward fractional charges arising from the ME-ESIE.

## 2.4 Continuum solvation

Solvent effects can be conveniently described by continuum solvation models, which assume the solute molecule to be embedded in a cavity within the bulk dielectric of the solvent.<sup>62,63</sup> In the widely adopted apparent surface charge (ASC) method, the solute-solvent interaction is modeled as the interaction between the solute's electrostatic potential and the reaction potential of the bulk dielectric, represented by effective charges on the cavity surface.<sup>62,64,65</sup>

A vertical electronic excitation corresponds to a non-equilibrium situation, in which the electronic degrees of freedom of the solvent relax to the solute's excited-state electrostatic potential, while the nuclear degrees of freedom remain equilibrated with the ground-state potential. Within the ASC method, the non-equilibrium situation is treated by splitting the ASC operator  $\mathbf{P}$  into an electronic (fast) component  $\mathbf{P}_e$  and a nuclear (slow) component  $\mathbf{P}_n$ ,<sup>62</sup>

$$\mathbf{P} = \mathbf{P}_e + \mathbf{P}_n, \quad (15)$$

and evaluating the contributions from  $\mathbf{P}_e$  and  $\mathbf{P}_n$  accordingly. We note that  $\mathbf{P}_e$  is related to the optical dielectric constant of the solvent, whereas  $\mathbf{P}_n$  is connected to its static dielectric constant.

Importantly, the commonly used linear-response (LR) formalism is not admissible for vertical CT excitations with significant change in the solute's electrostatic potential and in such cases the state-specific (SS) formalism should be employed

instead.<sup>62,66,67</sup> By comparative analysis of the two formalisms, approximate expressions can be derived for the vertical excitation energies  $E_{ve}^{(lr)}$  and  $E_{ve}^{(ss)}$  within the LR and SS formalism,<sup>68,69</sup>

$$E_{ve}^{(lr)} = E_{ve}^{(g)} - g_e^{(d)} \mu_{ct-gs}^2, \quad (16)$$

$$E_{ve}^{(ss)} = E_{ve}^{(g)} - \frac{1}{2} g_e^{(s)} (\mu_{ct}^2 - \mu_{gs}^2). \quad (17)$$

The vertical excitation energy in the gas phase is denoted as  $E_{ve}^{(g)}$ ,  $g_e^{(d)}$  and  $g_e^{(s)}$  are the dynamic (d) and static (s) electronic response functions, respectively,  $\mu_{ct-gs}$  is the transition dipole moment of the CT excitation, and  $\mu_{ct}$  and  $\mu_{gs}$  are the corresponding permanent dipole moments. It becomes evident that CT excitation energies within the LR formalism can be severely overestimated, since  $E_{ve}^{(lr)}$  is corrected in terms of the transition dipole moment,  $\mu_{ct-gs}$ , which vanishes for pure CT excitations. We note that while the SS formalism is suited for assessing CT excitation energies in solution, it does not allow for the computation of transition properties, such as  $\mu_{ct-gs}$ .<sup>70</sup>

## 2.5 Radiative decay

For completeness, we also consider the rate constant  $k_r$  for radiative decay by fluorescence. The widely known Einstein eq.,

$$k_r = \frac{(\Delta E)^3 f_{loc} \mu_{ct-gs}^2}{3\epsilon_0 \pi \hbar^4 c^3} \quad (18)$$

provides a simple expression for the rate constant, where  $f_{loc}$  is a correction factor accounting for the effect of the local electric field.<sup>71,72</sup> We determine the transition dipole moment  $\mu_{ct-gs}$  on the basis of a continuum solvation model in the LR formalism, effectively including the local electric field, and set  $f_{loc} = 1$ .

# 3 Computational details

To reduce computational cost, we assume that neither the gradient and curvature of the potential energy surfaces (PESs) nor the non-adiabatic coupling between electronic states of the same spin multiplicity are influenced by solvation effects. Consideration of solvation effects therefore only introduces a (constant) global shift of the PESs relative to one another.

## 3.1 Density functional

We used the CAM-Becke, 3-parameter, Lee-Yang-Parr<sup>56,73-76</sup> (CAM-B3LYP) exchange-correlation functional throughout our study and varied the range-separation parameter  $\gamma$  to adjust the balance between eX and semi-local exchange. The default values  $\alpha = 19\%$  and  $\beta = 46\%$  were adopted for the other two CAM parameters.<sup>56</sup> The default value of  $\gamma$  corresponds to  $\gamma = 0.33a_0^{-1}$ .<sup>56,57</sup>

## 3.2 Geometries and frequencies

Geometry optimizations and Hessian calculations were performed with a def2-SVP basis set<sup>77</sup> for the gas phase at each particular value of  $\gamma$ . No dispersion corrections were added, as changing functional parameters generally requires a re-adjustment of the dispersion coefficients.<sup>78</sup> The resolution-of-identity (RI)



approximation was employed throughout.<sup>79</sup> Geometry optimizations of the first excited electronic state (CT state) were carried out at the corresponding TD-DFT level. The calculations were conducted with version 7.8.1 of the Turbomole program package<sup>80</sup> using the m3 grid for numerical integration.<sup>81,82</sup>

### 3.3 Excitation energies

Vertical excitation energies in solution were determined from single-point energy calculations at the respective ground-state equilibrium geometries using the corresponding TD-DFT level with the polarizable continuum model (PCM)<sup>63</sup> and the solvent parameters of cyclohexane. Three different variants of PCM were considered: linear-response (PCM-LR),<sup>83</sup> corrected linear-response<sup>84</sup> (PCM-cLR) and state-specific<sup>85,86</sup> (PCM-SS). We note that PCM-SS establishes self-consistency between the solute's excited-state electrostatic potential and the solvent's reaction field, whereas PCM-cLR applies first-order perturbation theory. The calculations were performed with version Rev. C.01 of the Gaussian 16 software<sup>87</sup> using a Lebedev integration grid with 45 radial and 434 angular grid points.<sup>88</sup> The response-property-optimized def2-SVPD basis set<sup>89</sup> was employed throughout and imported into Gaussian 16 *via* the basis set exchange library.<sup>90</sup>

### 3.4 Excitonic and non-adiabatic couplings

To access vibronic couplings, we evaluated the electronic gradient **G** of the first excited (CT) state and the derivative coupling vector **D** at the electronic ground state equilibrium geometry.<sup>91</sup> In addition, **D** was also evaluated at the CT state equilibrium geometry.<sup>91</sup> The calculations were performed at the corresponding TD-DFT level using a def2-SVP basis set<sup>77</sup> and the RI approximation.<sup>79</sup> Relative mode displacements  $\Delta_\mu$ , Huang–Rhys factors  $S_\mu$  and non-adiabatic coupling vectors **C** were obtained from a normal coordinate transformation of **G** and **D**, respectively, using the corresponding electronic ground-state Hessian, see Section S2.2 of the SI. Correctly determining the directions of  $\Delta_\mu$  requires properly orienting the Hessian eigenvectors, as outlined in Section S2.1 of the SI. The calculations were carried out with version 7.8.1 of the Turbomole program package<sup>80</sup> using the m3 integration grid.<sup>81,82</sup>

### 3.5 Density of states

The total FCWD,  $\rho_{\text{cha}}$ , was computed with a recursive algorithm for direct state counting, see Section S2.3 of the SI for details.<sup>93,94</sup> Modes with frequencies up to  $300 \text{ cm}^{-1}$  were treated classically, while the harmonic approximation was applied to modes up to  $400 \text{ cm}^{-1}$ . Modes with higher frequencies were modeled as displaced Morse oscillators (*cf.* Section 2.1). The energy gap  $\Delta E$  of the non-radiative transition was determined by subtracting the total reorganization energy  $A$  for displaced harmonic oscillators (in the gas phase) from the vertical excitation energy  $E_{\text{ve}}$  in solution,

$$\Delta E = E_{\text{ve}} - A. \quad (19)$$

Note that we choose the PCM-SS excitation energy  $E_{\text{ve}}^{(\text{ss})}$  for evaluating the total FCWD  $\rho_{\text{cha}}$ . The average  $\langle \rho_{\text{cha}} \rangle_{(\Delta E)}$  of

$\rho_{\text{cha}}$ —which omits local fluctuations of the FCWD around  $\Delta E$ —was obtained using a normal distribution  $g_{(\Delta E)}$  centered at  $\Delta E$ ,

$$\langle \rho_{\text{cha}} \rangle_{(\Delta E)} = \int dE' \rho_{\text{cha}}(E') g_{(\Delta E)}(E'; \sigma_g), \quad (20)$$

$$g_{(\Delta E)}(E'; \sigma_g) = \frac{1}{\sigma_g \sqrt{2\pi}} \exp \left[ -\frac{1}{2} \left( \frac{E' - \Delta E}{\sigma_g} \right)^2 \right]. \quad (21)$$

The width was set to  $\sigma_g = 100 \text{ cm}^{-1}$  and the stability analysis of  $\langle \rho_{\text{cha}} \rangle_{(\Delta E)}$  in Section S2.4 of the SI shows that the results are practically not affected by the averaging.

## 4 Results and discussion

We consider in detail non-radiative decay and vibronic couplings in the three donor-functionalized radical emitters illustrated in Fig. 1. In Sections 4.1–4.3 and 4.6, we examine the relevance of various theoretical aspects to the mode-resolved modeling, *viz.*, balancing eX and semi-local exchange, solvation effects, and anharmonic contributions. This is supplemented by a model calibration in Section 4.4 and an illustrative application linking non-radiative decay to molecular design in Sections 4.5, 4.7 and 4.8.

Non-radiative decay in TTM-TPA and TBTM-TPA was previously characterized using a semi-quantitative approach.<sup>12</sup> Experimentally, the additional chlorination in TBTM-TPA was found to suppress non-radiative decay relative to TTM-TPA, which can be attributed to the perpendicular orientation between the donor and acceptor planes.<sup>12</sup> In the present study, the suppressed decay is explicated by a quantitative, mode-resolved analysis, demonstrating a reasonable overall agreement with experiment.

### 4.1 Long-range exact exchange

First, we recall the effects of the ME-ESIE on the CT properties in the gas phase, as they play a key role to understand the corresponding effects on (continuum) solvation and vibronic structure. Fig. 2a displays the trend in the CT properties of TTM-1Cz with respect to  $\gamma$  (separation of short-range and long-range eX) and supporting results for TTM-TPA and TBTM-TPA are provided in Section S3.1 of the SI. As to be expected, frontier

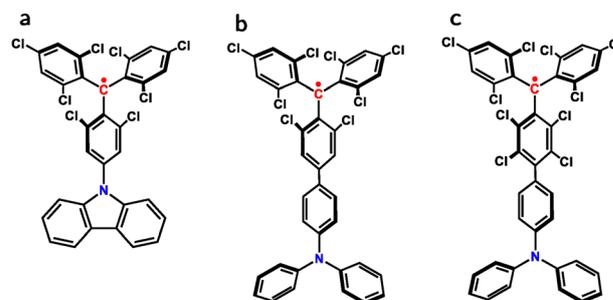


Fig. 1 Chemical structural formulas of TTM-1Cz (a), TTM-TPA (b) and TBTM-TPA (c). The radical centers are highlighted in red.



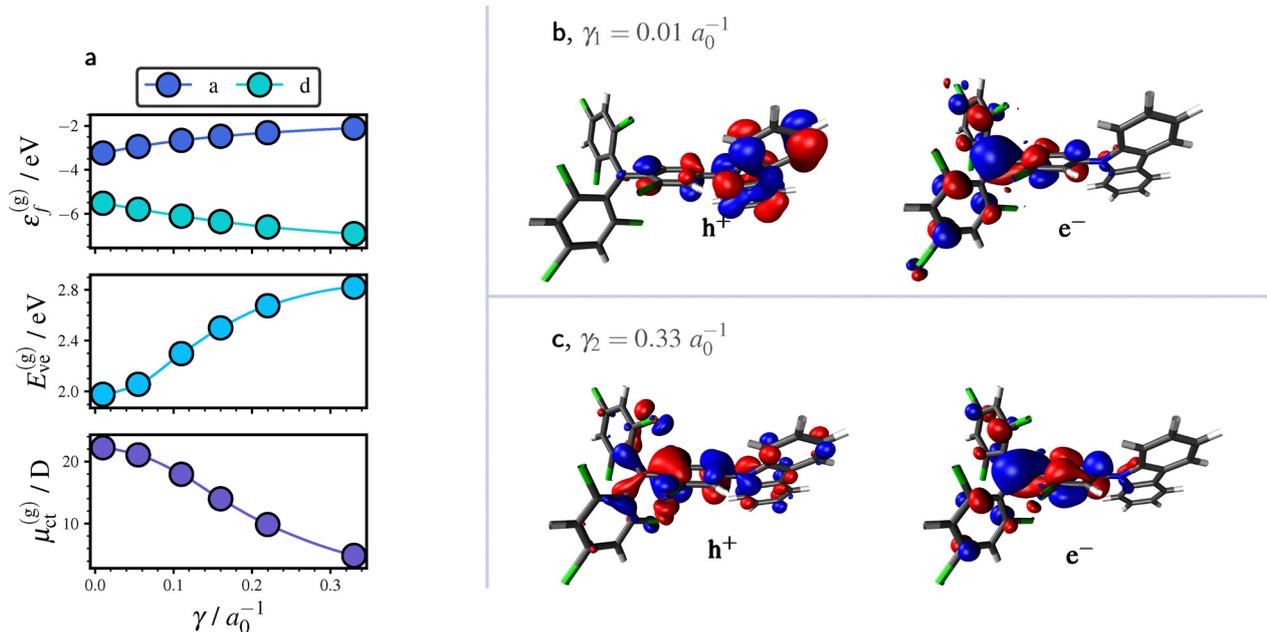


Fig. 2 CT properties of TTM-1Cz in the gas phase (g) in dependence on the range-separation parameter  $\gamma$ . (a) Frontier orbital energy  $\epsilon_f^{(g)}$  (acceptor: a, donor: d), vertical excitation energies  $E_{ve}^{(g)}$  and CT state dipole moment  $\mu_{ct}^{(g)}$ . (b) and (c), Hole ( $h^+$ ) and particle ( $e^-$ ) of the CT exciton obtained by a natural transition orbital analysis<sup>92</sup> at  $\gamma_1 = 0.01a_0^{-1}$  (contrib.: 99%) and  $\gamma_2 = 0.33a_0^{-1}$  (contrib.: 62%). All properties were evaluated at the respective ground state equilibrium geometries.

orbital energy gap and vertical excitation energy  $E_{ve}^{(g)}$  increase with  $\gamma$ . For the CT state dipole moment,  $\mu_{ct}^{(g)}$ , we observe a notable decline with practically no CT at large  $\gamma$  values.

As can be seen from Fig. 2c, hole and particle are in close proximity at  $\gamma_2 = 0.33a_0^{-1}$ , which can be comprehended in terms of an increased electrostatic attraction due to long-range eX, cf. eqn (14).<sup>61</sup> Accordingly, we observe at  $\gamma_1 = 0.01a_0^{-1}$  a confinement of hole and particle to donor and acceptor unit, respectively, as shown Fig. 2b.

The radical emitters therefore clearly demonstrate that long-range eX governs the separation between hole and particle, as reflected in the magnitude of  $\mu_{ct}^{(g)}$ .<sup>61</sup> The trend in  $\mu_{ct}^{(g)}$  is relevant for solvation effects, and the localization of hole and particle is also expected to have important consequences for excitonic couplings.

#### 4.2 Solvation effects

Large dipole moments  $\mu_{ct}^{(g)}$  lead to mutual polarization of solvent and solute and result in a considerable decrease of the CT excitation energy  $E_{ve}$  in solution. In Fig. 3, we compare the performance of three different versions of PCM, viz. PCM-LR, PCM-cLR and PCM-SS (see also Section 3.3). Note that PCM-cLR yields the same dipole moment  $\mu_{ct}$  as PCM-LR, because it applies first-order perturbation theory to PCM-SS and thus leaves the wavefunction unperturbed. For TBTM-TPA (Fig. 3c) we observe significant solvation effects, which can only be captured with PCM-SS. This can be attributed to enhanced particle-hole separation caused by the perpendicular orientation of donor and acceptor planes, along with stronger solute-solvent polarization resulting from an increased CT-state dipole moment. The results

for the other systems, TTM-1Cz and TTM-TPA, likewise support the need for state-specific continuum solvation, although the solvation effects are less pronounced. We note that abandoning the state-specific formalism in combination with small long-range eX can potentially lead to error cancellation. In this scenario, excitonic couplings are particularly likely to be underestimated, yielding misleading mode contributions to non-radiative decay, see also Section 4.3.

Concerning TBTM-TPA, evaluating excited-state electronic gradients and derivative coupling vectors in the gas phase may introduce considerable errors due to significant solvation effects. In Section S3.2 of the SI, a stability analysis demonstrates that the dominant contribution is the excitation energy, which is properly recovered in our framework. In addition, trends in key quantities, as discussed in Section S3.4 of the SI, indicate that solvation likely induces minor changes in the excitonic couplings and is therefore of secondary importance.

#### 4.3 Vibronic couplings and range-separation

We now examine the relationship between vibronic couplings in the radical emitters and the separation of short-range and long-range eX. We first present a stability analysis of the total rate constant  $k_{nr}^{(cha)}$  with respect to  $\gamma$ , followed by an assessment of the excitonic couplings. TTM-1Cz is used as illustrative example and supporting results for TTM-TPA and TBTM-TPA are provided in Section S3.2 of the SI.

The two main contributions to  $k_{nr}^{(cha)}$  are given by the non-adiabatic coupling  $|C|$  and the averaged total FCWD  $\langle \rho_{cha} \rangle_{(\Delta E)}$ , cf. eqn (3). As exemplified by eqn (7), the average  $\langle \rho_{cha} \rangle_{(\Delta E)}$  consists of two principal contributions, the excitonic couplings



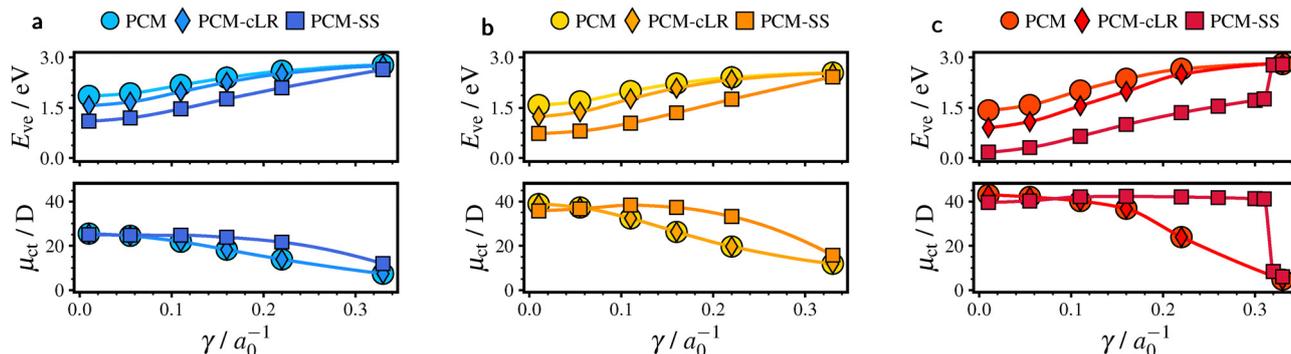


Fig. 3 Vertical excitation energies  $E_{ve}$  and CT state dipole moments  $\mu_{ct}$  in dependence on the range-separation parameter  $\gamma$  obtained from different versions of PCM: linear response (PCM-LR), corrected linear response (PCM-cLR) and state-specific (PCM-SS). Results are displayed for TTM-1Cz (a), TTM-TPA (b) and TBTM-TPA (c).

$S_\mu$  and the energy gap  $\Delta E$ . Fig. 4 illustrates the trend of the average  $\langle \rho_{cha} \rangle_{(\Delta E)}$  in relation to the energy gap  $\Delta E$  and the excitonic couplings, together with a comparison to the averaged FCWD  $\langle \rho_{ch} \rangle_{(\Delta E)}$ , where all high-frequency modes are treated harmonically.

In Fig. 4a, we observe a clear exponential decay of  $\langle \rho_{ch} \rangle_{(\Delta E)}$  and  $\langle \rho_{cha} \rangle_{(\Delta E)}$  with the gap  $\Delta E$ . Even small variations in  $\Delta E$  lead to changes by several orders of magnitude. This exponential behavior is consistent with the energy-gap law for non-radiative decay, since the corresponding rate constant  $k_{nr}$  differs from the FCWD average only by a constant factor (if non-adiabatic couplings are held fixed). Furthermore, the deviation between a harmonic and an anharmonic treatment of the high-frequency modes increases with  $\Delta E$  and becomes significant at physically relevant gaps of about 2 eV (see also Section 4.6).

Considering the influence of excitonic couplings in Fig. 4b,  $\langle \rho_{cha} \rangle_{(\Delta E)}$  is shifted relative to  $\langle \rho_{ch} \rangle_{(\Delta E)}$  by approximately a constant offset. In both cases, the FCWD average increases markedly with  $\gamma$  and appears to approach a saturation limit at large  $\gamma$ .

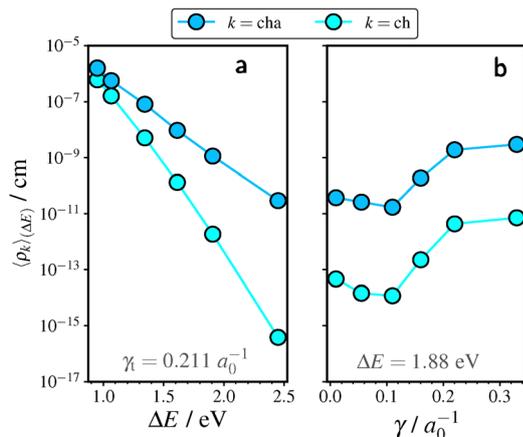


Fig. 4 Stability analysis of the FCWD averages  $\langle \rho_k \rangle_{(\Delta E)}$  for TTM-1Cz. (a) Dependence on the energy gap  $\Delta E$  with excitonic couplings fixed at range-separation parameter  $\gamma_i$ . (b) Dependence on the excitonic couplings evaluated at various  $\gamma$  with constant gap  $\Delta E$ .

The combined effect of  $\Delta E$  and excitonic couplings on the averages  $\langle \rho_k \rangle_{(\Delta E)}$  as a function of  $\gamma$  is illustrated in Fig. 5a. We note that the gaps  $\Delta E$  were computed from the corresponding PCM-SS excitation energies, cf. Fig. 3a. The overall trend is dominated by the gap  $\Delta E$ , leading to a decrease of  $\langle \rho_k \rangle_{(\Delta E)}$  by several orders of magnitude with increasing  $\gamma$ . The non-adiabatic coupling shown in Fig. 5b increases with  $\gamma$ , although it remains within the same order of magnitude. Consequently, the overall trend in  $k_{nr}^{(k)}$  with  $\gamma$  in Fig. 3c can be understood as determined by  $\langle \rho_{cha} \rangle_{(\Delta E)}$  and hence dominated by the gap  $\Delta E$ . We note that the stability analyses of TTM-TPA and TBTM-TPA in Section S3.2 of the SI lead to the same conclusions.

As pointed out in Section 4.7, non-radiative decay in the radical emitters is promoted by a single high-frequency mode owing to its significant contribution to  $\langle \rho_{cha} \rangle_{(\Delta E)}$ . To clarify the

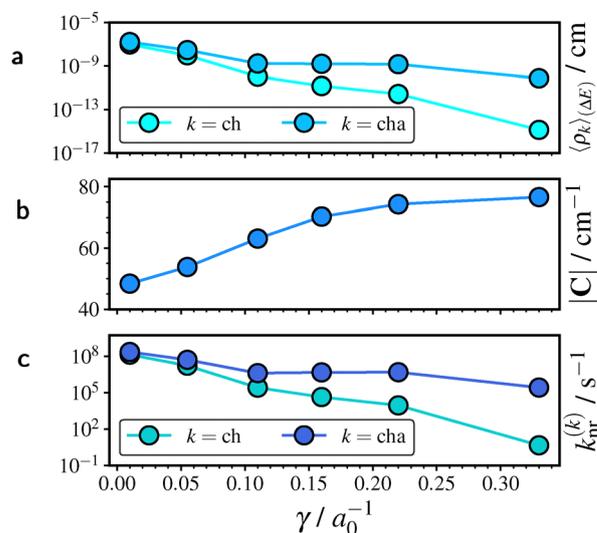


Fig. 5 Stability analysis of the rate constant  $k_{nr}^{(k)}$  of TTM-1Cz with respect to range-separation parameter  $\gamma$ . (a) Overall dependence of the averages  $\langle \rho_k \rangle_{(\Delta E)}$  on  $\gamma$ . Energy gaps  $\Delta E$  were evaluated using the vertical excitation energies at the PCM-SS level. (b) Dependence of the non-adiabatic coupling  $|C|$  on  $\gamma$ , evaluated at the electronic ground-state equilibrium geometry. (c) Dependence of  $k_{nr}^{(k)}$  on  $\gamma$ .



relationship between long-range eX and excitonic coupling, we examine the mode displacement  $\Delta_\lambda$  and HR factor  $S_\lambda$  (coupling strength) for the promoting mode  $\lambda$  of TTM-1Cz, as shown in Fig. 6. Supporting results for TTM-TPA and TBTM-TPA are provided in Section S3.3 of the SI.

The promoting mode of TTM-1Cz at approximately  $1650\text{ cm}^{-1}$  is localized on the acceptor moiety in close proximity to the donor unit (see Fig. S9a of the SI). As discussed in detail in Section S3.4 of the SI, the promoting mode perturbs hole and particle simultaneously more readily at larger  $\gamma$ , owing to their increased spatial adjacency resulting from long-range eX. In Fig. 6, we therefore observe a pronounced increase in the positive displacement  $\Delta_\lambda$  and in  $S_\lambda$  with increasing  $\gamma$ , both approaching a saturation limit at large  $\gamma$ . This trend is mirrored by that of  $\langle\rho_k\rangle_{(\Delta E)}$  in Fig. 4b.

Indeed, the results in Section 4.7 show that, with increasing  $\gamma$ , non-radiative decay in the radical emitters is governed almost exclusively by the positively displaced promoting mode. In combination with the trends in the PCM-SS excitation energy shown in Fig. 3, these findings suggest that an accurate description of the CT state energy and the vibronic structure of donor-functionalized radical emitters requires a considerable amount of long-range eX. These findings are supported by the data presented in Section 3.5 of the SI, showing that the observed trends extend to other exchange–correlation functionals.<sup>56,95–97</sup>

#### 4.4 Model calibration

In Sections 3.1–3.3, we demonstrated that  $\gamma$  significantly influences various properties of radical emitters relevant to the non-radiative decay of the CT state and is hence the central parameter of our model. In the present study, an empirical tuning of  $\gamma$  is performed by requiring that the PCM-SS vertical excitation energy  $E_{\text{ve}}^{(\text{ss})}$ , cf. Section 3.3, reproduce the experimental absorption maximum. To this end,  $E_{\text{ve}}^{(\text{ss})}$  is interpolated as a function of  $\gamma$ , and the tuned values  $\gamma_t$  are obtained by inverting the interpolant and supporting details and an extended discussion are provided in Section S3.6 of the SI.<sup>56,57,73–76,95–99</sup> Using  $\gamma_t$ , equilibrium geometries, Hessians, excited-state electronic gradients, and derivative coupling vectors are evaluated for the

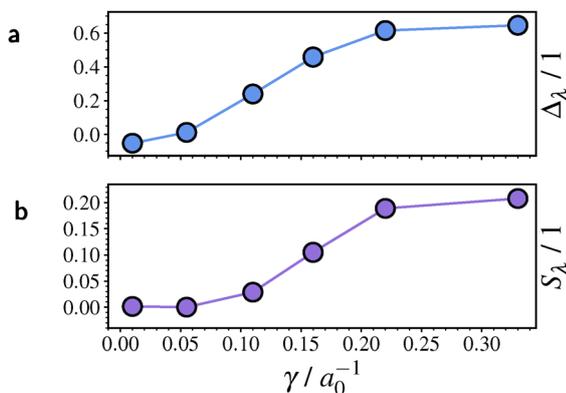


Fig. 6 Dependence of the signed mode displacement  $\Delta_\lambda$  (a) and HR factor  $S_\lambda$  (b) of the promoting mode  $\lambda$  in TTM-1Cz on the range-separation parameter  $\gamma$ .

Table 1 Comparison of the experimental (exp) ref. 12 and 100 with theoretical results: vertical excitation energies  $E_{\text{ve}}$ , calculated energy gaps  $\Delta E$  (cf. eqn (19)) and radiative and non-radiative rate constants  $k_r$  and  $k_{\text{nr}}$  calculated using the tuned range-separation parameters  $\gamma_t$ .  $E_{\text{abs}}^{(\text{exp})}$  and  $E_{\text{ems}}^{(\text{exp})}$  refer to the maximum of absorption and emission, respectively

	TTM-1Cz	TTM-TPA	TBTM-TPA
$\gamma_t/a_0^{-1}$	0.211	0.251	0.310
$E_{\text{abs}}^{(\text{exp})}/\text{eV}$	2.06	1.95	1.97
$E_{\text{ve}}^{(\text{ss})}/\text{eV}$	2.06	1.95	1.77
$E_{\text{ems}}^{(\text{exp})}/\text{eV}$	1.97	1.70	1.70
$\Delta E/\text{eV}$	1.88	1.66	1.57
$k_r^{(\text{exp})}/10^7\text{ s}^{-1}$	1.28	1.88	2.51
$k_r/10^7\text{ s}^{-1}$	1.12	1.19	1.30
$k_{\text{nr}}^{(\text{exp})}/\text{s}^{-1}$	$1.1 \times 10^7$	$1.4 \times 10^8$	$3.3 \times 10^7$
$k_{\text{nr}}^{(\text{cha})}/\text{s}^{-1}$	$1.0 \times 10^7$	$6.8 \times 10^8$	$1.4 \times 10^7$
$k_{\text{nr}}^{(\text{ch})}/\text{s}^{-1}$	$2.1 \times 10^4$	$2.1 \times 10^7$	$1.3 \times 10^6$

gas phase as described in Section 3. For TBTM-TPA, only an approximate tuning of  $\gamma$  was feasible due to computational difficulties arising from its perpendicular arrangement (see also Section S3.6 of the SI).

Table 1 compiles relevant properties obtained with the calibrated models together with experimental reference values. While  $\gamma$  was adjusted *a posteriori* to match the excitation energies, the model also shows good agreement with other experimental properties. In particular, the experimental trend in the non-radiative decay rates is reproduced correctly by the  $k_{\text{nr}}^{(\text{cha})}$  values. This indicates that the calibrated model provides a reasonable description of the systems. An empirical tuning of  $\gamma$  enables a detailed analysis of the experimental results and is employed here for convenience. Trend analyses of the FCWD (Fig. 4) and, in particular, the coupling between the promoting mode and the CT exciton (Fig. 6) reveal a consistent trend with respect  $\gamma$  suggesting that a non-empirical description could be feasible. Accordingly, alongside a non-empirical tuning of  $\gamma$ , comparative trend analyses of relevant excitonic couplings across different systems may suffice to predict trends in non-radiative decay.

Anharmonic contributions are necessary to reach quantitative agreement with experiment, cf. Table 1. Determining the non-adiabatic coupling  $|C|$  at the CT state equilibrium geometry is physically sound, as decay proceeds from this geometry and, according to exploratory calculations (not shown), further improves agreement with experiment.

The assumption of distinct accepting and promoting modes was employed in deriving the quantum-mechanical FCWDs<sup>37,38</sup> and is well fulfilled by the calibrated models, as shown in Section S3.7 of the SI. We also note that structural relaxation upon non-radiative transition is presumably small and can be reasonably described within the displaced harmonic oscillator framework, as supported by the considerations in Section S3.8 of the SI. To account for anharmonic contributions arising from highly excited vibrational states, the displaced Morse oscillator model is required.

#### 4.5 Mode-resolved analysis

An analysis of vibronic couplings can be performed by examining the spectra of the non-adiabatic couplings  $C_{\mu}$ , excitonic



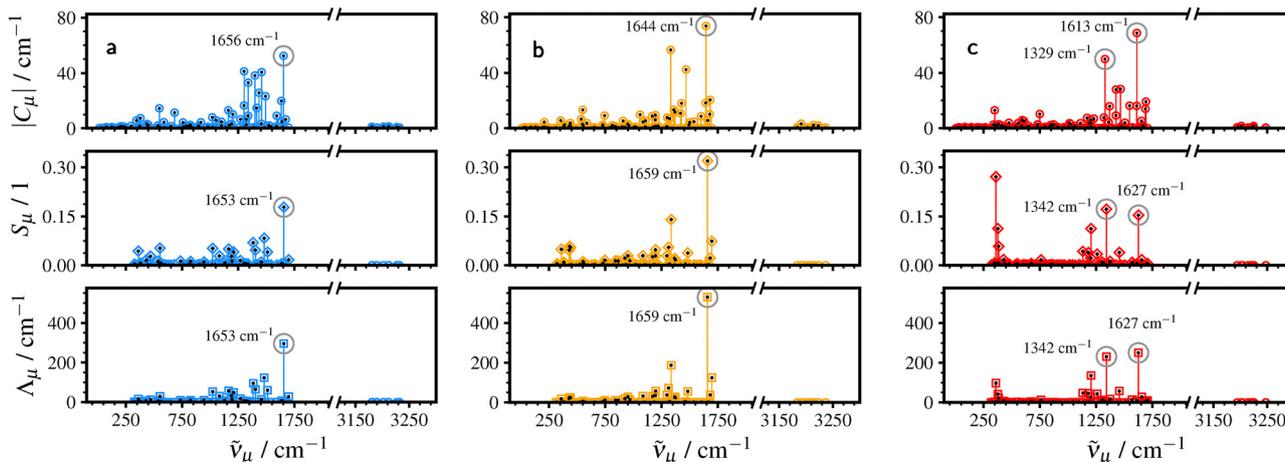


Fig. 7 Mode-resolved analysis of the unsigned non-adiabatic couplings  $|C_\mu|$ , excitonic couplings  $S_\mu$  and contributions  $A_\mu$  to the reorganization energy  $A$  related to a normal mode  $\mu$  with wavenumber  $\tilde{\nu}_\mu$ . Results are displayed for TTM-1Cz (a), TTM-TPA (b) and TBTM-TPA (c). Selected more contributions are highlighted. The required calculations were performed with the tuned range-separation parameters  $\gamma_t$  compiled in Table 1.

couplings  $S_\mu$ , and reorganization energies  $A_\mu$  with respect to the normal modes  $\mu$ , as shown in Fig. 7 (with  $S_\mu$  omitted for classical modes). The couplings  $C_\mu$  were evaluated at the CT equilibrium geometry, with further details provided in Section S3.9.<sup>101</sup>

The non-radiative transition in the radical emitters is induced by high-frequency skeletal vibrations, with exemplary illustrations provided in Section S3.10 of the SI. For both TTM-1Cz and TTM-TPA (Fig. 7a and b), a single high-frequency mode at around  $1650\text{ cm}^{-1}$  is found that exhibits significant excitonic coupling  $S_\mu$  and contributes substantially to the reorganization energy. For TBTM-TPA in Fig. 7c, we observe a reduced yet still notable excitonic coupling of the mode at  $1627\text{ cm}^{-1}$ , together with a marked coupling at  $1342\text{ cm}^{-1}$ .

A comparative analysis of vibronic couplings indicates that variations in non-radiative decay rates are governed by excitonic couplings  $S_\mu$ , whereas the non-adiabatic couplings  $C_\mu$  are comparable among the emitters studied. In particular, the suppressed mode in TBTM-TPA at  $1627\text{ cm}^{-1}$  is analogous to the strongly coupled modes of TTM-1Cz and TTM-TPA at approximately  $1650\text{ cm}^{-1}$  (see Section S3.4 of the SI for illustrations). The suppression of non-radiative decay can thus be attributed to the additional chlorination of the TTM moiety. While inspection of the spectra for  $S_\mu$  already provides insight into the experimental trend of  $k_{\text{nr}}$ , a more refined analysis can be obtained by accounting for anharmonic contributions, as discussed in Section 4.6.

A notable feature in the spectra of Fig. 7 is the absence of both large  $C_\mu$  and significant  $S_\mu$  for proton stretching vibrations (above  $3000\text{ cm}^{-1}$ ). This contrasts with the common view that non-radiative decay proceeds predominantly *via* such vibrations.<sup>9,28,40,41,102</sup> For the radical emitters considered, proton-stretching vibrations—typically localized at the outer edges of the donor or acceptor—have only minor effects on the CT state energy, giving rise to negligible excitonic couplings and making them essentially irrelevant for non-radiative decay

(see also Section S3.11). Moreover, previous studies on non-radiative decay in metal-organic complexes emphasize the key role of high-frequency skeletal vibrations.<sup>103–107</sup>

#### 4.6 Analysis of anharmonic contributions

The analysis in Section 4.5 is restricted to displaced harmonic oscillators. As shown in Table 1, accounting for anharmonic contributions to  $k_{\text{nr}}$  is necessary to achieve quantitative agreement with experiment. We therefore analyze the anharmonic contributions for TTM-1Cz, with particular focus on the promoting mode at  $1656\text{ cm}^{-1}$ , and provide a comparison with the harmonic approximation. Supporting results for TTM-TPA and TBTM-TPA are provided in Section S3.12 of the SI. Fig. 8a shows that the promoting mode  $\lambda$  possesses a comparatively large displacement  $\Delta_\lambda$  with positive orientation. We recall that both the magnitude and the orientation of  $\Delta_\lambda$  determine the anharmonic FC factor near the energy gap and thus the relevance of a given mode to nonradiative decay, *cf.* Section 2.1.

For a positive displacement  $\Delta_\lambda$ , the vibronic transition terminates near the inner turning point of the final Morse potential (see also Section S1.2 of the SI), resulting in a relatively slow decrease of the anharmonic FC factor  $F_{\lambda,v_\lambda}$  with the vibrational quantum number  $v_\lambda$  as shown in Fig. 8b. By contrast, the anharmonic  $F_{\lambda,v_\lambda}$  for a hypothetical negative displacement  $\Delta_\lambda$ , as well as the harmonic  $F_{\lambda,v_\lambda}$ , are smaller by several orders of magnitude in the relevant range  $v_\lambda \geq 9$ . Consequently, the orientation of promoting mode displacement plays an important role in achieving a quantitative description of non-radiative decay.

Moreover, the increasing deviation of the harmonic  $F_{\lambda,v_\lambda}$  with  $v_\lambda$  underscores the relevance of an anharmonic model for highly excited vibrational states.

The high-frequency mode in TBTM-TPA at  $1612\text{ cm}^{-1}$  not only exhibits a reduced displacement  $\Delta_\lambda$  (weaker excitonic coupling  $S_\lambda$ ) but also a negative orientation of this displacement as shown in Section S3.2 of the SI. Both effects lead to a reduction



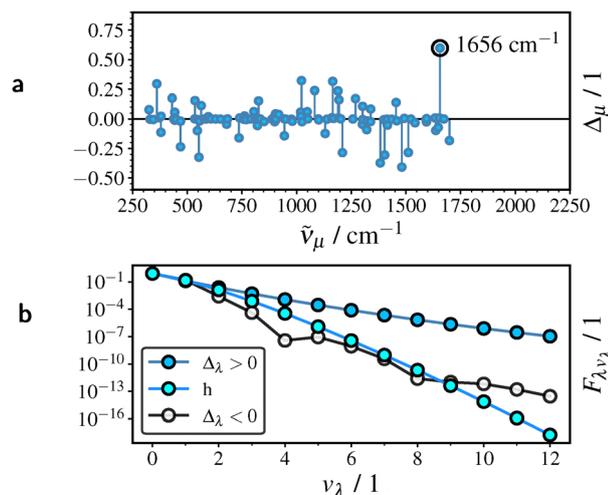


Fig. 8 Anharmonic contributions in TTM-1Cz. (a) Displacement  $\Delta_\mu$  of modes with wavenumbers  $\tilde{\nu}_\mu$ . (b) Dependence of the FC factor  $F_{\lambda,\nu_\lambda}$  of the promoting mode on vibrational quantum number  $\nu_\lambda$ . Anharmonic  $F_{\lambda,\nu_\lambda}$  with  $\Delta_\mu > 0$  and  $\Delta_\mu < 0$ , and harmonic (h)  $F_{\lambda,\nu_\lambda}$  are shown. Calculations were performed with the tuned parameter  $\gamma = 0.211a_0^{-1}$ .

of the (anharmonic) FC factor at relevant vibrational energies. Conversely, the promoting mode in TTM-TPA ( $1659\text{ cm}^{-1}$ ) is positively displaced, leading to faster non-radiative decay. In addition to a reduced excitonic coupling of the promoting mode, non-radiative decay in TBTM-TPA is suppressed due to a

diminished anharmonic contribution ( $\Delta_\lambda < 0$ ), resulting in a reduced vibronic transition probability. Qualitatively, the reduced  $S_\lambda$  value for TBTM-TPA suffices to explain the suppressed non-radiative decay, as the experimental trend is reproduced correctly at the qualitative level within the harmonic approximation, cf. the  $k_{\text{nr}}^{(\text{ch})}$  values in Table 1.

#### 4.7 Identification of the promoting mode

The relevance of a quantum-mechanical mode  $\lambda$  to non-radiative decay can be quantified through a sensitivity analysis of the total FCWD average  $\langle \rho_{\text{cha}} \rangle_{(\Delta E)}$ . We define the enhancement factor  $f_{e,\mu}$  of a mode  $\mu$  as

$$f_{e,\mu} = \frac{\langle \rho_{\text{cha}} \rangle_{(\Delta E)}}{\langle \rho_{\text{cha}}^{(\mu)} \rangle_{(\Delta E)}}, \quad (22)$$

where  $\langle \rho_{\text{cha}}^{(\mu)} \rangle_{(\Delta E)}$  denotes the FCWD average evaluated with mode  $\mu$  omitted. If the mode contributes significantly to non-radiative decay, its omission will markedly reduce the FCWD average and yield a large  $f_{e,\mu}$  value.

The spectra of enhancement factors  $f_{e,\mu}$  in Fig. 9 are dominated by a single promoting mode in each case, and the corresponding nuclear motion is visualized. TTM-1Cz and TTM-TPA share the same promoting mode around  $1650\text{ cm}^{-1}$ , which is suppressed in TBTM-TPA ( $1612\text{ cm}^{-1}$ ) due to additional chlorination. Instead, non-radiative decay in TBTM-TPA proceeds through a less efficient mode at  $1342\text{ cm}^{-1}$ . Thus, sensitivity

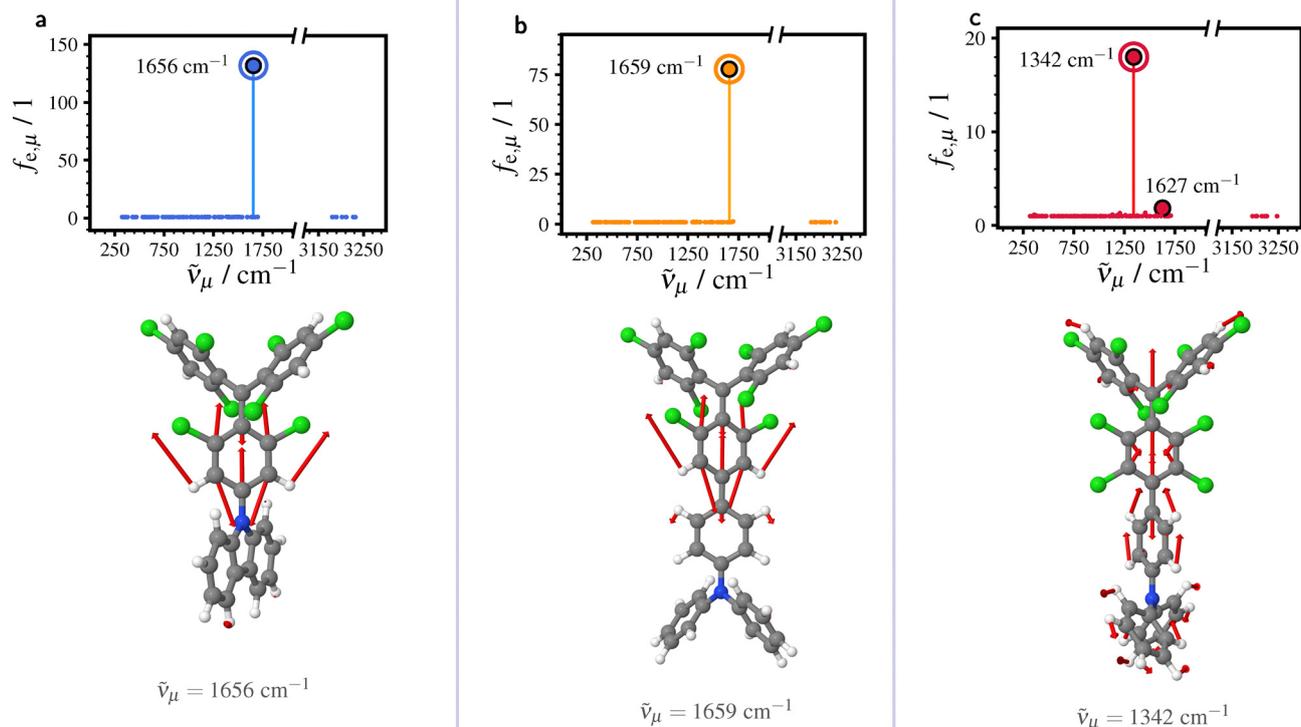


Fig. 9 Enhancement factors  $f_{e,\mu}$  measuring the contribution of a single mode  $\mu$  with wavenumber  $\tilde{\nu}_\mu$  to the total FCWD average  $\langle \rho_{\text{cha}} \rangle$ . Results are displayed for TTM-1Cz (a), TTM-TPA (b) and TBTM-TPA (c). The vibrational motions of the dominant modes are visualized. Calculations were performed with the tuned parameters  $\gamma_t$  compiled in Table 1.



analysis of  $\langle \rho_{\text{cha}} \rangle_{(\Delta E)}$  provides a reliable means of identifying the promoting mode, whereas simply examining the excitonic couplings  $S_{\mu}$  (cf. Fig. 7) may qualitatively agree with the identified promoting modes but does not necessarily make their dominant character sufficiently evident.

As the FCWD measures the joint probability of all allowed vibronic transitions at a given energy gap  $\Delta E$  (see Section 2.1), the results of the present section indicate that there is only a single mode with a significant FC factor near  $\Delta E$ . This large FC factor can be attributed to the large magnitude of the displacement  $\Delta_{\lambda}$  (i.e., a large HR factor), combined with its positive orientation (see Section 4.7). Intuitively, the dominant role of the promoting mode may be understood from its ability to effectively quench the CT exciton, as discussed in Section 3.4 of the SI.

Since non-radiative decay in the radical emitters is dominated by a single mode, a one-effective-mode model is potentially viable. Section S3.13 of the SI shows that the experimental trend in  $k_{\text{nr}}$  can be quantitatively reproduced with an extended MLJ model, treating the promoting mode as the effective mode. However, this requires explicitly accounting for anharmonic contributions.

It is also worth noting that, the sensitivity analysis of  $\langle \rho_{\text{cha}} \rangle_{(\Delta E)}$  can be applied to groups of (quantum-mechanical) modes. For example, excluding all proton stretching vibrations further supports the conclusion that this class of vibrations is insignificant (see Section S3.14 of the SI).

Finally, we note that long-range eX is crucial for correctly identifying the promoting mode. Insufficient long-range eX, or the use of a global hybrid (corresponding to  $\gamma = 0$ ), leads to a fundamentally different non-radiative decay mechanism in which no single mode contributes dominantly, cf. Section S3.15 of the SI.

#### 4.8 Notes on molecular design

A key advantage of the mode-resolved approach is that it enables non-radiative decay to be directly linked to specific structural features through molecular vibrations. Here, we discuss the findings of Sections 4.5–4.7 in the context of molecular design.

Our results suggest that the promoting high-frequency mode in donor-functionalized TTM radicals is located at the phenyl ring adjacent to the donor moiety. This mode is strongly coupled to the CT exciton and further accelerates non-radiative decay *via* anharmonic contributions ( $\Delta_{\lambda} > 0$ ). Together, these effects yield a substantial FC factor near the transition gap, which underlies the dominant role of the promoting mode. In donor-functionalized TBTM radicals, both the excitonic coupling and the anharmonic contribution of this mode are suppressed by the additional chlorination ( $\Delta_{\lambda} < 0$ ), resulting in a diminished FC factor. Non-radiative decay is hence reduced and likely proceeds through a (delocalized) less efficient mode. PTM-based emitters do not exhibit further suppression of non-radiative decay compared to the TBTM analogues,<sup>12,108</sup> as the promoting mode is already attenuated, consistent with our findings. Results for additional emitters in Section S3.16

support extending the explanation of suppressed non-radiative decay from TBTM-TPA to the TBTM and PTM series. This section also includes a comprehensive discussion on possible effects influencing non-radiative decay in the radical emitters.

Although localization of vibrations to either the donor or the acceptor moiety may reduce excitonic couplings to some extent,<sup>11,13</sup> our study demonstrates that a localized high-frequency vibration can still significantly perturb the CT exciton, since hole and particle may retain considerable overlap due to mutual electrostatic attraction. The introduction of bulky substituents near the donor–acceptor junction can further reduce the excitonic coupling and thereby suppress non-radiative decay.

Moreover, unlike in our earlier study,<sup>12</sup> we find that inclusion of contributions from the second, locally excited electronic state is not necessary to achieve a sufficient description of the non-radiative decay.

## 5 Conclusions

Our study presents a mode-resolved model for the non-radiative decay of the CT state in donor-functionalized radical emitters. The decisive role of additional long-range exact exchange (eX) in density functionals for governing this decay is clearly demonstrated. An adequate amount of eX is essential for a consistent description of excitation energies in solution as well as the couplings of relevant modes to the CT exciton. Importantly, eX amplifies the excitonic coupling of the promoting mode, which becomes evident only at sufficiently large eX contributions. Based on our theoretical analysis, it is likely that the role of exact exchange in governing vibronic transitions from charge-transfer (CT) states also extends to other systems, including closed-shell emitters. We emphasize, however, that confirming this hypothesis would require further computational verification. Moreover, since the non-radiative transition in the radical emitters terminates in an excited vibrational state of the promoting mode, explicit inclusion of anharmonic contributions is imperative for quantitative accuracy.

In this study, the model was calibrated through empirical tuning of a functional parameter, with trend analyses suggesting that a non-empirical approach may also be feasible. The calibrated model agrees well with experimental results and accounts for the suppressed decay in the examined radical emitters as a result of the perpendicular orientation of donor and acceptor planes. The sterically demanding substituents required for this arrangement hinder the vibrational motion of the promoting mode, substantially reducing both its coupling to the CT exciton and its anharmonic contribution. In this case, non-radiative decay proceeds *via* an alternative, less efficient mode. The promoting mode can be identified unambiguously by a sensitivity analysis of the averaged Franck–Condon weighted density of states (FCWD), explicitly accounting for anharmonic contributions. Our approach for identifying modes relevant to non-radiative decay through analysis of the quantum-mechanical FCWD is in principle transferable to



other systems, including decay from other types of excited states, e.g., a locally excited state. Such transferability would, in particular, require the Condon approximation to hold and would need further computational verification. We highlight that the key strength of this approach lies in its ability to link non-radiative decay processes to structural features *via* specific vibrational modes.

## Conflicts of interest

There are no conflicts to declare.

## Data availability

The data that support the findings of this study are available upon reasonable request from the corresponding author, RT.

The data supporting this article have been included as part of the supplementary information (SI). Supplementary information: supplementary theoretical and computational details, supplementary results, extended discussion and optimized geometries. See DOI: <https://doi.org/10.1039/d5cp03613e>.

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## Notes and references

- S. R. Forrest and M. E. Thompson, *Chem. Rev.*, 2007, **107**, 923–925.
- M. Xu, X. Li, S. Liu, L. Zhang and W. Xie, *Mater. Chem. Front.*, 2023, **7**, 4744–4767.
- L. Lan and H. Zhang, *Angew. Chem., Int. Ed.*, 2025, **64**, e202509140.
- R. H. Friend, R. Gymer, A. Holmes, J. Burroughes, R. Marks, C. Taliani, D. Bradley, D. D. Santos, J.-L. Bredas and M. Lögdlund, *et al.*, *Nature*, 1999, **397**, 121–128.
- Q. Peng, A. Obolda, M. Zhang and F. Li, *Angew. Chem., Int. Ed.*, 2015, **54**, 7091–7095.
- A. Obolda, X. Ai, M. Zhang and F. Li, *ACS Appl. Mater. Interfaces*, 2016, **8**, 35472–35478.
- I. Ratera, J. Vidal-Gancedo, D. MasPOCH, S. T. Bromley, N. Crivillers and M. Mas-Torrent, *J. Mater. Chem. C*, 2021, **9**, 10610–10623.
- R. Englman and J. Jortner, *Mol. Phys.*, 1970, **18**, 145–164.
- R. Englman and J. Jortner, *J. Lumin.*, 1970, **1**, 134–142.
- X. Ai, E. W. Evans, S. Dong, A. J. Gillett, H. Guo, Y. Chen, T. J. Hele, R. H. Friend and F. Li, *Nature*, 2018, **563**, 536–540.
- P. Ghosh, A. M. Alvertis, R. Chowdhury, P. Murto, A. J. Gillett, S. Dong, A. J. Sneyd, H.-H. Cho, E. W. Evans and B. Monserrat, *et al.*, *Nature*, 2024, **629**, 355–362.
- M. E. Arnold, R. Toews, L. Schneider, J. Schmid, M. H. Putra, M. Busch, A. Groß, F. Deschler, A. Köhn and A. J. Kuehne, *Adv. Opt. Mater.*, 2024, 2500296.
- M. Maiuri, *Nature*, 2024, **629**, 289–290.
- R. A. Marcus, *J. Chem. Phys.*, 1956, **24**, 966–978.
- R. Marcus, *Pure Appl. Chem.*, 1997, **69**, 13–30.
- V. Levich and R. Dogonadze, *Dokl. Akad. Nauk SSSR*, 1959, 123–126.
- V. Levich, *Adv. Electrochem. Electrochem. Eng.*, 1966, **4**, 249–371.
- J. Jortner, *J. Chem. Phys.*, 1976, **64**, 4860–4867.
- H. Abroshan, V. Coropceanu and J.-L. Brédas, *Adv. Funct. Mater.*, 2020, **30**, 2002916.
- E. Cho, V. Coropceanu and J.-L. Brédas, *J. Am. Chem. Soc.*, 2020, **142**, 17782–17786.
- E. Cho, V. Coropceanu and J.-L. Brédas, *J. Mater. Chem. C*, 2021, **9**, 10794–10801.
- H. Abroshan, P. Winget, H. S. Kwak, C. T. Brown and M. D. Halls, *Phys. Chem. Chem. Phys.*, 2022, **24**, 16891–16899.
- P. M. Morse, *Phys. Rev.*, 1929, **34**, 57.
- P. A. M. Dirac, *Proc. R. Soc. London, Ser. A*, 1927, **114**, 243–265.
- E. Fermi, *Nuclear physics: a course given by Enrico Fermi at the University of Chicago*, University of Chicago Press, 1950.
- A. Messiah, *Quantum mechanics*, North-Holland Publishing Company, Amsterdam, 1962, vol. 2.
- M. Born and J. Oppenheimer, *Ann. Phys.*, 1927, **389**, 457–484.
- R. Valiev, B. Merzlikin, R. Nasibullin, A. Kurtzevitch, V. Cherepanov, R. Ramazanov, D. Sundholm and T. Kurtén, *Phys. Chem. Chem. Phys.*, 2023, **25**, 6406–6415.
- G. Herzberg and E. Teller, *Z. Phys. Chem.*, 1933, **21**, 410–446.
- E. Condon, *Phys. Rev.*, 1926, **28**, 1182.
- E. U. Condon, *Phys. Rev.*, 1928, **32**, 858.
- E. Condon, *Phys. Rev.*, 1930, **36**, 1121.
- S. Mukamel, *Principles of nonlinear optical spectroscopy*, Oxford University Press, 1995.
- F. Duschinsky, *Acta Physicochim. URSS*, 1937, **7**, 551–566.
- K. Huang and A. Rhys, *Proc. R. Soc. London, Ser. A*, 1950, **204**, 406–423.
- E. S. Medvedev and V. I. Osherov, *Radiationless transitions in polyatomic molecules*, Springer, 1995, vol. 57.
- L. Shi, X. Xie and A. Troisi, *J. Chem. Phys.*, 2022, **157**, 134106.
- R. P. Fornari, J. Aragón and A. Troisi, *J. Chem. Phys.*, 2015, **142**, 184105.
- J. Franck and E. Dymond, *Trans. Faraday Soc.*, 1926, **21**, 536–542.
- R. Valiev, R. Nasibullin, V. Cherepanov, G. V. Baryshnikov, D. Sundholm, H. Ågren, B. F. Minaev and T. Kurtén, *Phys. Chem. Chem. Phys.*, 2020, **22**, 22314–22323.
- V. Plotnikov, *Int. J. Quantum Chem.*, 1979, **16**, 527–541.
- J. López V, A. Rivera, Y. F. Smirnov and A. Frank, *Int. J. Quantum Chem.*, 2002, **88**, 280–295.
- E. Laguerre, *Bull. Soc. Math.*, 1878, **6**, 72–78.
- E. Whittaker and G. Watson, *A course of modern analysis*, Courier Dover Publications, 2020.



- 45 S. Yeganeh and M. A. Ratner, *J. Chem. Phys.*, 2006, **124**, 044108.
- 46 K. M. Ervin, S. Gronert, S. Barlow, M. K. Gilles, A. G. Harrison, V. M. Bierbaum, C. H. DePuy, W. Lineberger and G. B. Ellison, *J. Am. Chem. Soc.*, 1990, **112**, 5750–5759.
- 47 T. M. Henderson, B. G. Janesko and G. E. Scuseria, *J. Phys. Chem. A*, 2008, **112**, 12530–12542.
- 48 J. P. Perdew, R. G. Parr, M. Levy and J. L. Balduz Jr, *Phys. Rev. Lett.*, 1982, **49**, 1691.
- 49 J. F. Janak, *Phys. Rev. B:Condens. Matter Mater. Phys.*, 1978, **18**, 7165.
- 50 O. A. Vydrov, G. E. Scuseria and J. P. Perdew, *J. Chem. Phys.*, 2007, **126**, 154109.
- 51 A. J. Cohen, P. Mori-Sánchez and W. Yang, *Phys. Rev. B:Condens. Matter Mater. Phys.*, 2008, **77**, 115123.
- 52 A. Dreuw and M. Head-Gordon, *J. Am. Chem. Soc.*, 2004, **126**, 4007–4016.
- 53 A. L. Sobolewski and W. Domcke, *Chem. Phys.*, 2003, **294**, 73–83.
- 54 A. Dreuw, G. R. Fleming and M. Head-Gordon, *J. Phys. Chem. B*, 2003, **107**, 6500–6503.
- 55 D. J. Tozer, R. D. Amos, N. C. Handy, B. O. Roos and L. Serrano-Andres, *Mol. Phys.*, 1999, **97**, 859–868.
- 56 T. Yanai, D. P. Tew and N. C. Handy, *Chem. Phys. Lett.*, 2004, **393**, 51–57.
- 57 Y. Tawada, T. Tsuneda, S. Yanagisawa, T. Yanai and K. Hirao, *J. Chem. Phys.*, 2004, **120**, 8425–8433.
- 58 T. Leininger, H. Stoll, H.-J. Werner and A. Savin, *Chem. Phys. Lett.*, 1997, **275**, 151–160.
- 59 R. Baer, E. Livshits and U. Salzner, *Annu. Rev. Phys. Chem.*, 2010, **61**, 85–109.
- 60 M. E. Casida, *Recent Advances In Density Functional Methods: (Part I)*, World Scientific, 1995, pp. 155–192.
- 61 A. Dreuw, J. L. Weisman and M. Head-Gordon, *J. Chem. Phys.*, 2003, **119**, 2943–2946.
- 62 J. Tomasi and M. Persico, *Chem. Rev.*, 1994, **94**, 2027–2094.
- 63 J. Tomasi, B. Mennucci and R. Cammi, *Chem. Rev.*, 2005, **105**, 2999–3094.
- 64 P. Shaw, *Phys. Rev. A:At., Mol., Opt. Phys.*, 1985, **32**, 2476.
- 65 P. Shaw, *Phys. Rev. A:At., Mol., Opt. Phys.*, 1987, **35**, 2254.
- 66 E. McRae, *J. Phys. Chem.*, 1957, **61**, 562–572.
- 67 N. Mataga and T. Kubota, *Molecular interactions and electronic spectra*, M. Dekker, 1970.
- 68 S. Corni, R. Cammi, B. Mennucci and J. Tomasi, *J. Chem. Phys.*, 2005, **123**, 134512.
- 69 R. Cammi, S. Corni, B. Mennucci and J. Tomasi, *J. Chem. Phys.*, 2005, **122**, 104513.
- 70 D. A. Phan Huu, R. Dhali, C. Pieroni, F. Di Maiolo, C. Sissa, F. Terenziani and A. Painelli, *Phys. Rev. Lett.*, 2020, **124**, 107401.
- 71 D. Toptygin, *J. Fluoresc.*, 2003, **13**, 201–219.
- 72 A. Einstein, *Phys. Z.*, 1917, **18**, 121.
- 73 P. J. Stephens, F. J. Devlin, C. F. Chabalowski and M. J. Frisch, *J. Phys. Chem.*, 1994, **98**, 11623–11627.
- 74 A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648–5652.
- 75 C. Lee, W. Yang and R. G. Parr, *Phys. Rev. B:Condens. Matter Mater. Phys.*, 1988, **37**, 785.
- 76 S. H. Vosko, L. Wilk and M. Nusair, *Can. J. Phys.*, 1980, **58**, 1200–1211.
- 77 F. Weigend and R. Ahlrichs, *Phys. Chem. Chem. Phys.*, 2005, **7**, 3297–3305.
- 78 M. Friede, S. Ehlert, S. Grimme and J.-M. Mewes, *J. Chem. Theory Comput.*, 2023, **19**, 8097–8107.
- 79 A. M. Burow, M. Sierka and F. Mohamed, *J. Chem. Phys.*, 2009, **131**, 214101.
- 80 *TURBOMOLE V7.7 2022*, a development of University of Karlsruhe and Forschungszentrum Karlsruhe GmbH, 1989–2007, TURBOMOLE GmbH, <https://www.turbomole.com>.
- 81 O. Treutler and R. Ahlrichs, *J. Chem. Phys.*, 1995, **102**, 346–354.
- 82 K. Eichkorn, F. Weigend, O. Treutler and R. Ahlrichs, *Theor. Chem. Acc.*, 1997, **97**, 119–124.
- 83 G. Scalmani and M. J. Frisch, *J. Chem. Phys.*, 2010, **132**, 114110.
- 84 M. Caricato, B. Mennucci, J. Tomasi, F. Ingrosso, R. Cammi, S. Corni and G. Scalmani, *J. Chem. Phys.*, 2006, **124**, 124520.
- 85 R. Improta, V. Barone, G. Scalmani and M. J. Frisch, *J. Chem. Phys.*, 2006, **125**, 054103.
- 86 R. Improta, G. Scalmani, M. J. Frisch and V. Barone, *J. Chem. Phys.*, 2007, **127**, 074504.
- 87 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. V. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery, Jr., J. E. Peralta, F. Ogliaro, M. J. Bearpark, J. J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman and D. J. Fox, *Gaussian ~ 16 Revision C.01*, Gaussian Inc., Wallingford CT, 2016.
- 88 V. Lebedev, *Theory of Cubature Formulas and Computational Mathematics*, 1980.
- 89 D. Rappoport and F. Furche, *J. Chem. Phys.*, 2010, **133**, 134105.
- 90 B. P. Pritchard, D. Altarawy, B. Didier, T. D. Gibson and T. L. Windus, *J. Chem. Inf. Model.*, 2019, **59**, 4814–4820.
- 91 F. Furche and R. Ahlrichs, *J. Chem. Phys.*, 2002, **117**, 7433–7447.
- 92 R. L. Martin, *J. Chem. Phys.*, 2003, **118**, 4775–4777.
- 93 S. E. Stein and B. Rabinovitch, *J. Chem. Phys.*, 1973, **58**, 2438–2445.
- 94 T. Beyer and D. Swinehart, *Commun. ACM*, 1973, **16**, 379.



- 95 M. A. Rohrdanz, K. M. Martins and J. M. Herbert, *J. Chem. Phys.*, 2009, **130**, 054112.
- 96 T. M. Henderson, B. G. Janesko and G. E. Scuseria, *J. Chem. Phys.*, 2008, **128**, 194105.
- 97 J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1996, **77**, 3865.
- 98 F. N. Fritsch and J. Butland, *SIAM J. Sci. Stat. Comput.*, 1984, **5**, 300–304.
- 99 H. Iikura, T. Tsuneda, T. Yanai and K. Hirao, *J. Chem. Phys.*, 2001, **115**, 3540–3544.
- 100 A. Abdurahman, T. J. Hele, Q. Gu, J. Zhang, Q. Peng, M. Zhang, R. H. Friend, F. Li and E. W. Evans, *Nat. Mater.*, 2020, **19**, 1224–1229.
- 101 J. R. Reimers, *J. Chem. Phys.*, 2001, **115**, 9103–9109.
- 102 R. R. Valiev, V. N. Cherepanov, R. T. Nasibullin, D. Sundholm and T. Kurten, *Phys. Chem. Chem. Phys.*, 2019, **21**, 18495–18500.
- 103 A. Köhler, A. L. Khan, J. S. Wilson, C. Dosche, M. K. Al-Suti, H. H. Shah and M. S. Khan, *J. Chem. Phys.*, 2012, **136**, 094905.
- 104 D. G. Thompson, J. R. Schoonover, C. J. Timpson and T. J. Meyer, *J. Phys. Chem. A*, 2003, **107**, 10250–10260.
- 105 S. D. Cummings and R. Eisenberg, *J. Am. Chem. Soc.*, 1996, **118**, 1949–1960.
- 106 E. M. Kober, J. V. Caspar, R. S. Lumpkin and T. J. Meyer, *J. Phys. Chem.*, 1986, **90**, 3722–3734.
- 107 J. V. Caspar, E. M. Kober, B. P. Sullivan and T. J. Meyer, *J. Am. Chem. Soc.*, 1982, **104**, 630–632.
- 108 S. Dong, W. Xu, H. Guo, W. Yan, M. Zhang and F. Li, *Phys. Chem. Chem. Phys.*, 2018, **20**, 18657–18662.

