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ARTICLE

Sodium-based Donor-Acceptor Assemblies Featuring Thermally Activated Delayed Fluorescence Enabled by Highly Efficient Through-Space Charge Transfer

Received 00th January 20xx,
Accepted 00th January 20xx

DOI: 10.1039/x0xx00000x

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Exploring more economical and innovative alternatives to precious 4d and 5d metal complexes used in photocatalysis, OLEDs, or energy conversion has primarily focused on 3d metals, leaving highly abundant alkali metals underexplored. We show that under stoichiometry control of 1,10-phenanthroline-5-carbonitrile (**^{CN}phen**) and sodium 2,6-bis(trimethylsilyl)benzenethiolate Na(**^{TMS}BT**), sodium complexes can be obtained that either form the neutral linear assembly [Na(THF)(^{CN}phen)(^{TMS}BT)]$_N$ (**Na1D**), or {[Na(^{CN}phen)$_4$](^{TMS}BT)}$_N$ (**Na2D**) as a 2D-polymeric arrangement with octahedrally coordinated Na, similar to 18 valence electron transition metal complexes, and the thiolate acting as counter anion. **Na1D** and **Na2D** feature bright visible light absorption and thermally activated delayed fluorescence (TADF) from intra- and intermolecular through-space charge-transfer ($^{1/3}$TSCT) excited states, respectively, with high radiative rates k_{TADF} up to 4.5 × 10^5 s^{-1}, unprecedented for alkali metal-based luminophores. Solution studies revealed extensive dynamic behavior, including reversible metal–ligand bond dissociation. However, the Na assemblies show $^{1/3}$TSCT emission ($\lambda_{em,max}$ = 555 nm) in solution and for the first time we have successfully employed Na-based compounds as visible light photosensitizers in Dexter energy transfer catalysis. This study demonstrates the critical role of TSCT states in constructing photoactive coordination complexes and indicates an underexplored, yet significant potential of sodium-based luminophores as TADF emitters and earth-abundant photocatalysts.

Introduction

The photoluminescent (PL) phenomenon is essential in many application fields, including organic light-emitting diodes,[1, 2] photocatalysis,[3, 4] sensing,[5-7] and molecular imaging.[8, 9] Particular emphasis of current research has been laid on developing new photocatalysts and emitters with high radiative rate constants k_r for electroluminescent devices, applications for which the efficient generation of electronically excited triplet states is highly beneficial. For example, while the formally spin-forbidden intersystem-crossing (ISC) $S_n \rightarrow T_1$ promotes photoreactivity due to an extended excited state lifetime in comparison to the fluorescent S_1 state, triplet state emission allows for unitary exciton utilization in devices for maximum efficiency.[10] An interesting emission mechanism for many of the abovementioned applications is thermally activated delayed fluorescence (TADF),[11] which involves thermally-driven reverse ISC (rISC) $T_1 \rightarrow S_1$, ultimately leading to fast, spin-allowed emission from the higher-lying singlet excited state. Thus, spin-forbidden phosphorescence is bypassed, but triplet excitons can still be harvested. TADF is only enabled for systems with a small singlet-triplet excited state energy gap (ΔE_{ST}), typically below 200 meV.[12] This can be achieved by constructing a donor-acceptor (D–A) motif that leads to charge transfer (CT) excited states with a spatially well-separated electron-hole pair, reducing their exchange-interaction integral that is proportional to ΔE_{ST}.

Recent efforts to explore new PL complexes based on earth-abundant 3d metals led to the discovery of several groundbreaking examples that can compete with traditionally employed compounds based on precious heavy 4d and 5d transition metals, either due to their high luminescent performance or photo(redox)catalysis-relevant properties. Although complexes of 3d metals in d^1-d^9 electron configuration currently do not offer photophysical properties suitable for the application in electroluminescent devices due to the fast non-radiative decay via metal-centered (MC) states,[13] a handful of examples based on, e.g., Cr, Co or Fe metal cores were engaged in relevant photochemistry, including photoredox catalysis and photo-induced energy transfer catalysis, displaying in some instances activity competitive to heavy metal-based photosensitizers.[14-16] A d^{10} configuration, on the other hand, allows to reach highly emissive complexes with k_r that might overcome the second generation of phosphorescent OLED emitters. Cu^I compounds with archetypal carbene-copper-

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amide (CCA) structural motifs are well-explored in this field. A TADF mechanism was observed for CCA complexes with highly efficient emission featuring near-unity quantum yield.[17] Notably, extensive tunability of photophysical properties is characteristic of this type of emitters. For instance, the emission wavelength can be manipulated by modulating the energy of the LUMO,[18, 19] comprising the π^* orbital located on the carbene. Additionally, the radiative rate constant might be improved by reducing ΔE_{ST}.[20, 21] Besides Cu^I, carbene zinc(II) thiolates (CZT) TADF emitters are currently on the rise.[22, 23] Very recently we demonstrated that structural control of the relative ligand orientation in trigonal planar CZT compounds enables TADF from ligand-to-ligand/ligand-to-metal $^{1/3}$(LL/LM)CT excited states with k_r in the range of 10^6 s^{-1}, faster than commercially used Ir-based emitter.[23] Luminescent 3d metal complexes are experiencing a golden era; however, additional highly abundant metals should also be considered as potential substitutes or complements to expensive heavy transition metal luminophores.

In principle, the alkali metal sodium can be considered a very promising candidate for constructing new photoluminescent complexes due to its very high abundance of 2.36% in the Earth's crust. Sodium complexes feature a flexible coordination sphere and show great structural diversity, which are additional significant benefits for exploring novel photoluminescent motifs. This chemical behavior can be ascribed to the negligible covalent character of the Na^+–L bonds, leaving Coulombic interactions the dominant attractive force responsible for the less energetic and more flexible bonding between the metal ion and ligands. Unfortunately, such high molecular flexibility is often detrimental to photoluminescence due to pronounced vibrational relaxation to the ground state S_0. Thus, sodium-based molecules have barely been considered as luminescent materials in literature.

One of the rare examples of a sodium-based luminescent compound was reported recently by Hinz *et al.*, who observed prompt singlet excited state emission from toluene-coordinated Na^+ carbazolide complexes in the solid state with no evidence for ISC (Fig. 1, **A**).[24] The latter finding suggests that spin-orbit coupling (SOC) is insufficient in these light metal complexes to foster the formation of triplet states. Roesky and co-workers prepared a dimeric sodium complex of iminophosphonamides **B** (Fig. 1) featuring TADF, for which triplet formation was attributed to symmetry-breaking intra-ligand CT.[25] Although this strategy can promote ISC for specific dimeric molecules,[26, 27] it comes at the price of reduced oscillator strength and a long S_1 lifetime of τ = 230 ns for **B**, resulting in a relatively low k_{TADF} of 5×10^4 s^{-1}, as well as limited structural flexibility for photophysical modification.

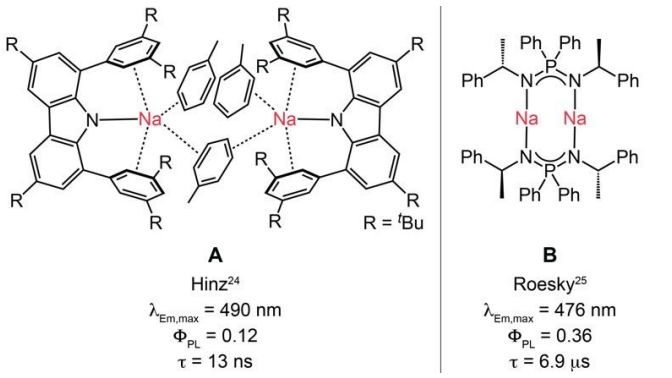

Fig. 1 Literature-known Na^+-based photoluminescence systems. Fluorescent (**A**) and TADF (**B**) complexes of sodium featuring intra-ligand CT and selected photophysical data in the solid state at room temperature.

Herein, we focused on a fundamentally different strategy to achieve ISC for triplet excited state formation and emission in sodium complexes, which is based on introducing spatially separated electronic donor and acceptor moieties in the first coordination sphere of the Na^+ ion. The acceptor unit used in this work comprises 1,10-phenanthroline-5-carbonitrile (**^{CN}phen**), whereas the donor fragment is based on 2,6-bis(trimethylsilyl)benzenethiolate (**^{TMS}BT**). In the case of **^{CN}phen**, the nitrile function is expected to strengthen π-electrophilicity due to a strong electron-withdrawing effect. In addition, the steric demand of the **^{TMS}BT** should lead to the rigidification of the system, hindering detrimental vibrational relaxation. Moreover, TMS groups with high electron-donating properties make ^{TMS}BT an exceptionally strong π-donor.

Results and discussion

Synthesis and molecular structures.

Our investigation of sodium-based donor-acceptor systems was commenced by mixing different ratios of ^{CN}phen and Na^{TMS}BT (Fig. 2A) in tetrahydrofuran (THF), which affords orange to red solutions from initially colorless reagents. Vapor diffusion of *n*-pentane into THF solution containing a 1:1 ratio of ^{CN}phen:Na^{TMS}BT yielded yellow single crystals, whereas, for a 2:1 ratio, the formation of deep-red single-crystalline material was observed. Both types of crystals are sensitive to moisture as decolorization occurs under ambient conditions. X-ray diffraction analysis of the yellow sample revealed an unprecedented linear assembly (**Na1D**, Fig. 2B). In contrast, the red crystals consist of a two-dimensional coordination assembly denoted as **Na2D** (Fig. 2C).

For **Na1D**, each Na^+ ion is coordinated by one THF molecule, ^{TMS}BT, a chelating ^{CN}phen ligand, and a bridging –CN group from the neighboring unit, leading to a distorted square-pyramidal geometry around the metal ion. Interatomic distances Na–N(^{CN}phen) of 2.5106(17) and 2.4734(18) Å are in the range of values reported for phenanthroline complexes of sodium with various coordination geometries.[28-32] Likewise, the Na–NC(^{CN}phen) bond length of 2.515(2) Å shows minor distinctions compared to values reported for diverse sodium complexes bearing aliphatic[33, 34] or aromatic[35-37] nitriles, typically falling in the 2.39-2.52 Å range. The Na–S bond length of 2.7971(9) Å is slightly elongated by ca. 0.05-0.1 Å in comparison to sodium complexes containing only solely coordinated thiolate ligands, e.g., dimeric [Na^{Ar}BT]$_2$ (^{Ar}BT = 2,6-bis(2,4,6-triisopropylphenyl)benzenethiolate, or 2,6-bis(2,4,6-trimethylphenyl)benzenethiolate).[38, 39] In contrast, sodium benzenethiolates bearing additional σ-donors, such as multidentate aliphatic amines, exhibit Na–S distances typically

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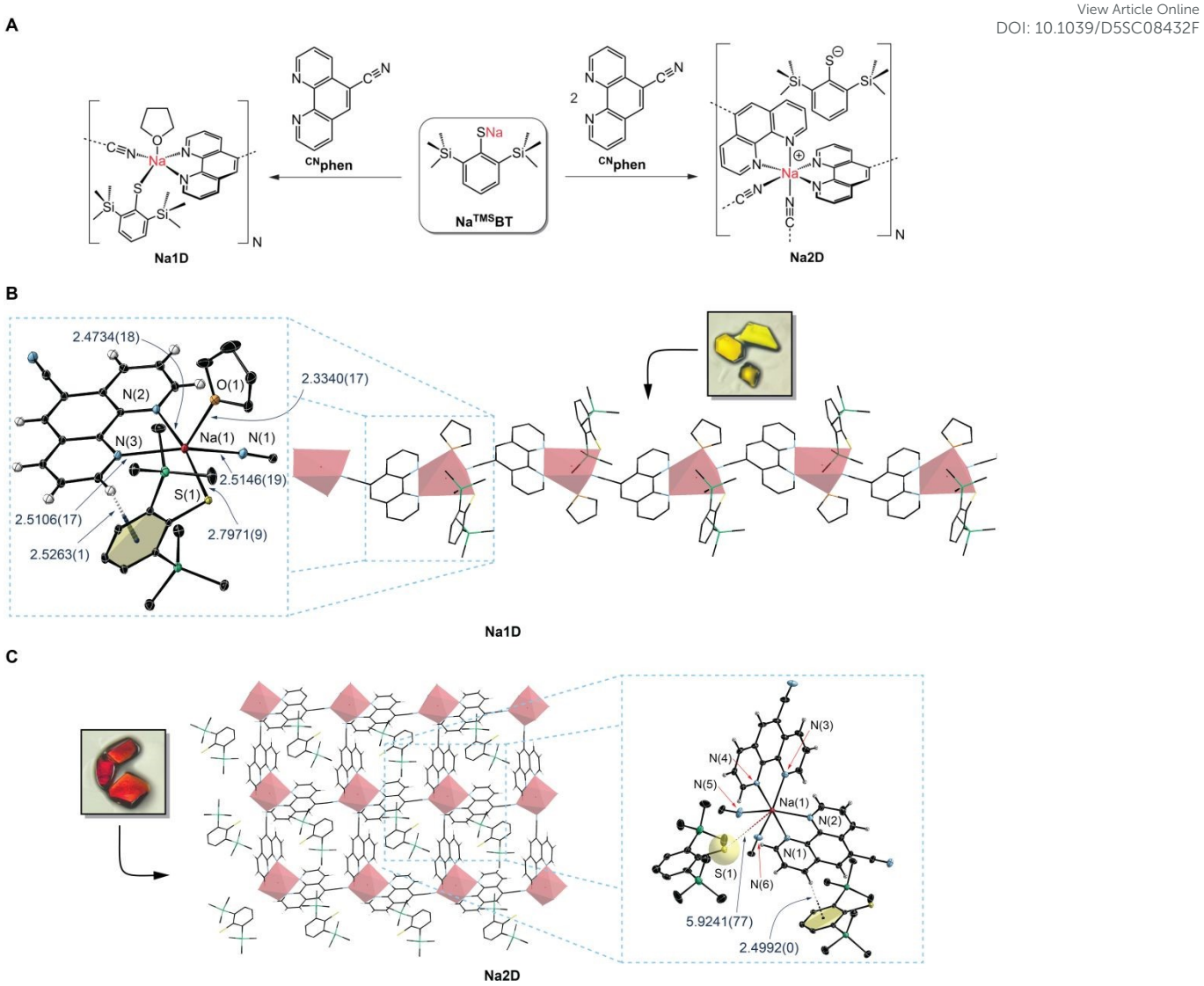

Fig. 2 Synthesis, structures, and macromolecular architectures of **Na1D** and **Na2D**. **A**: Synthesis of the sodium-based assemblies. **B**-**C**: Molecular structures of monomeric units of **Na1D** and **Na2D** and their macromolecular architectures. Insets show photographs of single crystals of each assembly. Thermal ellipsoids are drawn at a 30% probability level. Crystal solvents (*n*-pentane for **Na1D** and THF for **Na2D**) are omitted for clarity. Selected interatomic distances are given in Å.

in the range of 2.78-2.84 Å that are slightly elongated compared to **Na1D**.[40] The **Na2D** assembly is based on a monomeric unit comprising two ^{CN}phen coordinated to Na^+ in a typical chelating fashion, with two additional –CN bridging groups from neighboring units completing the pseudo-octahedral coordination environment. While the Na–N(^{CN}phen) and Na–NC(^{CN}phen) interatomic distances are comparable to the **Na1D** assembly, the ^{TMS}BT acts as an outer sphere counter anion in the voids of the two-dimensional network, as evidenced by the long S–Na interatomic distance of 5.9241(77) Å (Fig. 2B).

Non-covalent interactions of Na1D and Na2D.

We observed that the ^{CN}phen and ^{TMS}BT ligands in **Na1D** adopt a close-to-perpendicular orientation with a dihedral angle of 78.6°, presumably as a consequence of a weak non-covalent C–*H*···π interaction as indicated by the short distance of 2.5263(1) Å between the hydrogen atom in the 2-position of the phenanthroline ligand and the benzene centroid of ^{TMS}BT (Fig. 2, B). Although the ^{TMS}BT anions in the network voids of **Na2D** do not interact with the sodium ions, their orthogonal orientation relative to the ^{CN}phen ligands allows for dense packing with T-shape geometries and the closest spatial separation of 2.4992(0) Å. Such a short distance also suggests significant C–*H*···π non-covalent intermolecular interaction. In addition, the ^{CN}phen ligands of **Na2D** also interact non-covalently with neighboring ^{CN}phen as indicated by the tilted T-shape (TT) arrangements and C–*H*···π separation in the range of 2.95-3.45 Å (for details, see Supporting Information). Within each monomeric unit of **Na2D**, this type of TT interaction is always maintained for the pair of ^{CN}phen ligands, one bound in a bidentate fashion and the second via the –CN group to the same Na^+ cation. We note that the N–Na–N angle defined by the nitrile and heterocyclic nitrogen in *trans*-position of the nitrile is only 152° because the intermolecular C–*H*···π

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interactions are distorting the octahedral coordination geometry of the Na^+ ion. The realization of ultra-short D–A distances below 2.8 Å for strong through-space electronic communication (TSEC) is challenging, especially for organic systems, where donor and acceptor fragments are linked via a rigid, covalent spacer. Compared with organic and organometallic TADF emitters for which non-covalent D–A coupling was rationally designed,[41-46] the C–*H*···π distances between ^{CN}phen and ^{TMS}**BT** of *ca*. 2.5 Å in **Na1D** and **Na2D** are exceptionally short and close to the calculated separation for a benzene T-shaped dimer, providing substantial stabilization energy.[47] The fact that bonding in sodium complexes mainly arises from Columbic interactions which are generally weaker and more flexible than covalent bonds results in a structure-determining effect of the ^{TMS}BT···^{CN}phen and ^{CN}phen···^{CN}phen interactions, allowing for the ultra-short distances between these D–A pairs. This aspect of Na-based complexes has a game-changing potential for constructing molecular systems where enhanced non-covalent interactions are required, for instance, in material science or drug design.[48, 49] TSEC between donor and acceptor ligands also bears great potential for the development of new emitter classes as it promotes the formation of through-space charge transfer (TSCT) excited states,[41, 43, 44, 50, 51] which commonly enhances rISC by reducing ΔE_{ST} due to the absence of through-bond D–A conjugation. As a result, non-radiative deactivation can be suppressed in favor of improved PL performance.[42, 44, 52, 53] However, the decisive parameter affecting the efficiency of TSEC is the spatial D–A separation since a distance beyond ca. 3.0 Å may lead to a substantial decrease in CT rate.[50]

Photophysical properties and emission mechanism in the solid

Surprisingly, upon 450 nm light irradiation, single crystals of **Na1D** and **Na2D** show bright orange to deep red luminescence at room temperature with very broad spectral appearance and $\lambda_{em,max}$ of 645 and 715 nm, respectively, without tailing into the near-infrared region (NIR) (Table 1 and Fig. 3A-B and inset of Fig. 3F). Such low emission energies are highly unusual not only for luminescent complexes of sodium but in general for alkali and earth-alkali metal-based compounds that typically feature blue to green fluorescence with $\lambda_{em,max}$ below *ca*. 540 nm.[24, 25, 54, 55] According to our time-dependent density functional theory (TD-DFT) calculation (see below), this unique photophysical characteristic of **Na1D** and **Na2D** stems from a ^{TMS}BT→^{CN}phen TSCT mechanism, which commonly red-shifts emission profiles due to weak electronic coupling and stabilization of the ground state via non-covalent interactions. Such modulation of emission energy is potentially of interest for future developments regarding applications where red-to-NIR emission is beneficial, such as bio-imaging[56] or telecommunication.[57] The relatively low photoluminescence quantum yields Φ_{PL} of 0.036 (**Na1D**) and 0.013 (**Na2D**) (Table 1) result from vibrational coupling between the ground and excited states according to the energy gap law,[58] presumably enhanced by significant photo-initiated structural distortions of the assemblies as evidenced by the large apparent Stokes shifts and full widths at half maximum of 3956 (**Na1D**) and 3225 cm^{-1} (**Na2D**). Lifetime (LT) measurements revealed biexponential decays (Fig. 3C) with average values $\langle\tau\rangle$ of 221 and 29 ns, affording estimated k_r of 1.6 (**Na1D**) and 4.5 × 10^5 s^{-1} (**Na2D**) indicative for spin-forbidden triplet excited states being involved.

Variable-temperature (VT) measurements for **Na1D** reveal an increase of $\langle\tau\rangle$ and Φ_{PL} to 2956 ns and 0.16, respectively. The resulting slight decrease of k_r = 0.5 × 10^5 s^{-1} in conjunction with the hypsochromic shift of the emission maximum from 645 to 607 nm (ΔE = 970 cm^{-1}) at 77 K suggests temperature-dependent rigidification and change of the Frank-Condon factors of the rather flexible emitter system. Upon further cooling, however, an enormous elongation of excited state lifetime by more than three orders of magnitude compared to room temperature is observed to reach ca. 0.4 ms at 7 K (Fig. 3E left and Table 1). This cannot be explained by additional rigidification but by changing the nature of the emitting excited state. Importantly, the emission spectrum at 7 K with $\lambda_{em,max}$ = 605 nm shows only a minor additional hypsochromic shift by 55 cm^{-1} compared to the spectrum recorded at 77 K with no signs of vibrational progression, i.e., the CT character of the emissive state is maintained even at very low temperature. Such a photophysical behavior is typical for a TADF mechanism with a very small energy gap ΔE_{ST} that allows the detection of emissive triplets only at very low temperatures, which we conclude to stem from the formation of interconverting $^{1/3}$TSCT states. A very similar behavior of the VT data was also found for **Na2D** (Fig. 3E right). It is important to note that for a wide temperature range, the experimentally obtained luminescence lifetime decays are bi-exponential, analogously as for the 297 K data shown in Fig. 3C. However, we observed an additional short-lived component of ca. 0.9 to 1.0 ns (Supplementary Figures S47 and S48), of which the contribution becomes larger with decreasing temperature in particular below ca. 170 K (see Fig. 3D and inset). Since sulfur is the heaviest atom in the **Na1D** and **Na2D** systems, it is reasonable to expect the (r)ISC processes to be dominated by a spin-vibronic coupling mechanism. At low temperatures, where the vibrational motions are significantly suppressed, the rate/efficiency of spin-vibronic-driven ISC would be expected to be reduced. Thus, we attributed the short-lived components to residual prompt fluorescence from the ^{1}TSCT state and excluded it from the $\langle\tau\rangle^{TADF}$ calculation, and estimate the rate constant k_{ISC} at room

Table 1 Selected photophysical data for **Na1D** and **Na2D** (see also Supporting Information).

	T [K]	λ_{em} [nm]	$\langle\tau\rangle$ [ns] [a]	Φ_{PL}	k_r [s^{-1}] [b]
Na1D	297	645	221	0.036	1.6×10^5
solid[c]	77	607	2956	0.16	0.5×10^5
	7	605	435×10^3	-	-
Na2D	297	715	29	0.013	4.5×10^5
solid[c]	77	656	251	0.053	2.1×10^5
	7	656	125×10^3	-	-
Na1D THF	297	555	-	-	-

[a] amplitude average lifetimes, [b] $k_r = \Phi_{PL}/\langle\tau\rangle$, [c] single-crystalline form

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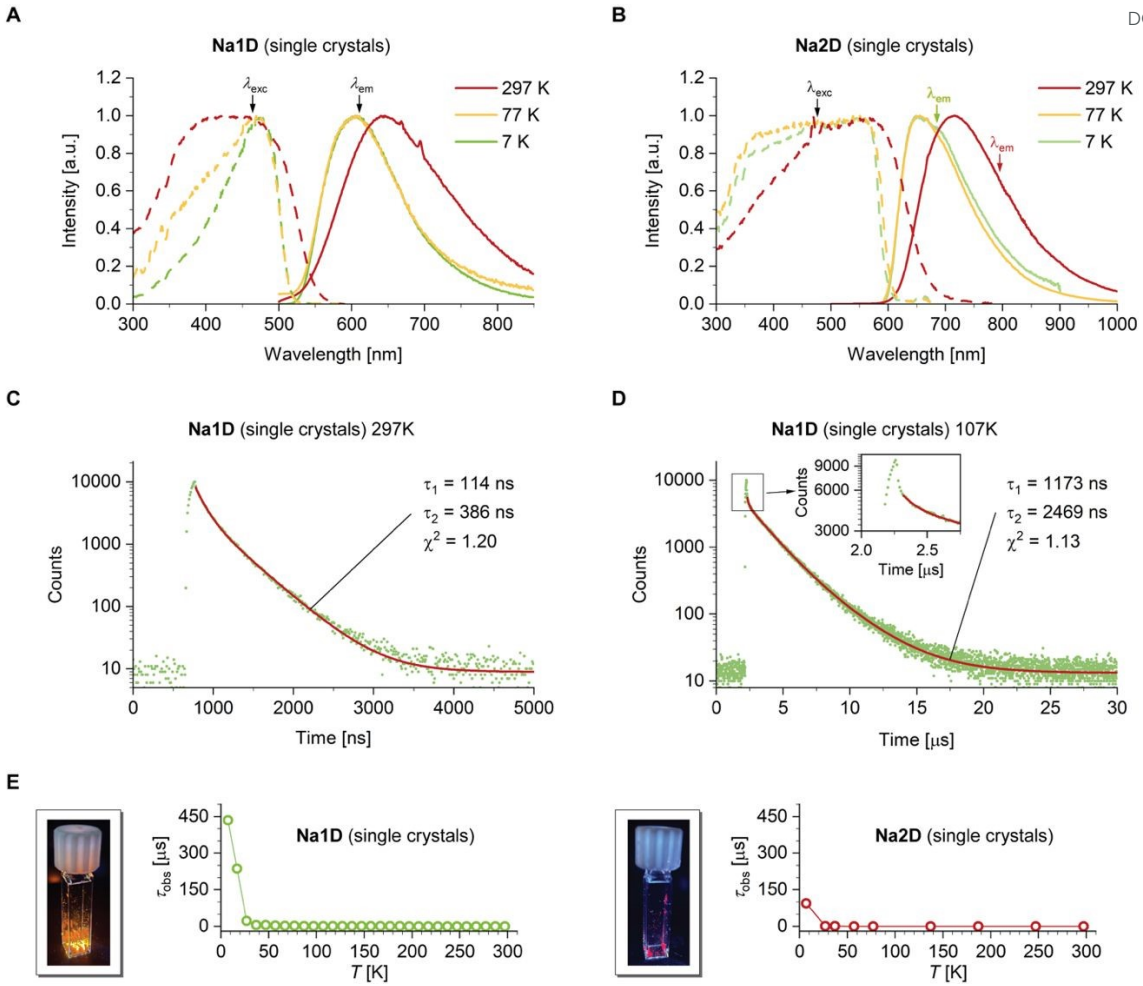

Fig. 3 Photophysical characterization of **Na1D** and **Na2D**. Top: variable-temperature emission (solid lines) and excitation (dashed lines) spectra of single-crystalline samples of **Na1D** (**A**) and **Na2D** (**B**). Middle: time-resolved excited state decays of **Na1D** (λ_{ex} = 450 nm, λ_{em} = 600 nm) recorded at 297 (**C**) and 107 K (**D**). Inset: detail of short-lived component observed at lower temperatures. Bottom: temperature-dependence of the observed lifetime recorded for single-crystalline samples of **Na1D** (**E**, left) and **Na2D** (**E**, right) and photographs of the luminescence of single crystals of **Na1D** and **Na2D** at room temperature upon 365 nm irradiation.

temperature to be ca. 10^9 s^{-1}. Assuming that prompt fluorescence occurs with Φ_{PL} of 0.001-0.01, the resulting fluorescence rate k_F of 10^6-10^7 s^{-1} leaves us to conclude that reverse ISC occurs with k_{rISC} = 10^7-10^8 s^{-1} for **Na1D** and **Na2D**, which are compatible with established organic TADF emitters. The estimated values appear reasonable as the oscillator strength for prompt fluorescence from a ^{1}TSCT state would be expected to be small, while a very small energy gap ΔE_{ST} would foster fast ISC and also rISC processes.

To gain deeper insight into the photophysical properties of **Na1D** and **Na2D**, we performed TD-DFT calculations on the monomeric units extracted from X-ray diffraction data with the CN-coordinating phen ligand being substituted by acetonitrile as simplified models of the polymeric structures (for full details, see Supporting information), assuming localization of the excitons (see below). Under these approximations, the TD-DFT calculations provided energetically nearly identical S_1 and T_1 states (Fig. 4 and Supporting Fig. 49) with ΔE_{ST} values of only 2 (**Na1D**) and 31 meV (**Na2D**), which coincide with the proposed close-to-isoenergetic $^{1/3}$TSCT excited states, allowing for efficient (r)ISC processes in a TSCT-TADF mechanism (see above). Considering the polymeric nature of **Na1D** and **Na2D**, we further investigated whether the transport of optically excited charges in the form of exciton hopping can occur. For both compounds we obtained large single crystals suitable for time-resolved 400 nm pump terahertz (THz) transmission probe spectroscopic experiments with femtosecond time resolution.

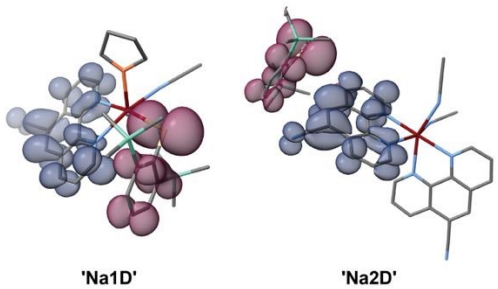

Fig. 4 TD-DFT difference densities of the $S_0 \rightarrow S_1$ excitations of monomeric units isolated from X-ray diffraction data of **Na1D** and **Na2D**. Areas losing electron density are depicted in purple, and areas gaining in blue.

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This is a well-established approach to investigating charge carrier dynamics in nonequilibrium states.[59, 60] As presented in Fig. 5, the change of the transmitted THz electric field ΔE is recorded as a function of a time delay between the 400 nm excitation laser pulse and the peak electric field of the THz probe pulse. For a variety of pump fluences from 35 to 583 μJ/cm^2, which is well above the threshold for the observation of photoluminescence, we cannot resolve any change in the THz probe pulse. This is in clear contrast to the situation of, e.g., a semiconductor, where an optical excitation creates additional charge carriers (i.e., electrons and holes), whose transport leads to enhanced absorption of the THz field.[60] The observed absence of THz absorption indicates that the optically excited charges in **Na1D** and **Na2D** are mostly localized. Otherwise, the transport of these charges will not only result in an absorption of the THz field but also an enhanced absorption at higher pump fluences.

Intramolecular TSCT-derived emission related to metal complexes is very rare and underexplored. Usually, the high degree of metal–ligand bond covalency restricts the molecular geometry and prevents its distortion from optimizing the D–A spatial separation and non-covalent interactions, limiting TS electronic coupling between donor and acceptor moieties. Consequently, TSCT excited states with transition metal complexes are restricted to very few specific molecular designs with limited chemical tunability. For instance, a rapid TSCT-TADF process due to extremely low ΔE_{ST} of *ca*. 7 meV and rISC rate constant up to 10^7 s^{-1} was recently reported for a new type of CCA complex, consisting of a carbazolide donor and functionalized N-heterocyclic carbene (NHC).[42] In this case, however, TSCT is only enabled for NHCs decorated with phenylsulfonyl flanking units that act as acceptors due to the strongly electron-withdrawing sulfonyl function,[42] while the herein-reported sodium-based assemblies provide ample possibilities for further excited state modification.

Coordination behavior in solution

An additional important photophysical feature of our TSCT assemblies Na1D and Na2D presents visible light absorption (excitation spectra in Fig. 3A-B), critical for many applications, such as photocatalysis. The direct comparison with the previously reported Na-based TADF emitter (Fig. 1, B), which can only be excited by UV light,[25] clearly demonstrates the potential of TSCT donor–acceptor motifs to access luminescent alkaline metal complexes with visible-light absorption. Thus, we were further interested in whether a molecular species based on the assemblies featuring TS electronic donor-acceptor communication might persist in a solution for potential photocatalysis.

Despite the diverse solid-state structures of **Na1D** and **Na2D**, ^{1}H and ^{13}C{^{1}H} NMR spectra in THF-d$_8$ have an analogous character with only minor variations in chemical shifts (Supporting Fig. 25). In the ^{1}H NMR spectra, the only significant spectral differences were found in the integral intensities of the observed resonances with the expected ratios for the ^{TMS}BT and ^{CN}phen ligands of 1:1 and 1:2 for **Na1D** and **Na2D**, respectively. Interestingly, the spectra show signals of one molecule THF (3.62 and 1.78 ppm) originating from a non-coordinated solvent.[61] For **Na2D**, this is expected as single crystals used for NMR measurements contain one molecule of co-crystallized THF in the unit cell, while for **Na1D,** this implies either an exchange of coordinated THF for THF-d$_8$ or a more complicated solution behavior of both assemblies, which appears to be independent of the ^{TMS}BT:^{CN}phen ratio. ^{1}H diffusion-ordered spectroscopy (DOSY) of **Na1D** dissolved in THF-d$_8$ revealed extensive ligand dissociation and the formation of free ^{CN}phen, ^{TMS}BT, and THF (for detailed ^{1}H DOSY NMR analysis, see Supporting Fig. S28). Notably, the ^{1}H-^{1}H Nuclear Overhauser Effect spectroscopy (NOESY) revealed NOE between the protons of the TMS group and those of ^{CN}phen in position 9 or 2.

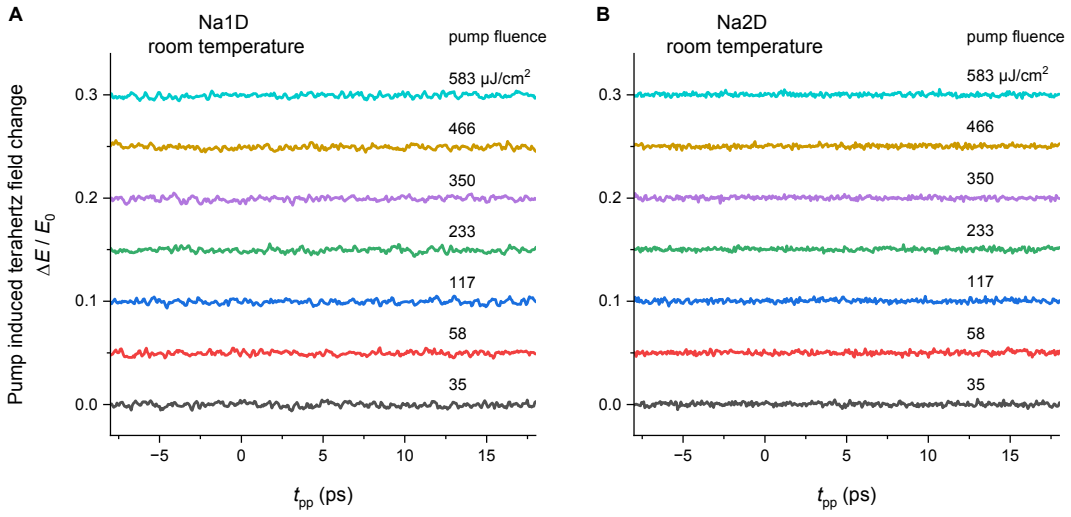

Fig. 5 400 nm optical pump terahertz transmission probe spectroscopy of **Na1D** (A) and **Na2D** (B) at 297 K. The pump-induced change of the transmitted terahertz electric field Δ E/E_0 is measured as a function of pump-probe time delay t_{pp} for various fluences of the 400 nm pump pulses, where E_0 denotes the peak electric field of the terahertz probe pulse without the optical pump.

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This observation suggests a close proximity of the respective protons similar to the ligand arrangement found in the solid state with a distance of only ca. 3.2 Å, and thus reversible dissociation from/association to sodium in a solution of ^{CN}phen and ^{TMS}BT. The exact Na^+ complex solution structure is unclear and present only in very low concentrations as this equilibrium is shifted towards mostly dissociated ligands, preventing identification by 1D NMR techniques or ^{1}H DOSY NMR spectroscopy at room temperature. However, at –100°C, the original ^{1}H NMR signal of the protons in positions 2/9 on ^{CN}phen (9.12 ppm at 297 K) splits into two doublets with chemical shifts of 9.32 and 9.28 ppm (Supplementary Fig. 27), indicating a temperature-dependent reversible complexation–dissociation equilibrium that can be shifted towards the sodium complex under kinetic control. The ^{15}N NMR resonance of heterocyclic nitrogen atoms is shifted upfield by 9.3 ppm at –100°C to a value of –88.4 ppm, which is also expected upon the coordination of ^{CN}phen with metal ions.[62]

Absorption, emission, and photocatalytic triplet energy transfer properties in solution

At concentrations below *ca*. 10^{-4} M, the steady-state UV/Vis spectra of Na1D dissolved in THF show solely high-energy excitations below ca. 370 nm (Fig. 6A), which we attribute by comparison to π→π* and n(E)→π*(phen) (E = N, S) transitions of free colorless ^{CN}phen and ^{TMS}BT, respectively (Fig. 6B). However, at concentrations > 5 × 10^{-4} M, an additional broad band in the visible between 410-650 nm with $\lambda_{abs,max}$ = 428 nm is observed that we assign to CT excitation between ^{CN}phen and ^{TMS}BT due to a concentration-dependent shift of the equilibria investigated by NMR spectroscopy (*vide supra*). Interestingly, previously reported coordination complexes bearing phen and benzenethiolate ligands, for example, [Zn(BT)$_2$(phen)],[63, 64] are typically colorless with only near-UV LLCT excitations. Considering the pronounced tendency to form C–*H*···π non-covalent interactions in the solid state and the observed NOE between ^{CN}phen and ^{TMS}BT, proving the proximity of these two ligands in solutions even at room temperature, the occurrence of electrostatic-controlled low energy ^{TMS}BT→^{CN}phen TSCT facilitated by the sodium ion would explain the shift into the visible observed for dissolved Na1D. It is important to note that such intense low-energy absorption bands were not observed for pure solutions of ^{CN}phen or Na^{TMS}BT, even at higher concentrations (Supporting Fig. 40A,B), excluding the formation of ^{CN}phen- or Na^{TMS}BT-based solvent exciplexes. In line with the VT-NMR data discussed above, cooling THF solutions of Na1D also affects the color of the solution from light orange to red due to the equilibrium shift towards the sodium coordination complex (Fig. 6C). As shown in Fig 6A, at 173 K (red traces), the low energy band shows significant bathochromic shift to 450 nm with the apparent shoulder at *ca*. 550 nm and considerable broadening, leading to offset of the band up to 670 nm. Notably, the absorption intensity increases significantly by factors of *ca*. 2.5 and 10 at 450 and 550 nm, respectively. These findings further prove preferable complexation at low temperatures, for

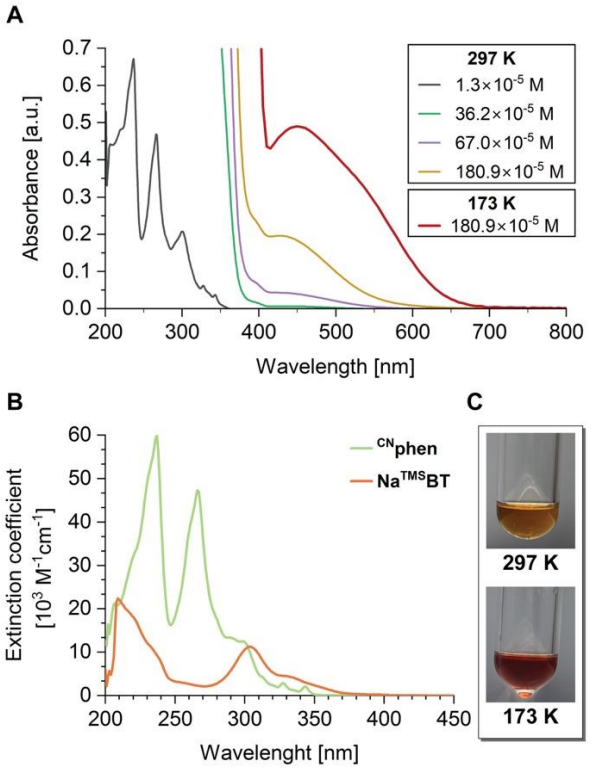

Fig. 6. A: concentration-dependent absorption spectra recorded upon dissolving **Na1D** in THF. **B**: absorption spectra of ^{CN}phen (green) and Na^{TMS}BT (orange) in THF solution. **C**: photographs of solutions of **Na1D** (~180 × 10^{-5} M) at 297 K (top) and 173 K (bottom).

which more pronounced ^{TMS}BT→^{CN}phen TSCT visible-light excitations are expected.

Excitation into the visible absorption band enabled by TSCT transitions in the range of 470-500 nm resulted in very weak broad luminescence with $\lambda_{em,max}$ = 555 nm (Table 1 and Fig. 7A), similar to the properties of **Na1D** in the solid state. Although we could not deduce the emission lifetime or quantum yield, the existence of triplet excited states in single crystals of **Na1D** under visible light excitation led us to assume that the luminescence of the assembly in solution is also TADF in nature. This would imply that dissolved **Na1D** may be suitable for homogenous photocatalysis via Dexter energy transfer.

Photocatalytic (*E*)-to-(*Z*) isomerization via triplet energy transfer (EnT) is an important strategy for accessing stereodefined (*Z*)-alkenes as relevant building blocks in synthetic chemistry, which are typically thermodynamically less stable compared to their more common (*E*)-counterparts.[65, 66] The key concept behind the isomerization is the promotion of electron density from π to π* orbitals accompanied by lowering of the C=C bond order, allowing rotation and subsequent relaxation to the (*Z*)-alkene configuration. However, since the lowest π→π* singlet excitations typically require high-energy UV light, transition metal complexes that can be excited with visible light are typically used for sensitization of the alkene ^{3}ππ* states.[67, 68] Bearing in mind the assumed triplet excited state energy of **Na1D** in solution of ~2.23 eV ($\lambda_{em,max}$ ~ 555 nm), we employed our novel sodium-

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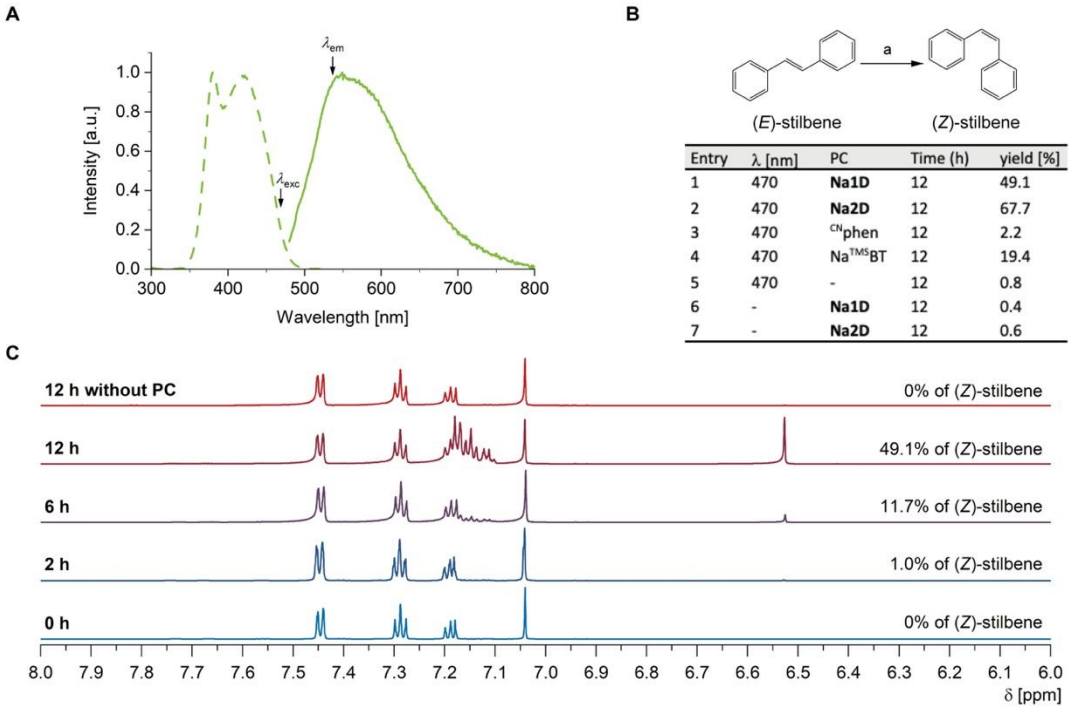
based systems for photo-driven isomerization of (*E*)-stilbene exhibiting a triplet energy of 2.13 eV as a proof of concept test reaction. Irradiation of degassed THF solutions containing (*E*)-stilbene and 5 mol-% of dissolved **Na1D** or **Na2D** with 470 nm blue light promote (*E*)→(*Z*) conversion within 12 h to give the desired product in 50% and 68% yield, respectively (Fig. 7B-C). Importantly, control experiments without light or using only ^{CN}phen as a catalyst gave only negligible formation of (*E*)-stilbene under otherwise identical reaction conditions. Interestingly, we observed up to 19% conversion with Na^{TMS}BT, presumably due to very weak direct $S_0 \rightarrow T_1$ absorption of the ^{TMS}BT enabled by SOC from the sulfur and subsequent EnT.

Recently, efficient energy and electron transfer photocatalysis using a Coulombic dyad composed of tris(1,10-phenanthroline)ruthenium(II) bound non-covalently to pyrene-1,3,6,8-tetrasulfonate was reported by Kerzig et al.[69] Our findings demonstrate for the first time that a similar approach is applicable also for simple organic alkaline metal salts, such as NaTMSBT, which might act as photoactive species in triplet EnT processes. Additional topical studies have shown that organic and transition metal photocatalysts can decompose within the first short period of the overall reaction time, although the conversion continues.[70] Bearing in mind the coexistence of various salts in the reaction mixtures of typical photocatalytic applications, our study indicates that they may be involved in the decisive photoinitiation steps. In addition, the activity and required excitation wavelength can be efficiently modulated by employing or formation of donor–acceptor motifs, which enhances visible-light absorption and likely promotes ISC due to lowering ΔE_{ST} as demonstrated by the application of **Na1D** and **Na2D**.

Conclusion

In contrast to previous studies focusing on prompt fluorescence or TADF originating from intra-ligand CT states in sodium compounds, we have shown a new approach to foster emission involving triplet excited states in abundant alkaline metal complexes by generating through-space CT. The critical design motif we deduce from our case study is a D/A orientation around the Na ion that allows for non-covalent interactions. Generally, sodium complexes are dominantly stabilized by the Columbic character of the Na–ligand bonds, which are geometrically more flexible than typically observed in transition metal complexes bearing a much higher degree of covalency. Consequently, the coordination geometry of the sodium ion in our compounds is distorted to accommodate the interligand interactions.

As a far-reaching consequence, a strong D–A through-space coupling is observed for single-crystalline one- (**Na1D**) or two-dimensional (**Na2D**) coordination entities composed of Na^{TMS}BT and ^{CN}phen, leading to efficient ISC and rISC between the $^{1/3}$TSCT states due to their very low experimentally determined ΔE_{ST} of only ca. 55 cm^{-1} (7 meV) as found for **Na1D**. The enabled TADF process leads to deep red to NIR luminescence, which on



Entry	λ [nm]	PC	Time (h)	yield [%]
1	470	**Na1D**	12	49.1
2	470	**Na2D**	12	67.7
3	470	^{CN}phen	12	2.2
4	470	Na^{TMS}BT	12	19.4
5	470	-	12	0.8
6	-	**Na1D**	12	0.4
7	-	**Na2D**	12	0.6

Fig. 7 Solution luminescence and energy transfer. **A**: luminescence (solid) and excitation (dashed) spectra of **Na1D** in THF (λ_{ex} = 470 nm, λ_{em} = 525 nm). **B**: photoisomerization of (*E*)-stilbene, a = photocatalytic conditions are summarized in the table. **C**, Example of the low-field part of ^{1}H NMR spectra showing photocatalytic conversion of (*E*)-stilbene to (*Z*)-stilbene using **Na1D** solution and 470 nm irradiation.

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its own is highly unusual for alkali- or earth-alkaline metal compounds, [24, 25, 54, 55] and high radiative rate constants up to 4.5 × 10^5 s^{-1} that are unprecedented for sodium-based emitters involving triplet excited states. Despite the rather complicated solution behavior, TSCT was also confirmed for the system obtained by dissolving **Na1D** in THF, giving emission at λ_{max} = 555 nm, leading for the first time to the successful application of a sodium complex as a photosensitizer for visible-light-driven triplet EnT, as evidenced by the photoisomerization of (*E*)-stilbene. Despite their insignificant SOC, our case study indicates great potential for highly abundant alkaline and earth-alkaline complexes in future photonic applications involving TADF or red-to-NIR luminescence, such as electroluminescent devices, bio-imaging or telecommunication, and sensitizer compounds for photocatalytic conversions.

Author contributions

O. M. designed the research, wrote the paper, and performed experimental work, including syntheses, photophysical measurements, and data analysis; T. H. performed experimental work, including syntheses, photophysical measurements, and data analysis; A. B. carried out X-ray diffraction analysis; L. H. and I. S. contributed to synthetic work and photophysical measurements; P. P. and Z. W. performed THz measurements and wrote the respective section of the manuscript; A. S. designed the research, analyzed the data, supervised the project and wrote the paper. All authors discussed the results and commented on the manuscript.

Conflicts of interest

There are no conflicts to declare.

Data availability

CCDC 2288161 and 2288162 contain the supplementary crystallographic data for this paper. Supplementary information: synthesis details, characterisation data for compounds including NMR spectroscopy, mass spectrometry, single crystal X-ray diffraction data, theoretical studies, and photophysical data.

Acknowledgements

We thank Martin Hejda (University of Pardubice) for valuable discussions and advice on NMR spectroscopy, and we acknowledge helpful discussions with S. Kovalev and C. Zhu (TU Dortmund University).

References

1. Yersin, H., et al., *The triplet state of organo-transition metal compounds. Triplet harvesting and singlet harvesting for efficient OLEDs.* Coordination Chemistry Reviews, 2011. **255**(21): p. 2622-2652.
2. Yang, X., G. Zhou, and W.-Y. Wong, *Functionalization of phosphorescent emitters and their host materials by main-group elements for phosphorescent organic light-emitting devices.* Chemical Society Reviews, 2015. **44**(23): p. 8484-8575.
3. Welin, E.R., et al., *Photosensitized, energy transfer-mediated organometallic catalysis through electronically excited nickel(II).* Science, 2017. **355**(6323): p. 380-385.
4. Arias-Rotondo, D.M. and J.K. McCusker, *The photophysics of photoredox catalysis: a roadmap for catalyst design.* Chemical Society Reviews, 2016. **45**(21): p. 5803-5820.
5. Krämer, J., et al., *Molecular Probes, Chemosensors, and Nanosensors for Optical Detection of Biorelevant Molecules and Ions in Aqueous Media and Biofluids.* Chemical Reviews, 2022. **122**(3): p. 3459-3636.
6. Zhou, X., et al., *Recent Progress on the Development of Chemosensors for Gases.* Chemical Reviews, 2015. **115**(15): p. 7944-8000.
7. Wenger, O.S., *Vapochromism in Organometallic and Coordination Complexes: Chemical Sensors for Volatile Organic Compounds.* Chemical Reviews, 2013. **113**(5): p. 3686-3733.
8. Yang, Y., et al., *Luminescent Chemodosimeters for Bioimaging.* Chemical Reviews, 2013. **113**(1): p. 192-270.
9. Zhang, K.Y., et al., *Long-Lived Emissive Probes for Time-Resolved Photoluminescence Bioimaging and Biosensing.* Chemical Reviews, 2018. **118**(4): p. 1770-1839.
10. Baldo, M.A., et al., *Highly efficient phosphorescent emission from organic electroluminescent devices.* Nature, 1998. **395**(6698): p. 151-154.
11. Parker, C.A. and C.G. Hatchard, *Triplet-singlet emission in fluid solutions. Phosphorescence of eosin.* Transactions of the Faraday Society, 1961. **57**(0): p. 1894-1904.
12. Dos Santos, J.M., et al., *The Golden Age of Thermally Activated Delayed Fluorescence Materials: Design and Exploitation.* Chemical Reviews, 2024. **124**(24): p. 13736-14110.
13. McCusker, J.K., *Electronic structure in the transition metal block and its implications for light harvesting.* Science, 2019. **363**(6426): p. 484-488.
14. Sinha, N., et al., *Cobalt(III) Carbene Complex with an Electronic Excited-State Structure Similar to Cyclometalated Iridium(III) Compounds.* Journal of the American Chemical Society, 2022. **144**(22): p. 9859-9873.
15. Chan, A.Y., et al., *Exploiting the Marcus inverted region for first-row transition metal–based photoredox catalysis.* Science, 2023. **382**(6667): p. 191-197.

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16. Leis, W., et al., *A Photoreactive Iron(II) Complex Luminophore.* Journal of the American Chemical Society, 2022. **144**(3): p. 1169-1173.
17. Hamze, R., et al., *Eliminating nonradiative decay in Cu(I) emitters: >99% quantum efficiency and microsecond lifetime.* Science, 2019. **363**(6427): p. 601-606.
18. Gernert, M., et al., *Cyclic (Amino)(aryl)carbenes Enter the Field of Chromophore Ligands: Expanded π System Leads to Unusually Deep Red Emitting CuI Compounds.* Journal of the American Chemical Society, 2020. **142**(19): p. 8897-8909.
19. Shi, S., et al., *Highly Efficient Photo- and Electroluminescence from Two-Coordinate Cu(I) Complexes Featuring Nonconventional N-Heterocyclic Carbenes.* Journal of the American Chemical Society, 2019. **141**(8): p. 3576-3588.
20. Hamze, R., et al., *"Quick-Silver" from a Systematic Study of Highly Luminescent, Two-Coordinate, d10 Coinage Metal Complexes.* Journal of the American Chemical Society, 2019. **141**(21): p. 8616-8626.
21. Muniz, C.N., et al., *π-Extended Ligands in Two-Coordinate Coinage Metal Complexes.* Journal of the American Chemical Society, 2022. **144**(39): p. 17916-17928.
22. Mrózek, O., et al., *An Air- and Moisture-stable Zinc(II) Carbene Dithiolate Dimer Showing Fast Thermally Activated Delayed Fluorescence and Dexter Energy Transfer Catalysis.* Chemistry – A European Journal, 2023. **29**(23): p. e202203980.
23. Mitra, M., et al., *Structural Control of Highly Efficient Thermally Activated Delayed Fluorescence in Carbene Zinc(II) Dithiolates.* Angewandte Chemie International Edition, 2024. **63**(7): p. e202316300.
24. Kaiser, M., et al., *Efficient fluorescence of alkali metal carbazolides.* Inorganic Chemistry Frontiers, 2023. **10**(10): p. 2987-2994.
25. Feuerstein, T.J., et al., *Alkali metal complexes of an enantiopure iminophosphonamide ligand with bright delayed fluorescence.* Chemical Science, 2019. **10**(18): p. 4742-4749.
26. Das, S., et al., *Manipulating Triplet Yield through Control of Symmetry-Breaking Charge Transfer.* The Journal of Physical Chemistry Letters, 2018. **9**(12): p. 3264-3270.
27. Cruz, C.D., et al., *Sulfur-Bridged Terthiophene Dimers: How Sulfur Oxidation State Controls Interchromophore Electronic Coupling.* Journal of the American Chemical Society, 2015. **137**(39): p. 12552-12564.
28. Tseng, C.-K., et al., *Copper(I)–Anilide Complex [Na(phen)3][Cu(NPh2)2]: An Intermediate in the Copper-Catalyzed N-Arylation of N-Phenylaniline.* Chemistry – A European Journal, 2011. **17**(9): p. 2716-2723.
29. Doi, R., M. Ohashi, and S. Ogoshi, *Copper-Catalyzed Reaction of Trifluoromethylketones with Aldehydes via a Copper Difluoroenolate.* Angewandte Chemie International Edition, 2016. **55**(1): p. 341-344.
30. Buttery, J.H.N., et al., *Complexes of Group 1 Salts with N,N′-Aromatic Bidentate Ligands, of Mononuclear ('Molecular') 1 : 2 Salt : Base Ratio.* Zeitschrift für anorganische und allgemeine Chemie, 2006. **632**(10-11): p. 1829-1838.
31. Marek, J., P. Kopel, and Z. Travnicek, *Tris(1,10-phenanthroline)sodium 2,4,6-trimercapto-1,3,5-triazin-1-ide.* Acta Crystallographica Section C, 2003. **59**(10): p. m399-m401.
32. Shah, S.R., et al., *Sodium, Potassium, and Lithium Complexes of Phenanthroline and Diclofenac: First Report on Anticancer Studies.* ACS Omega, 2019. **4**(25): p. 21559-21566.
33. Müller, U. and A. Noll, *Crystal structure of hexakis(acetonitrile)sodium hexachlorotantalate(V), [Na(NCCH3)6][TaCl6].* Zeitschrift für Kristallographie - New Crystal Structures, 2000. **215**(1): p. 191-192.
34. Avent, A.G., et al., *Reactions between a sodium amide Na[N(SiMe3)R1] (R1 = SiMe3, SiMe2Ph or But) and a cyanoalkane RCN (R = Ad or But).* Dalton Transactions, 2006(7): p. 919-927.
35. Clegg, W., et al., *Structurally Defined Reactions of Sodium TMP–Zincate with Nitrile Compounds: Synthesis of a Salt-Like Sodium Sodiumdizincate and Other Unexpected Ion-Pair Products.* Angewandte Chemie International Edition, 2008. **47**(4): p. 731-734.
36. Konarev, D.V., et al., *Experimental observation of C60 LUMO splitting in the C602− dianions due to the Jahn–Teller effect. Comparison with the C60˙− radical anions.* Physical Chemistry Chemical Physics, 2013. **15**(23): p. 9136-9144.
37. Berryman, O.B., et al., *Structural Criteria for the Design of Anion Receptors: The Interaction of Halides with Electron-Deficient Arenes.* Journal of the American Chemical Society, 2007. **129**(1): p. 48-58.
38. Pratt, J., et al., *Effects of Remote Ligand Substituents on the Structures, Spectroscopic, and Magnetic Properties of Two-Coordinate Transition-Metal Thiolate Complexes.* Inorganic Chemistry, 2018. **57**(11): p. 6491-6502.
39. Niemeyer, M. and P.P. Power, *Donor-Free Alkali Metal Thiolates: Synthesis and Structure of Dimeric, Trimeric, and Tetrameric Complexes with Sterically Encumbered Terphenyl Substituents.* Inorganic Chemistry, 1996. **35**(25): p. 7264-7272.
40. Englich, U., S. Chadwick, and K. Ruhlandt-Senge, *Sodium and Potassium Triisopropylbenzenethiolates: Influence on Solid-State Structure by Metal and Donor.* Inorganic Chemistry, 1998. **37**(2): p. 283-293.
41. Kumar, S., et al., *Investigation of Intramolecular Through-Space Charge-Transfer States in Donor–Acceptor Charge-Transfer Systems.* The Journal of Physical Chemistry Letters, 2021. **12**(11): p. 2820-2830.
42. Ruduss, A., et al., *Carbene–Metal Complexes As Molecular Scaffolds for Construction of through-Space Thermally Activated Delayed Fluorescence Emitters.* Inorganic Chemistry, 2022. **61**(4): p. 2174-2185.
43. Tsujimoto, H., et al., *Thermally Activated Delayed Fluorescence and Aggregation Induced Emission with Through-Space Charge Transfer.* Journal of the American Chemical Society, 2017. **139**(13): p. 4894-4900.
44. Tang, X., et al., *Highly efficient luminescence from space-confined charge-transfer emitters.* Nature Materials, 2020. **19**(12): p. 1332-1338.
45. Wu, C., et al., *Face-to-Face Orientation of Quasiplanar Donor and Acceptor Enables Highly Efficient Intramolecular Exciplex Fluorescence.* Angewandte Chemie International Edition, 2021. **60**(8): p. 3994-3998.
46. Xiong, J., et al., *Synthesis, Structures, Photophysics, and Electroluminescence of Dinuclear Carbene-Au(I)-Amide*

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Complexes Featuring the Through-Space Charge Transfer Excited State. Inorganic Chemistry, 2025. **64**(27): p. 13623-13634.
47. Pitoňák, M., et al., *Benzene Dimer: High-Level Wave Function and Density Functional Theory Calculations.* Journal of Chemical Theory and Computation, 2008. **4**(11): p. 1829-1834.
48. Scholes, G.D. and G. Rumbles, *Excitons in nanoscale systems.* Nature Materials, 2006. **5**(9): p. 683-696.
49. Nakano, T., *Synthesis, structure and function of π-stacked polymers.* Polymer Journal, 2010. **42**(2): p. 103-123.
50. Batra, A., et al., *Quantifying through-space charge transfer dynamics in π-coupled molecular systems.* Nature Communications, 2012. **3**(1): p. 1086.
51. Zhang, B., et al., *Facile Tuned TSCT-TADF in Donor-Acceptor MOF for Highly Adjustable Photonic Modules based on Heterostructures Crystals.* Angewandte Chemie International Edition, 2023. **62**(32): p. e202303262.
52. Feng, Z.-Q., et al., *Indirect Control of Donor/Acceptor Interactions for Highly Efficient Space-Confined Thermally Activated Delayed Fluorescence Emitters.* Advanced Functional Materials, 2023. **33**(6): p. 2209708.
53. Wang, X.-Q., et al., *Multi-Layer π-Stacked Molecules as Efficient Thermally Activated Delayed Fluorescence Emitters.* Angewandte Chemie International Edition, 2021. **60**(10): p. 5213-5219.
54. Bestgen, S., et al., *Intensely Photoluminescent Diamidophosphines of the Alkaline-Earth Metals, Aluminum, and Zinc.* Angewandte Chemie International Edition, 2018. **57**(43): p. 14265-14269.
55. Pinter, P., et al., *Bright luminescent lithium and magnesium carbene complexes.* Chemical Science, 2021. **12**(21): p. 7401-7410.
56. Smith, A.M., M.C. Mancini, and S. Nie, *Second window for in vivo imaging.* Nature Nanotechnology, 2009. **4**(11): p. 710-711.
57. Harrison, M.T., et al., *Colloidal nanocrystals for telecommunications. Complete coverage of the low-loss fiber windows by mercury telluride quantum dot.* Pure and Applied Chemistry, 2000. **72**(1-2): p. 295-307.
58. Caspar, J.V., et al., *Application of the energy gap law to the decay of charge-transfer excited states.* Journal of the American Chemical Society, 1982. **104**(2): p. 630-632.
59. Reinhoffer, C., et al., *Relaxation dynamics of the optically driven nonequilibrium states in the electron- and hole-doped topological-insulator materials (Bi1-xSbx)2Te3.* Physical Review Materials, 2020. **4**(12): p. 124201.
60. Zhu, C., et al., *Ultrafast dynamics of optically excited charge carriers in the room-temperature antiferromagnetic semiconductor α—MnTe.* Physical Review Materials, 2023. **7**(5): p. 054601.
61. Fulmer, G.R., et al., *NMR Chemical Shifts of Trace Impurities: Common Laboratory Solvents, Organics, and Gases in Deuterated Solvents Relevant to the Organometallic Chemist.* Organometallics, 2010. **29**(9): p. 2176-2179.
62. Pazderski, L., *15N NMR coordination shifts in Pd(II), Pt(II), Au(III), Co(III), Rh(III), Ir(III), Pd(IV), and Pt(IV) complexes with pyridine, 2,2′-bipyridine, 1,10-phenanthroline, quinoline, isoquinoline, 2,2′-biquinoline, 2,2′:6′, 2′-terpyridine and their alkyl or aryl derivatives.* Magnetic Resonance in Chemistry, 2008. **46**(S1): p. S3-S15.
63. Lüdtke, N., et al., *Revisiting Ligand-to-Ligand Charge Transfer Phosphorescence Emission from Zinc(II) Diimine Bis-Thiolate Complexes: It is Actually Thermally Activated Delayed Fluorescence.* ChemPhotoChem, 2023. **7**(1): p. e202200142.
64. Truesdell, K.A. and G.A. Crosby, *Observation of a novel low-lying excited state in zinc(II) complexes.* Journal of the American Chemical Society, 1985. **107**(6): p. 1787-1788.
65. Singh, K., S.J. Staig, and J.D. Weaver, *Facile Synthesis of Z-Alkenes via Uphill Catalysis.* Journal of the American Chemical Society, 2014. **136**(14): p. 5275-5278.
66. Siau, W.-Y., Y. Zhang, and Y. Zhao, *Stereoselective Synthesis of Z-Alkenes*, in *Stereoselective Alkene Synthesis*, J. Wang, Editor. 2012, Springer Berlin Heidelberg: Berlin, Heidelberg. p. 33-58.
67. Strieth-Kalthoff, F. and F. Glorius, *Triplet Energy Transfer Photocatalysis: Unlocking the Next Level.* Chem, 2020. **6**(8): p. 1888-1903.
68. Strieth-Kalthoff, F., et al., *Energy transfer catalysis mediated by visible light: principles, applications, directions.* Chemical Society Reviews, 2018. **47**(19): p. 7190-7202.
69. Schmitz, M., et al., *Efficient Energy and Electron Transfer Photocatalysis with a Coulombic Dyad.* Journal of the American Chemical Society, 2024. **146**(37): p. 25799-25812.
70. Bryden, M.A., et al., *Lessons learnt in photocatalysis – the influence of solvent polarity and the photostability of the photocatalyst.* Chemical Science, 2024. **15**(10): p. 3741-3757.

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DOI: 10.1039/D5SC08432F

Data availability

CCDC 2288161 and 2288162 contain the supplementary crystallographic data for this paper. Supplementary information: synthesis details, characterisation data for compounds including NMR spectroscopy, mass spectrometry, single crystal X-ray diffraction data, theoretical studies, and photophysical data.

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