RESEARCH ARTICLE

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Radical cations and dications of bis[1]benzothieno [1,4]thiazine isomers†

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Bis[1]benzothieno[1,4]thiazines (BBTT) are particularly electron-rich *S,N*-heteropentacenes and their radical cations and dications can be relevant intermediates in charge transport materials. All three regio-isomers of *N-p*-fluorophenyl-BBTT (*syn–syn*, *syn–anti*, and *anti–anti*) were studied. A reliable preparation of radical cations and dications using antimony pentachloride as an oxidant gives deeply colored salts. The electronic structure of the radical cations was assessed by EPR spectroscopy, whereas dicationic structures were characterized by NMR spectroscopy. In addition, a deeper insight into the elec-tronic structure was experimentally and computationally obtained by UV/Vis spectroscopy as well as UV/ Vis spectroelectrochemistry and DFT and TDDFT calculations. BBTTs can be considered as highly polariz-able donor π-systems with significant charge transfer contributions in neutral, cationic and dicationic state.
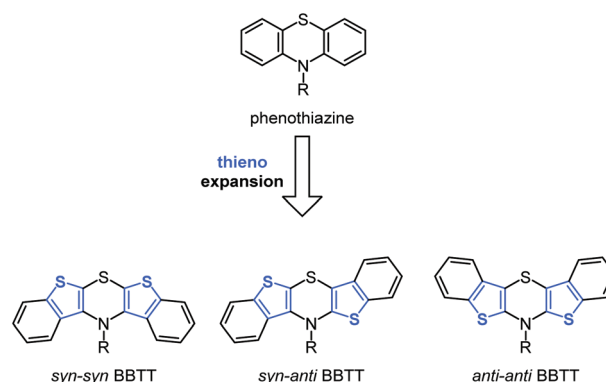
Introduction

Molecular electronics based upon organic molecules has become increasingly important over the past decade and the quest for tailor-made functional small molecules is an ongoing task for chemistry.[1,2] Since the efficiency of electronic devices correlates with electronic properties of the constituting molecules, a rational design of novel molecule based semiconductors commences with high lying HOMO levels, which in turn correspond to low oxidation potentials.[3–8]

Electron-rich heterocycles, such as phenothiazine, are pro-minent structural motifs used in electroactive materials for batteries,[9–11] OLEDs,[12–14] and DSSCs.[15–19] They possess low oxidation potentials with fully chemically reversible one-elec-tron oxidations, low band gaps, and luminescent properties.[20–22] In addition to their convenient accessibility, their intrinsic properties are easily fine tunable by synthetic functionalization.[22,23] Furthermore, the remarkable stability of their radical cations is crucial for providing efficient charge transfer as hole conductors.[24,25] Recently, we successfully proposed the two-fold thieno expansion of phenothiazine to

bis[1]benzothieno[1,4]thiazines (BBTT) that can be considered to be novel electron-enriched phenothiazine congeners (Scheme 1).[26]

The resulting compounds likewise possess discretely adjus-table properties that depend on the mode of anellation, as for dithienothiazines,[27] as well as on the substitution pattern. Different steric hindrance unequivocally causes the absence of measurable luminescence in *syn–syn* BBTTs (bis[1]benzo-thieno[2,3-*b*:3′,2′-*e*][1,4]thiazines), whereas inverse anellation of benzo[*b*]thiophene units leads to luminescence for *syn–anti* BBTTs (bis[1]benzothieno[2,3-*b*:2′,3′-*e*][1,4] thiazines) peaking for *anti–anti* BBTTs (bis[1]benzo-thieno[3,2-*b*:2′,3′-*e*][1,4]thia-zines) with fluorescence quantum yields drastically increased
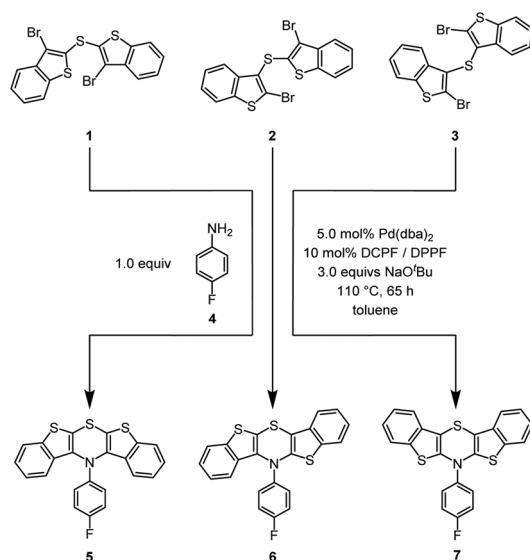
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Scheme 1 Regioisomeric BBTTs resulting by formal thieno expansion of phenothiazine.

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to 45%.[26–30] Furthermore, *anti–anti* BBTTs represent the first known fused thiazines with antiaromatic character in the crystalline solid state. Likewise, being identified as Weitz-type-redox systems,[31] BBTTs are outscoring their progenitors, *i.e.* phenothiazines, by significantly lower oxidation potentials with reversible two step oxidations, forming radical cations and dications.[26,28] Herein, we report the first synthesis of BBTTs' radical cations and dications by chemical and electrochemical oxidation as well as the detailed investigation of structural and electronic properties.
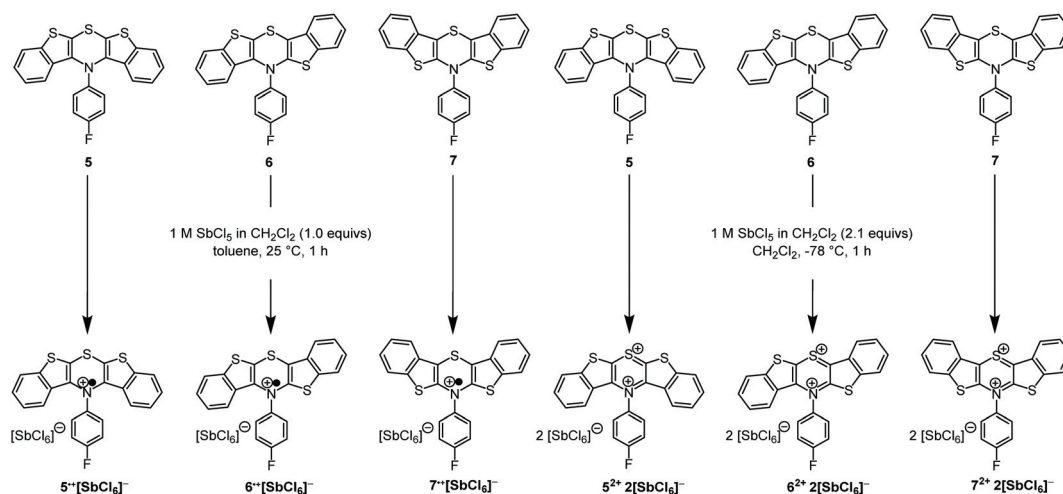
Results and discussion

Synthesis and ground state structure

N-p-Fluorophenyl-BBTTs were prepared according to our previously published procedure[28] by twofold Buchwald–Hartwig amination[32–34] (Scheme 2).

Starting from respective dibrominated dibenzo[*b*]thienyl sulfides **1**, **2**, or **3**[28,35–37] and *p*-fluoroaniline (**4**), all regioisomeric BBTTs **5–7** were reliably synthesized. The original syntheses were upscaled by a factor of ten and even upon reducing catalyst loading the yields for compounds **5** (49%), **6** (45%), and **7** (67%) were maintained. Fluorinated compounds were chosen due to their synthetic accessibility of the neutral BBTT, as well as the opportunity to monitor conversion and easily identify the products by ^{19}F NMR spectroscopy.

As supported previously by CV measurements, the BBTTs' radical cations (**5**: $E_0^{0/1+} = 0.21$ V, **6**: $E_0^{0/1+} = 0.05$ V, **7**: $E_0^{0/1+} = 0.01$ V; against ferrocene $E_0^{0/1+} = 0.00$ V) and dications (**5**: $E_0^{1+/2+} = 0.68$ V, **6**: $E_0^{1+/2+} = 0.72$ V, **7**: $E_0^{1+/2+} = 0.75$ V; against ferrocene $E_0^{0/1+} = 0.00$ V) can be readily generated in solution and they are stable over short periods.[26,28] In addition, the radical cations reveal high semiquinone formation constants K_{SEM} (**5**: 9.2×10^7, **6**: 2.3×10^{11}, **7**: 3.5×10^{12})[28] that claim remarkably stable radical cations against disproportionation into reduced BBTT and dication.[28,31] Consequently, we attempted the preparation of the oxidized species of all three BBTTs. Analogously to dithieno[1,4]thiazine (DTT)[27,30,38–41] chemical oxidation was carried out with antimony pentachloride as oxidant (Scheme 3).[41] By choice of solvent, temperature and equivalents of antimony pentachloride the oxidation stage can be reliably controlled.

BBTTs **5–7** are soluble in toluene and dichloromethane, whereas the corresponding hexachloroantimonate salts of the radical cations **5**$^{•+}$, **6**$^{•+}$, and **7**$^{•+}$ precipitated from toluene and thereby removed from the reaction equilibrium. Additionally, over-oxidation was prevented by a stoichiometric deficiency of

Scheme 2 Selective synthesis of regioisomeric *N-p*-fluorophenyl-BBTTs **5**, **6**, and **7** prepared by cyclizing Buchwald–Hartwig amination (for the synthesis of compounds **5** and **6** DCPF (1,1′-bis(dicyclohexylphosphano)ferrocene) was employed as a ligand, and for compound **7** DPPF (1,1′-bis(diphenylphosphano)ferrocene) was used as a ligand).[28]

Scheme 3 Controlled preparation of oxidized BBTT-compounds **5**$^{•+}$, **6**$^{•+}$, **7**$^{•+}$, and **5**$^{2+}$, **6**$^{2+}$, **7**$^{2+}$ as hexachloroantimonates using antimony pentachloride as an oxidant.

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oxidant. If dichloromethane was used as a solvent, the hexachloroantimonate salts of the radical cations remained soluble, forming dication salts upon increasing the amount of antimony pentachloride. The desired oxidation products precipitated from the respective reaction mixture.

For assessing the electronic structure, the molecules' geometries were calculated on the DFT level of theory using the uB3LYP functional[42,43] and the 6-311G** basis set[44,45] with SCRF (IEFPCM, CH_2Cl_2)[46–48] as implemented in the Gaussian 09 program package.[49] Indeed the BBTT units are entirely planarized with a perpendicular orientation of the N-p-fluorophenyl substituents for all three radical cations $5^{•+}$, $6^{•+}$, and $7^{•+}$. Depending on the anellation mode, different spin density distributions were found, however, with highest spin density located on the nitrogen atoms (Mulliken atomic spin density at N-atom: $5^{•+}$: 0.284, $6^{•+}$: 0.248, $7^{•+}$: 0.228). The highest delocalization of spin density is anticipated for the *anti–anti* derivative $7^{•+}$, whereas the lowest delocalization is predicted for the *syn–syn* derivative $5^{•+}$ (Fig. 1).

The radical nature of the obtained salts $5^{•+}$, $6^{•+}$, and $7^{•+}$ was proven by X-band EPR spectroscopy and by comparison with the simulated spectra (Fig. 2).
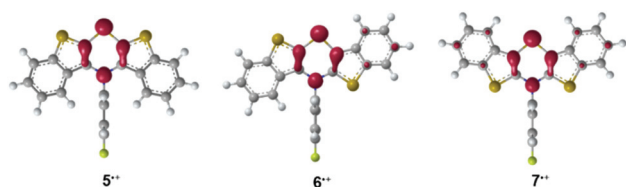

Fig. 1 DFT calculated spin density distribution plots of radical cations $5^{•+}$, $6^{•+}$, and $7^{•+}$ (uB3LYP/6-311++G**, IEFPCM CH_2Cl_2, isosurface value = 0.06 a.u.).
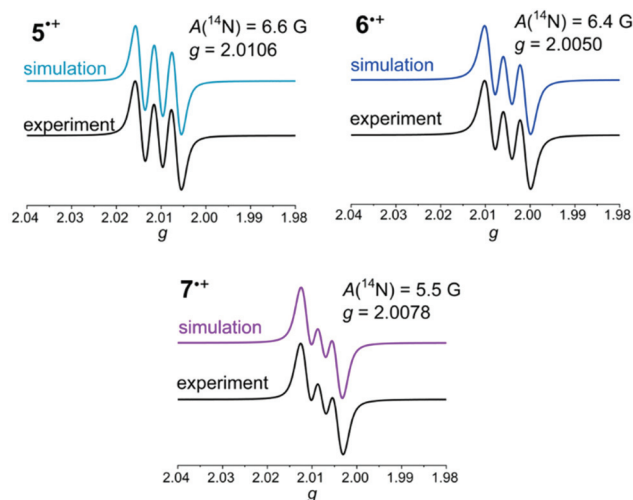

Fig. 2 Simulated and experimental EPR spectra of hexachloroantimonate salts of $5^{•+}$, $6^{•+}$, and $7^{•+}$ (recorded in THF, T = 298 K, g-factor referenced to external Mn^{2+} in ZnS).

Spectra with hyperfine coupling patterns of three equidistant lines reflect the coupling of the unpaired electron with the nitrogen nucleus in each case. Furthermore, the range of the g-factors ($5^{•+}$: 2.0106, $6^{•+}$: 2.0050, $7^{•+}$: 2.0078) indicates the predominant localization of the radicals on the nitrogen atoms.[41,50] The Mulliken atomic spin density on the nitrogen atoms decreases from *syn–syn* over *syn–anti* to *anti–anti* derivative, which is in accordance with the descending order of hyperfine coupling constants. Hence, an increasing spin delocalization across the BBTT units for *anti*-anellated benzo[b]thiophene wings can be deduced. The stronger delocalization matches with an increased stability of the corresponding radical cation $7^{•+}$, which is consistent with the experimentally determined semiquinone formation constants K_{SEM}, which also give largest values for *anti–anti* radical cation $7^{•+}$.

The delocalization of spin density of the radical cations is unequivocally supported by the calculated Wiberg bond orders (Fig. 3).[51]

The bond orders of the systems should gradually converge upon oxidation due to increasing orbital overlap by planarization resulting in an increased resonance stabilization. All three regioisomers show the most significant changes in bond orders in the 1,4-thiazine core, where CN and CS bond orders increase while CC bond orders decrease. This effect is most pronounced for the *syn–syn* regioisomer $5/5^{•+}$ while the *anti–anti* isomer $7/7^{•+}$ is most marginally affected. However, *anti*-fused benzo[b]-thiophene wings result in higher bond order changes on the fused thiophene in turn. This also confirms the previously observed increased spin delocalization upon introduction of *anti*-anellated wings. Nevertheless, even with *anti*-anellation, the spin density on the benzo ring is not high enough for causing observable coupling with the protons' nuclear spins. Although the calculated spin density distributions (Fig. 1) suggest this conclusion, no hyperfine coupling patterns of the protons are experimentally found.

For hexachloroantimonate salts of dications 5^{2+}, 6^{2+}, and 7^{2+}, DFT calculations claim formation of singlet dications being thermodynamically favored over triplet dications by energy differences of −92 kJ mol^{-1} (5^{2+}), −89 kJ mol^{-1} (6^{2+}), and −90 kJ mol^{-1} (7^{2+}). Consequently, EPR-silent but NMR-active target compounds should be present. NMR spectra of

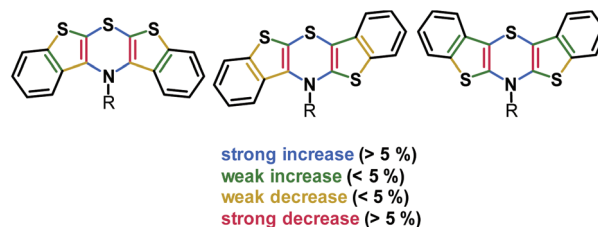**Fig. 3** Relative change of Wiberg bond orders $\Delta(BO_{X^{•+}}-BO_X)$ from neutral precursors 5, 6, and 7 to radical cations $5^{•+}$, $6^{•+}$, and $7^{•+}$ in the BBTT core (R = p-fluorophenyl; uB3LYP/6-311G**) (A detailed list of bond orders is given in the ESI†).
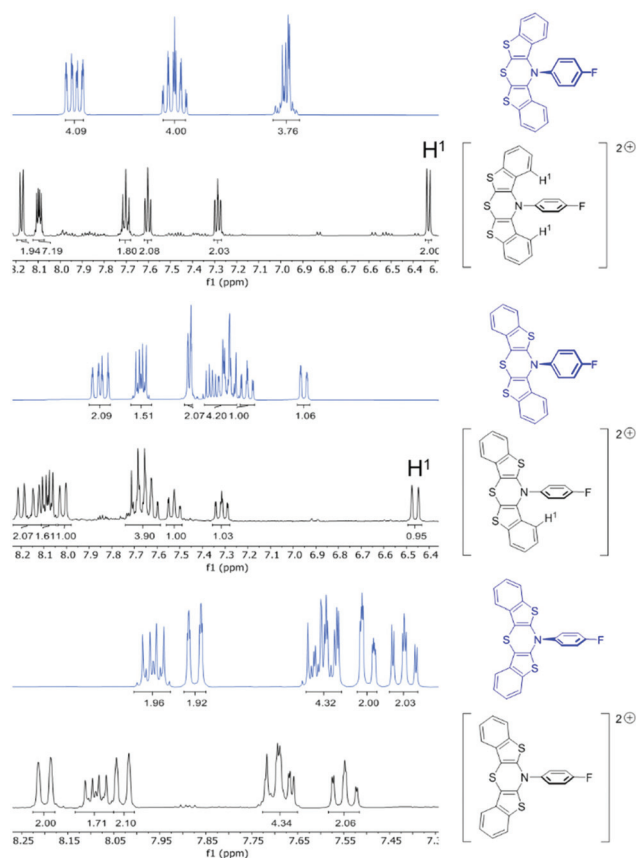
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Fig. 4 Sections of the ^{1}H NMR spectra of the neutral compounds **5**, **6**, and **7** (blue) and of the bis(hexachloro-antimonate) salts of **5^{2+}**, **6^{2+}**, and **7^{2+}** (black) (300 MHz, acetone-d_6, T = 298 K).

the bis(hexachloroantimonate) salts of **5^{2+}**, **6^{2+}**, and **7^{2+}** were recorded and compared to the spectra of the native compounds **5**, **6**, and **7**, thus, confirming the formation of diamagnetic dications, *i.e.* in their singlet ground states (Fig. 4).

As expected, the majority of the multiplets shift towards lower field, underlining the electron-deficient dicationic nature of the compounds. Due to structural differences of the regioisomers, parts of the signals appear shielded in high field. DFT-calculated geometry optimizations show planar BBTT units with orthogonally twisted *N-p*-fluorophenyl substi-

tuents, in analogy to the radical cations. *syn*-Anellated wings place benzo protons into the anisotropy cone of the *N*-aryl substituent's diamagnetic ring current. Thereby, their signals are shifted to high field.

Photophysical properties and spectroelectrochemistry

For further insight into electronic transitions of oxidized species, *i.e.* radical cations and dications, spectroelectrochemical measurements were performed in an OTTLE (Optically Transparent Thin Layer Electrochemical) cell, monitoring the *in situ* formation of radical cations and dications from the neutral precursors **5**, **6**, and **7** (Table 1). Hence, spectroelectrochemical UV/Vis absorption spectra were recorded by steadily increasing the applied potential in steps of 0.1 V.

Initially, UV/Vis spectra of all three parent regioisomers **5**, **6**, and **7** were recorded showing multiple absorptions bands in the UV region with the longest wavelength maxima at 309 nm (**5**) and 311 nm (**6**). For compound **7** the longest wavelength maximum is found in the visible at 424 nm (Fig. 5).

All oxidation processes proceed without formation of intermediates, as supported by the occurrence of isosbestic points. Moreover, the oxidations from the neutral BBTTs to the radical cations reflect the largest changes in the UV/Vis spectra and consequently in the electronic structure (*vide infra*). This originates from major geometry changes from butterfly structures
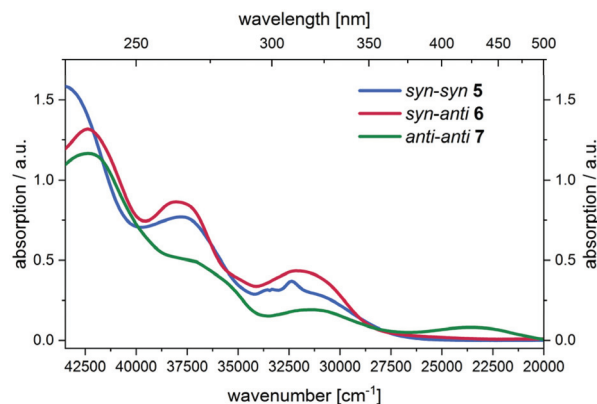

Fig. 5 UV/Vis spectra of compounds **5**, **6**, and **7** in the spectroelectrochemical cell at zero potential (CH$_2$Cl$_2$, T = 293 K).

Table 1 UV/Vis spectroelectrochemical data of *N-p*-fluorophenyl-BBTTs **5**, **6**, and **7**, corresponding radical cations **5$^{\bullet+}$**, **6$^{\bullet+}$**, and **7$^{\bullet+}$**, and dications **5^{2+}**, **6^{2+}**, and **7^{2+}** (CH$_2$Cl$_2$, T = 293 K)

Compound		Absorption maxima λ_{max} [nm]		Isosbestic points [nm]
syn–syn	**5**	230, 265, 300, 309		
	5$^{\bullet+}$	226, 234, 258, 284, 295 (sh), 311, 328, 382, 427, 494, 680	**5→5$^{\bullet+}$**	243, 276, 342, 357
	5^{2+}	227, 233, 272, 346, 494, 526, 672	**5$^{\bullet+}$→5^{2+}**	266, 279, 320, 372, 401
syn–anti	**6**	236, 263, 311		
	6$^{\bullet+}$	228, 262, 285, 315(sh), 328, 360 (sh), 377, 398, 566 (sh), 617 (sh), 694	**6→6$^{\bullet+}$**	271, 317, 351
	6^{2+}	224, 236 (sh), 253, 324, 349 (sh), 395, 595, 709 (sh)	**6$^{\bullet+}$→6^{2+}**	249, 303, 368, 418, 649, 763
anti–anti	**7**	236, 268 (sh), 317, 424		
	7$^{\bullet+}$	229, 259, 276, 293, 308, 318, 352, 369, 450 (sh), 490, 502, 534, 626 (sh), 717	**7→7$^{\bullet+}$**	255, 264, 275, 326, 361, 374, 459
	7^{2+}	223, 255, 271, 337, 561, 657	**7$^{\bullet+}$→7^{2+}**	316, 444, 542, 697

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Fig. 6 From left to right: photographs of the solutions of parent compounds **5**, **6**, and **7** in dichloromethane (c = 5.5 × 10^{-4} M, left), hexachloroantimonate salts of **5$^{•+}$**, **6$^{•+}$**, and **7$^{•+}$** in dichloromethane (c = 5.5 × 10^{-4} M, center) and of **5^{2+}**, **6^{2+}**, and **7^{2+}** in nitromethane (c = 5.5 × 10^{-4} M, right).

By gradual increase of the potential the formation of radical cations is accomplished, revealing significantly bathochromically shifted absorption bands (Fig. 7).

Therefore, compound **5$^{•+}$** shows an intense maximum at 680 nm. A strongly broadened absorption band ranging from 450 to 850 nm with shoulders at 566 and 617 nm in conjunction with a maximum at 694 nm is found for radical cation **6$^{•+}$**. The two prominent absorption maxima for oxidation product **7$^{•+}$** appear at 534 and 717 nm.

Further increase of the potential allows monitoring of the formation of dications (Fig. 8). For dication **5^{2+}**, the longest wavelength absorption maximum at 680 nm shifts slightly hypsochromically, resulting in a band at 672 nm while simultaneously a second band of lower intensity appears at 526 nm. In addition, the most intense band appears at 346 nm. A com-
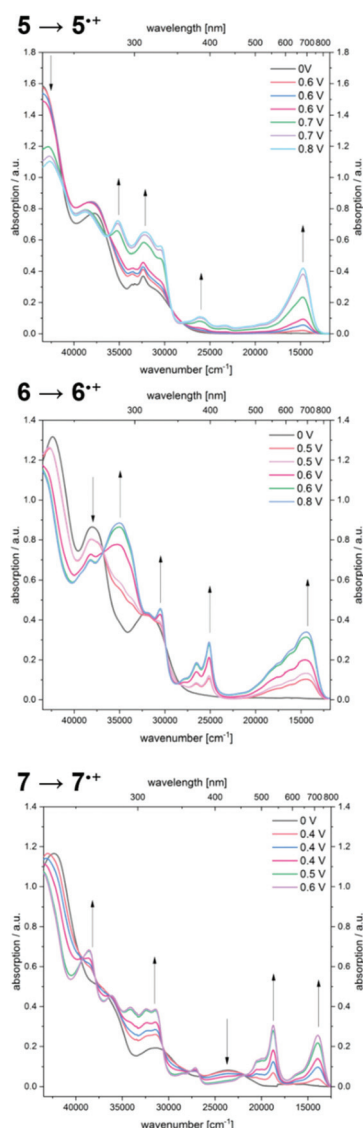

Fig. 7 UV/Vis spectroelectrochemical measurements of compounds **5**, **6**, and **7** forming the corresponding radical cations **5$^{•+}$**, **6$^{•+}$**, and **7$^{•+}$** (CH$_2$Cl$_2$, T = 293 K; electrolyte 0.1 M [^{n}Bu$_4$N][PF$_6$]; Pt working electrode, Pt counter electrode and Ag/AgCl pseudo reference electrode).
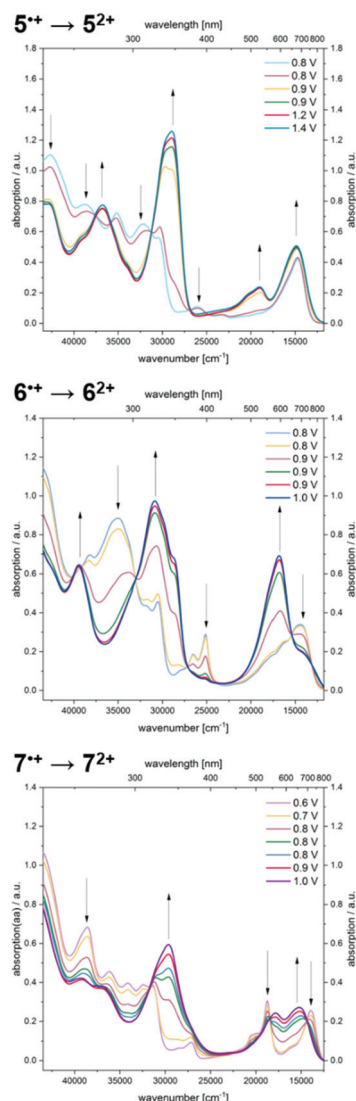
Fig. 8 UV/Vis spectroelectrochemical measurements of radical cations **5$^{•+}$**, **6$^{•+}$**, and **7$^{•+}$** forming the corresponding dications **5^{2+}**, **6^{2+}**, and **7^{2+}** (CH$_2$Cl$_2$, T = 293 K; electrolyte 0.1 M [^{n}Bu$_4$N][PF$_6$]; Pt working electrode, Pt counter electrode and Ag/AgCl pseudo reference electrode).

to the planar alignments of the BBTT units (*cf.* phenothiazines and DTT).[38,52,53] Consequently, oxidized compounds yielded by chemical as well as electrochemical oxidation are deeply colored (Fig. 6).

parable absorption band is found at 324 nm for dication 6^{2+}. The shoulders observed for radical cation $6^{•+}$ merge into a new maximum at 595 nm, whereas the maximum at 694 nm decreases to a shoulder at 709 nm. Upon oxidation of radical cation $7^{•+}$ to dication 7^{2+}, the two intense bathochromically shifted maxima of the radical cation result in the formation of two new maxima at 561 and 657 nm for the dication. Likewise, an intense band appears in the UV at 337 nm.

Finally, the spectra from spectroelectrochemical measurements were compared with the spectra of the synthesized hexachloroantimonate radical cation salts (Fig. 9).

The superposition shows that the salts unambiguously correspond to the spectroelectrochemically generated specimen and a good overall fit is obtained. Differences in the intensity of the absorption bands in the UV region of the spectra (230–380 nm) can be accounted to counter ion effects.[25] Likewise, the longest wavelength absorption maxima for the hexachloroantimonate salts of the radical cations are only slightly bathochromically shifted ($5^{•+}$ $SbCl_6^-$: $\Delta \tilde{\nu}$ = +171 cm^{-1}, $6^{•+}$ $SbCl_6^-$: $\Delta \tilde{\nu}$ = +305 cm^{-1}, $7^{•+}$ $SbCl_6^-$: $\Delta \tilde{\nu}$ = +173 cm^{-1}). As known for Weitz-type redox systems, the radical ions are remarkably stable due to their high semiquinone formation constants K_{SEM} (vide supra).[31] However, the dications could not be successfully measured due lacking stability under prolonged measuring conditions.

Electronic structure of the radical cations and dications

For a deeper insight into the electronic structure of the absorption behavior of neutral, radical cation and dication BBTTs TDDFT calculations[54,55] ((u)B3LYP[43,44,56]/6-311++G**, IEFPCM CH_2Cl_2) were performed.

First, the optical transitions of the experimentally observed absorption bands of the parent compounds 5, 6, and 7 were assigned (Table 2, Fig. 10).
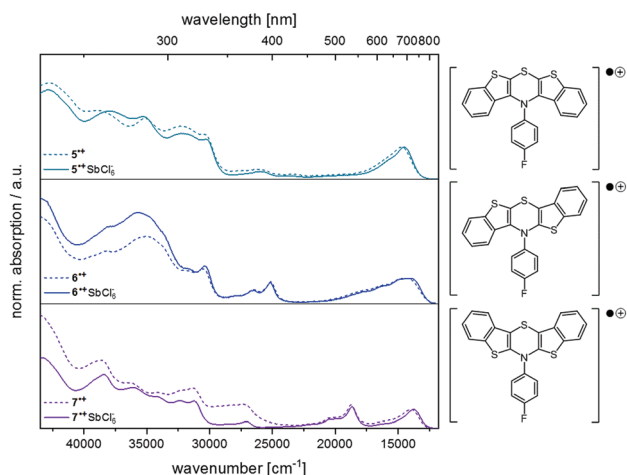
Fig. 9 Comparison of the UV/Vis spectra from measurements of the radical cation salts (solid lines) and from spectroelectrochemical measurements (dotted lines) (CH_2Cl_2, T = 293 K, for further details, see ESI†).

Table 2 Experimental and TDDFT calculated (B3LYP/6-311++G**, IEFPCM CH_2Cl_2) absorption spectra of neutral compounds 5, 6, and 7 (for further details, see ESI†)

Compound	$\lambda_{max,abs}$[a] [nm]	$\lambda_{max,calcd}$ (oscillatory strength) [nm]	Most dominant contributions
syn–syn 5	309	339 (0.075)	HOMO → LUMO (94%)
syn–anti 6	311	331 (0.143)	HOMO → LUMO+1 (68%)
anti–anti 7	424	422 (0.144)	HOMO → LUMO (95%)

[a] Recorded in CH_2Cl_2, c = 10^{-5} M, T = 293 K.
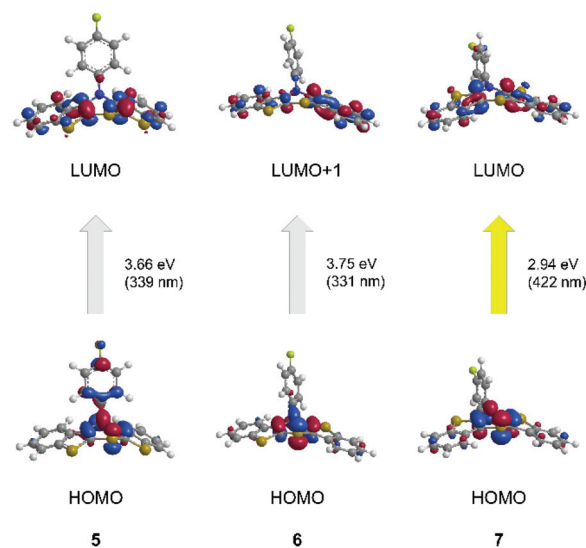
Fig. 10 TDDFT calculated frontier molecular orbital (FMO) transitions of the longest wavelength absorptions of the neutral compounds 5, 6, and 7 (B3LYP/6-311++G**, IEFPCM CH_2Cl_2, isosurface value 0.05 a.u.).

The longest wavelength absorption bands are reasonably well reproduced by transitions at 339 nm (5) and 422 nm (7). They predominantly consist of HOMO → LUMO transitions representing a charge transfer (CT) from the thiazine core to the benzo[b]thiophene wings. The calculated longest wavelength absorption band of 6 appears at 331 nm and can be predominantly assigned to a HOMO → LUMO+1 transition. Again, this transition is accompanied by a CT from the central thiazine to the benzo[b]thiophene units, with a dominant shift of coefficient density due to anti-anellation.

Likewise, for assigning the observed UV/Vis absorption of the radical cations, TDDFT calculations with the uB3LYP/6-311++G** functional were performed to prevent spin pairing and to cause separate orbitals occupation (Table 3, Fig. 11).

This leads to a singly occupied molecular orbital (SOMO). The longest wavelength absorption bands are almost accurately reproduced by HOMO−1 → SOMO transitions at wavelengths of 665 nm ($5^{•+}$), 714 nm ($6^{•+}$) and 701 nm ($7^{•+}$). For cation $7^{•+}$, the measured second absorption band at 534 nm is best represented by a HOMO−3 → SOMO transition at 538 nm, while

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Table 3 Experimental and TDDFT calculated (uB3LYP/6-311++G**, IEFPCM CH$_2$Cl$_2$) absorption spectra of the radical cations **5$^{•+}$**, **6$^{•+}$**, and **7$^{•+}$** (for further details, see ESI†)

Compound	$\lambda_{max,abs}$[a] [nm]	$\lambda_{max,calcd}$ (oscillatory strength) [nm]	Most dominant contributions
syn–syn **5$^{•+}$**	680	665 (0.191)	HOMO−1 → SOMO (100%)
syn–anti **6$^{•+}$**	694	714 (0.086)	HOMO−1 → SOMO (97%)
	617	617 (0.156)	HOMO−2 → SOMO (95%)
	566	577 (0.022)	HOMO−3 → SOMO (94%)
anti–anti **7$^{•+}$**	717	701 (0.154)	HOMO−1 → SOMO (98%)
	534	538 (0.111)	HOMO−3 → SOMO (100%)

[a] Recorded in CH$_2$Cl$_2$, $c = 10^{-5}$ M, $T = 293$ K.

Table 4 Experimental and TDDFT calculated (uB3LYP/6-311++G**, IEFPCM CH$_2$Cl$_2$) absorption spectra of the dications **5^{2+}**, **6^{2+}**, and **7^{2+}** in their singlet state (For further assignments, see ESI†)

Compound	$\lambda_{max,abs}$[a] [nm]	$\lambda_{max,calcd}$ (oscillatory strength) [nm]	Most dominant contributions
syn–syn **5^{2+}**	672	736 (0.226)	HOMO → LUMO (92%)
	526	517 (0.078)	HOMO−4 → LUMO (91%)
	346	325 (0.578)	HOMO−1 → LUMO+1 (91%)
syn–anti **6^{2+}**	650–850 (br)	817 (0.063)	HOMO → LUMO (78%)
	595	617 (0.345)	HOMO−1 → LUMO (64%)
	324	311 (0.704)	HOMO−1 → LUMO+1 (100%)
anti–anti **7^{2+}**	657	747 (0.163)	HOMO → LUMO (83%)
	561	579 (0.363)	HOMO−2 → LUMO (84%)
	337	317 (0.490)	HOMO−1 → LUMO+1 (100%)

[a] Recorded in CH$_2$Cl$_2$, $c = 10^{-5}$ M, $T = 293$ K.

Fig. 11 TDDFT calculated molecular orbital transitions of most bathochromic absorptions of radical cations **5$^{•+}$**, **6$^{•+}$**, and **7$^{•+}$** (uB3LYP/6-311++G**, IEFPCM CH$_2$Cl$_2$, isosurface value 0.05 a.u.) (For further molecular orbital transitions, see SI).

Fig. 12 TDDFT calculated molecular orbital transitions of most bathochromic absorptions of the dications **5^{2+}**, **6^{2+}**, and **7^{2+}** (uB3LYP/6-311++G**, IEFPCM CH$_2$Cl$_2$, isosurface value 0.05 a.u.) (For further molecular orbital transitions, see ESI†).
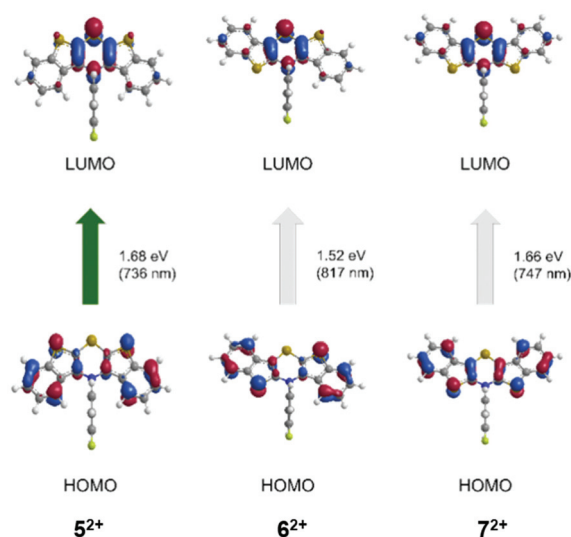
the shoulders of structure **6$^{•+}$** are best reproduced by calculated bands at 617 nm (HOMO−2 → SOMO) and 577 nm (HOMO−3 → SOMO). All these transitions are predominantly characterized by CT contributions from the anellated benzo[*b*]thiophene wings to the central thiazine core, while coefficient density on the *N*-aryl substituents is absent.

Finally, TDDFT calculations (uB3LYP/6-311++G**) were also carried out to assign the absorption bands of the dications **5^{2+}**, **6^{2+}**, and **7^{2+}** (Table 4, Fig. 12).

The longest wavelength absorption bands are best represented by calculated HOMO → LUMO transitions at wavelengths of 736 (**5^{2+}**), 709 (**6^{2+}**), and 747 (**7^{2+}**) that can be interpreted as CT transitions from the outer benzo[*b*]thiophene part to the central thiazine unit. The following experimental bands in the visible are assigned to transitions arising from HOMO−4 → LUMO at 517 nm (**5^{2+}**), HOMO−1 → LUMO at 617 nm (**6^{2+}**), and HOMO−2 → LUMO at 579 nm (**7^{2+}**).

These transitions are characterized by CT contributions from the *N*-aryl moiety to the thiazine core for structure **5^{2+}**, whereas for structure **6^{2+}** by a shift of coefficient density from the benzo[*b*]thiophene parts to the thiazine core. For structure **7^{2+}**, the transition can be accounted as a combination of both preceding coefficient density shifts.

The preceding calculations reveal that BBTT are strongly polarizable systems with large CT contributions. Accordingly, neutral BBTT are potent donors due to their electron richness, whereas the oxidized compounds are accompanied by strong acceptor properties.

Aromaticity of the dications

Furthermore, the dications' thiazine cores should inhere aromatic character due to their Weitz-type nature. Proving this,

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Table 5 Calculated NICS values of dications 5^{2+}, 6^{2+}, and 7^{2+} (B3LYP/6-311+G**). Ghost atoms respectively placed in center of the assigned rings

Ring no.	NICS (0/+1/−1)			Ring current	Aromaticity
	syn–syn 5^{2+}	*syn–anti* 6^{2+}	*anti–anti* 7^{2+}		
1a	−8.16/−9.83/−9.83	−8.26/−10.23/−10.14	−6.02/−8.20/−8.20	Diatropic	Aromatic
1b	−8.16/−9.83/−9.83	−5.99/−8.33/−8.35	−6.02/−8.20/−8.20	Diatropic	Aromatic
2a	−7.29/−6.17/−6.17	−7.38/−6.46/−6.35	−6.50/−5.96/−5.96	Diatropic	Aromatic
2b	−7.29/−6.17/−6.17	−6.39/−5.66/−5.72	−6.50/−5.96/−5.96	Diatropic	Aromatic
3	−6.94/−7.22/−7.22	−7.19/−7.43/−7.54	−6.64/−7.12/−7.12	Diatropic	Aromatic

starting with the previously mentioned optimized planar geometries, NICS (0) and NICS (1) (above (+1) and under (−1) the plane) calculations were performed. Using the GIAO protocol with B3LYP functional and 6-311+G** basis set[44,45] calculations were carried out for all five rings of the BBTT core, giving NICS values indicating diatropic ring current stating aromaticity (Table 5).

Experimental

^{1}H and ^{19}F NMR spectra of compounds **5**, **6**, **7**, 5^{2+}, 6^{2+}, and 7^{2+}, EPR spectra of **5**[•+], **6**[•+], and **7**[•+], spectroelectrochemistry of compounds **5**, **6**, and **7**, and TDDFT calculations of optimized structures **5**, **6**, **7**, **5**[•+], **6**[•+], **7**[•+], 5^{2+}, 6^{2+}, and 7^{2+} are compiled in the ESI.†

General procedure – synthesis of bis[1]benzothieno[1,4]thiazines (BBTTs)

In a dry *Schlenk* tube with a magnetic stir bar under Ar atmosphere were placed dibrominated sulfane **1**, **2**, or **3** (2.3 g, 5.0 mmol), Pd(dba)$_2$ (0.14 g, 5.0 mol%) and ligand (10 mol%). By glovebox technique sodium *tert*-butoxide (1.4 g, 15 mmol) was added into the reaction tube. Subsequently, *p*-fluoroaniline (0.47 mL, 5.0 mmol) und dry toluene (30 mL) were added by syringe before stirring the reaction mixture at 110 °C (oil bath) for 65 h. After cooling to room temp the reaction was terminated by addition of saturated aqueous sodium sulfite solution (400 mL). The aqueous phase was extracted with dichloromethane (3 × 300 mL), the combined organic phases were dried with anhydrous magnesium sulfate and the solvents were removed under vacuum. After adsorption to Celite® the residue was purified by chromatography on silica gel. After recrystallization from *n*-hexane the products **5**, **6**, or **7** were isolated as crystals. Deviations from this general procedure are individually highlighted.

N-(4-Fluorophenyl)bis[1]benzothieno[2,3-*b*:3′,2′-*e*][1,4]thiazine (*syn–syn* 5)[26]

According to the general procedure with bis(3-bromobenzo[*b*]thiophene-2-yl)sulfane (**1**) and 1,1′-bis(dicyclo-hexylphosphano)

ferrocene (0.29 g, 0.50 mmol) as a ligand and after chromatography (*n*-hexane) and recrystallization from *n*-hexane (250 mL) compound **5** (0.92 g, 49%) was obtained as colorless needles, Mp 215 °C (DSC measurement). ^{1}H NMR (300 MHz, acetone-d_6) δ 6.92–7.04 (m, 2 H), 7.42–7.55 (m, 5 H), 7.89–7.99 (m, 5 H); ^{19}F NMR (282 MHz, acetone-d_6) δ 53.42; ^{1}H NMR (600 MHz, THF-d_8) δ 6.85–6.93 (m, 4 H), 7.35–7.45 (m, 4 H), 7.80–7.88 (m, 4 H); ^{13}C{^{1}H} NMR (151 MHz, THF-d_8) δ 116.4 (CH, d, $^2J_{CF}$ = 22.7 Hz), 119.1 (CH, d, $^3J_{CF}$ = 7.7 Hz), 122.0 (CH), 124.0 (CH), 126.0 (CH), 126.0 (CH), 132.7 (C_{quat}), 136.1 (C_{quat}), 139.6 (C_{quat}), 141.0 (C_{quat}), 144.6 (C_{quat}, d, $^4J_{CF}$ = 2.4 Hz), 159.2 (C_{quat}, d, $^1J_{CF}$ = 239.4 Hz); ^{19}F NMR (565 MHz, THF-d_8) δ −124.4; HRMS(ESI) *m/z* calcd for [$C_{22}H_{12}FNS_3$]$^+$: 405.0110; Found: 405.0112; Anal. calcd for $C_{22}H_{12}FNS_3$ (405.5): C 65.16, H 2.98, N 3.45, S 23.72; Found: C 65.38, H 2.98, N 3.44, S 23.49.

N-(4-Fluorophenyl)bis[1]benzothieno[2,3-*b*:2′,3′-*e*][1,4]-thiazine (*syn–anti* 6)[26]

According to the general procedure with 2-bromo-3-((3-bromo-benzo[*b*]thiophen-2-yl)thio)benzo[*b*]thiophene (**2**) and 1,1′-bis(dicyclohexyl-phosphano)ferrocene (0.29 g, 0.50 mmol) as a ligand and after chromatography (*n*-hexane) and recrystallization from *n*-hexane (550 mL) compound **6** (0.92 g, 2.3 mmol, 45%) was obtained as yellow crystals, Mp 201 °C. ^{1}H NMR (300 MHz, acetone-d_6) δ 6.92–6.98 (m, 1 H), 7.17–7.24 (m, 1 H), 7.24–7.40 (m, 4 H), 7.44–7.48 (m, 2 H), 7.63–7.71 (m, 2 H), 7.80–7.85 (m, 1 H), 7.85–7.90 (m, 1 H); ^{19}F NMR (282 MHz, acetone-d_6) δ 61.08; ^{1}H NMR (600 MHz, THF-d_8) δ 6.93–6.96 (m, 1 H), 7.10–7.20 (m, 3 H), 7.20–7.24 (m, 1 H), 7.25–7.30 (m, 1 H), 7.35–7.43 (m, 2 H), 7.56–7.61 (m, 2 H), 7.69–7.77 (m, 2 H); ^{13}C{^{1}H} NMR (151 MHz, THF-d_8) δ 109.0 (C_{quat}), 117.3 (CH, d, $^2J_{CF}$ = 22.8 Hz), 120.6 (CH), 121.1 (C_{quat}), 121.8 (CH), 123.3 (CH), 123.6 (CH), 125.2 (CH), 125.3 (CH), 125.5 (CH), 126.1 (CH), 128.1 (CH, d, $^3J_{CF}$ = 8.4 Hz), 133.8 (C_{quat}), 135.1 (C_{quat}), 136.0 (C_{quat}), 136.3 (C_{quat}), 140.0 (C_{quat}), 143.3 (C_{quat}, d, $^4J_{CF}$ = 2.7 Hz), 145.3 (C_{quat}), 161.9 (C_{quat}, d, $^1J_{CF}$ = 245.2 Hz); ^{19}F NMR (565 MHz, THF-d_8) δ −116.7; HRMS (ESI) *m/z* calcd for [$C_{22}H_{12}FNS_3$]$^+$: 405.0110; Found: 405.0111; Anal. calcd for $C_{22}H_{12}FNS_3$ (405.5): C 65.16, H 2.98, N 3.45, S 23.72; Found: C 65.42, H 2.89, N 3.47, S 23.96.

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N-(4-Fluorophenyl)bis[1]benzothieno[3,2-*b*:2′,3′-*e*][1,4]thiazine (*anti–anti* 7)[26]

According to the general procedure with bis(2-bromobenzo[*b*]thiophen-3-yl)sulfane (**3**) and 1,1′-bis(diphenyl-hexylphosphano)ferrocene (0.29 mg, 0.50 mmol) as a ligand and after chromatography (*n*-hexane/dichloromethane/triethylamine 100 : 1 : 3) and recrystallization from *n*-hexane (450 mL) compound **7** (1.35 g, 67%) was obtained as orange crystals, Mp 177 °C (DSC measurement); ^{1}H NMR (300 MHz, acetone-d_6) δ 7.26–7.33 (m, 2 H), 7.35–7.39 (m, 2 H), 7.43–7.51 (m, 4 H), 7.72–7.77 (m, 2 H), 7.79–7.88 (m, 2 H); ^{19}F NMR (282 MHz, acetone-d_6) δ 65.73; ^{1}H NMR (600 MHz, THF-d_8) δ 7.18–7.22 (m, 2 H), 7.31–7.38 (m, 6 H), 7.58–7.62 (m, 2 H), 7.70–7.74 (m, 2 H); ^{13}C{^{1}H} NMR (151 MHz, THF-d_8) δ 100.3 (C$_{quat}$), 118.3 (d, $^2J_{CF}$ = 23.3 Hz, CH), 120.2 (CH), 122.9 (CH), 124.2 (CH), 126.2 (CH), 132.2 (d, $^3J_{CF}$ = 9.1 Hz, CH), 134.6 (C$_{quat}$), 137.3 (C$_{quat}$), 140.0 (d, $^4J_{CF}$ = 3.2 Hz, C$_{quat}$), 142.3 (C$_{quat}$), 164.1 (d, $^1J_{CF}$ = 249.0 Hz, C$_{quat}$); ^{19}F NMR (565 MHz, THF-d_8) δ −111.7; HRMS (ESI) *m/z* calcd for [C$_{22}$H$_{12}$FNS$_3$]$^+$: 405.0110; Found: 405.0110; Anal. calcd for C$_{22}$H$_{12}$FNS$_3$ (405.5): C 65.16, H 2.98, N 3.45, S 23.72; Found: C 65.31, H 2.97, N 3.43, S 23.53.

General procedure – synthesis of bis[1]benzothieno[1,4]thiazine radical cations (BBTT$^{•+}$)

In a dry *Schlenk* tube with a magnetic stir bar the *N*-4-fluorophenyl-BBTT **5**, **6**, or **7** (0.12 g, 0.30 mmol) was dissolved in dry toluene (60 mL). Under vigorous stirring, a 1 M solution of antimony pentachloride in dichloromethane (0.30 mL, 0.30 mmol) was added dropwise to the solution at room temp. After 45 min the precipitate was filtered off and washed with dry toluene (5 × 10 mL) and dry *n*-pentane (2 × 10 mL). Deviations from this general procedure are individually highlighted.

12-(4-Fluorophenyl)benzo[4,5]thieno[2,3-*b*]benzo[4,5]-thieno[3,2-*e*][1,4]thiazinyl-12-ium hexachloroantimonate (*syn–syn* 5$^{•+}$ SbCl$_6$$^−$)

According to the general procedure with *N*-(4-fluorophenyl)-bis[1]benzothieno[2,3-*b*:3′,2′-*e*][1,4]thiazine (**5**) compound **5$^{•+}$ SbCl$_6$$^−$** (0.12 g, 80%) was obtained as a dark turquoise solid. EPR (THF): *g* = 2.0106 (*A*(^{14}N) = 6.6 G); HRMS (ESI) *m/z* calcd for [C$_{22}$H$_{12}$FNS$_3$]$^+$: 405.0110; Found: 405.0106, ESI-MS *m/z* for [SbCl$_6$]$^−$: 334.7 [^{121}Sb^{35}Cl$_4$^{37}Cl$_2$/^{123}Sb^{35}Cl$_5$^{37}Cl]$^−$; IR: $\tilde{\nu}$ [cm^{-1}] = 3109 (w), 3067 (w), 1585 (w), 1548 (w), 1506 (w), 1443 (m), 1423 (w), 1358 (m), 1315 (s), 1304 (s), 1265 (m), 1233 (m), 1152 (w), 1134 (w), 1111 (w), 1063 (w), 1043 (w), 1018 (m), 856 (w), 772 (s), 754 (s), 719 (s), 653 (w), 619 (w); Anal. calcd for C$_{22}$H$_{12}$Cl$_6$NS$_3$Sb (739.98): C 35.71, H 1.63, N 1.89, S 13.00; Found: C 35.99, H 1.58, N 1.88, S 13.29.

12-(4-Fluorophenyl)benzo[4,5]thieno[2,3-*b*]benzo[4,5]thieno[2,3-*e*][1,4]thiazinyl-12-ium hexachloroantimonate (*syn–anti* 6$^{•+}$ SbCl$_6$$^−$)

According to the general procedure with *N*-(4-fluorophenyl)bis[1]benzothieno[2,3-*b*:2′,3′-*e*][1,4]thiazine (**6**) compound **6$^{•+}$ SbCl$_6$$^−$** (0.10 g, 71%) was obtained as an ink blue solid. EPR

(THF): *g* = 2.0050 (*A*(^{14}N) = 6.4 G); HRMS (ESI) *m/z* calcd for [C$_{22}$H$_{12}$FNS$_3$]$^+$: 405.0110; Found: 405.0114; ESI-MS *m/z* for [SbCl$_6$]$^−$: 334.7 [^{121}Sb^{35}Cl$_4$^{37}Cl$_2$/^{123}Sb^{35}Cl$_5$^{37}Cl]$^−$; IR: $\tilde{\nu}$ [cm^{-1}] = 2972 (w), 2901 (w), 1601 (w), 1584 (w), 1520 (w), 1506 (m), 1445 (m), 1427 (w), 1368 (w), 1335 (s), 1315 (s), 1291 (w), 1256 (m), 1240 (m), 1236 (w), 1221 (w), 1157 (m), 1111 (w), 1059 (w), 1029 (w), 980 (w), 843 (m), 756 (s), 727 (m), 717 (m), 632 (w), 610 (m), 602 (s); Anal. calcd for C$_{22}$H$_{12}$Cl$_6$NS$_3$Sb (739.98): C 35.71, H 1.63, N 1.89, S 13.00; Found: C 35.76, H 1.67, N 1.81, S 12.79.

6-(4-Fluorophenyl)benzo[4,5]thieno[3,2-*b*]benzo[4,5]thieno[2,3-*e*][1,4]thiazinyl-6-ium hexachloroantimonate (*anti–anti* 7$^{•+}$ SbCl$_6$$^−$)

According to the general procedure with *N*-(4-fluorophenyl)-bis[1]benzothieno[3,2-*b*:2′,3′-*e*][1,4]thiazine (**7**) and after additional washing of the collected precipitate with toluene/ethyl acetate (25 mL (10 : 1)) in an ultrasound bath, filtration and washing with toluene/ethyl acetate (20 mL (10 : 1), 2 mL (1 : 1)) compound **7$^{•+}$ SbCl$_6$$^−$** (0.14 g, 97%) was obtained as a dark violet solid. EPR (THF): *g* = 2.0078 (*A*(^{14}N) = 5.5 G); HRMS (ESI) *m/z* calcd for [C$_{22}$H$_{12}$FNS$_3$]$^+$: 405.0110; Found: 405.0115; ESI-MS *m/z* for [SbCl$_6$]$^−$: 334.7 [^{121}Sb^{35}Cl$_4$^{37}Cl$_2$/^{123}Sb^{35}Cl$_5$^{37}Cl]$^−$; IR: $\tilde{\nu}$ [cm^{-1}] = 3065 (w), 2990 (w), 2974 (w), 2899 (w), 1738 (m), 1585 (w), 1525 (w), 1501 (m), 1481 (m), 1433 (m), 1340 (w), 1329 (s), 1304 (m), 1250 (m), 1233 (s), 1186 (w), 1167 (w), 1153 (m), 1132 (w), 1089 (w), 1065 (w), 1045 (w), 1016 (w), 849 (m), 755 (m), 750 (s), 718 (m), 620 (w); Anal. calcd for C$_{22}$H$_{12}$Cl$_6$NS$_3$Sb (739.98): C 35.71, H 1.63, N 1.89, S 13.00; Found: C 37.07, H 1.60, N 1.82, S 12.98.

General procedure – of bis[1]benzothieno[1,4]thiazine dications (BBTT^{2+})

In a dry *Schlenk* tube with a magnetic stir bar was dissolved *N*-4-fluorophenyl-BBTT **5**, **6**, or **7** (0.12 g, 0.30 mmol) in dry dichloromethane (9 mL). Under vigorous stirring a 1 m solution of antimony pentachloride in dichloromethane (0.60 mL, 0.60 mmol) was added dropwise to the solution at −78 °C (dry ice/acetone). After 45 min the mixture was allowed to warm to room temp. The precipitate was filtered off under Ar atmosphere and washed with dry dichloromethane (5 × 10 mL). Deviations from this general procedure are individually highlighted.

12-(4-Fluorophenyl)benzo[4,5]thieno[2,3-*b*]benzo[4,5]thieno[3,2-*e*][1,4]thiazine-6,12-diium bishexachloroantimonate (*syn–syn* 5^{2+} (SbCl$_6$$^-$)$_2$)

According to the general procedure with *N*-(4-fluorophenyl)-bis[1]benzothieno[2,3-*b*:3′,2′-*e*][1,4]thiazine (**5**) and after additional washing with dry toluene (2 × 10 mL) and dry diethyl ether (2 × 10 mL) compound **5^{2+} (SbCl$_6$$^−$)$_2$** (0.10 g, 48%) was obtained as a dark blue solid. ^{1}H NMR (300 MHz, acetone-d_6) δ 6.32–6.34 (m, 1 H), 7.27–7.30 (m, 1 H), 7.59–7.62 (m, 1 H), 7.68–7.73 (m, 1 H), 8.07–8.13 (m, 1 H), 8.15–8.20 (m, 1 H); HRMS (ESI) calcd for [C$_{22}$H$_{12}$FNS$_3$]$^+$: 405.0110; Found: 405.0101; ESI-MS *m/z* for [SbCl$_6$]$^−$: 330.8 [^{121}Sb^{35}Cl$_6$]$^−$, 332.8 [^{121}Sb^{35}Cl$_5$^{37}Cl]$^−$, 334.8

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$[^{121}Sb^{35}Cl_4^{37}Cl_2]^-$, 336.8 $[^{121}Sb^{35}Cl_3^{37}Cl_3]^-$, 338.8 $[^{121}Sb^{35}Cl_2^{37}Cl_4]$, 340.8 $[^{121}Sb^{35}Cl^{37}Cl_5]$, 342.8 $[^{121}Sb^{37}Cl_6]$; IR: $\tilde{\nu}$ [cm^{-1}] = 3082 (w), 1578 (m), 1565 (w), 1501 (m), 1481 (w), 1451 (w), 1445 (w), 1431 (w), 1428 (w), 1416 (w), 1315 (w), 1285 (m), 1256 (w), 1212 (w), 1196 (s), 1184 (s), 1179 (s), 1155 (s), 1144 (m), 1125 (m), 1103 (w), 1076 (w), 1042 (s), 1036 (s), 1013 (w), 993 (w), 951 (w), 945 (w), 849 (w), 822 (w), 775 (s), 762 (s), 721 (s), 711 (m); Anal. calcd for $C_{22}H_{12}Cl_{12}FNS_3Sb_2$ (1074.4): C 24.59, H 1.13, N 1.30, S 8.95; found: C 24.82, H 1.12, N 1.34, S 8.82.

12-(4-Fluorophenyl)benzo[4,5]thieno[2,3-b]benzo[4,5]thieno[2,3-e][1,4]thiazine-6,12-diium bishexachloroantimonate (syn–anti 6^{2+} (SbCl$_6^-$)$_2$)

According to the general procedure with *N*-(4-fluorophenyl)-bis[1]benzothieno[2,3-b:2′,3′-e][1,4]thiazine (**6**) and after additional washing with dry tetrahydrofuran (2 × 2 mL) and dry pentane (2 × 10 mL) compound **6^{2+} (SbCl$_6^-$)$_2$** (0.13 g, 62%) was obtained as a blue solid. ^{1}H NMR (300 MHz, acetone-d_6) δ 6.42–6.49 (m, 1 H), 7.26–7.37 (m, 1 H), 7.49–7.56 (m, 1 H), 7.59–7.72 (m, 4 H), 7.98–8.04 (m, 1 H), 8.05–8.11 (m, 2 H), 8.11–8.16 (m, 1 H), 8.18–8.23 (m, 1 H); HRMS (ESI) calcd for $[C_{22}H_{12}FNS_3]^+$: 405.0110; Found: 405.0115; ESI-MS *m/z* for $[SbCl_6]^-$: 330.8 $[^{121}Sb^{35}Cl_6]^-$, 332.8 $[^{121}Sb^{35}Cl_5^{37}Cl]^-$, 334.8 $[^{121}Sb^{35}Cl_4^{37}Cl_2]^-$, 336.8 $[^{121}Sb^{35}Cl_3^{37}Cl_3]^-$, 338.8 $[^{121}Sb^{35}Cl_2^{37}Cl_4]^-$; IR: $\tilde{\nu}$ [cm^{-1}] = 3107 (w), 3052 (w), 1580 (m), 1573 (m), 1554 (w), 1508 (w), 1499 (m), 1485 (w), 1464 (m), 1442 (w), 1404 (w), 1346 (m), 1337 (m), 1320 (w), 1312 (w), 1275 (s), 1254 (m), 1190 (s), 1162 (m), 1150 (m), 1125 (m), 1076 (w), 1043 (m), 1012 (w), 988 (w), 845 (w), 754 (s), 756 (s), 729 (w), 716 (m), 621 (w), 604 (w); Anal. calcd for $C_{22}H_{12}Cl_{12}FNS_3Sb_2$ (1074.4): C 24.59, H 1.13, N 1.30, S 8.95; Found: C 24.88, H 1.28, N 1.27, S 9.06.

6-(4-Fluorophenyl)benzo[4,5]thieno[3,2-b]benzo[4,5]thieno[2,3-e][1,4]thiazine-6,12-diium bishexachloroantimonate (anti–anti 7^{2+} (SbCl$_6^-$)$_2$)

According to the general procedure with *N*-(4-fluorophenyl)bis[1]benzothieno[3,2-b:2′,3′-e][1,4]thiazine (**7**) after additional washing with dry pentane (2 × 10 mL) compound **7^{2+} (SbCl$_6^-$)$_2$** (0.18 g, 82%) was obtained as a dark violet solid. ^{1}H NMR (300 MHz, acetone-d_6) δ 7.52–7.58 (m, 2 H), 7.66–7.72 (m, 4 H), 8.00–8.05 (m, 2 H), 8.05–8.13 (m, 2 H), 8.18–8.22 (m, 2 H); HRMS (ESI) calcd for $[C_{22}H_{12}FNS_3]^+$: 405.0105; found: 405.0113; HRMS (ESI) calcd for $[SbCl_6]^-$: 330.7175; Found: 330.7131; IR: $\tilde{\nu}$ [cm^{-1}] = 3095 (w), 1578 (m), 1497 (m), 1454 (w), 1437 (w), 1420 (w), 1406 (w), 1325 (m), 1297 (w), 1269 (m), 1246 (w), 1211 (s), 1167 (w), 1155 (m), 1130 (m), 1092 (w), 1063 (w), 1024 (w), 993 (w), 943 (w), 866 (w), 845 (m), 756 (s), 712 (m), 670 (w), 764 (w), 635 (w), 627 (w), 620 (w); Anal. calcd for $C_{22}H_{12}Cl_{12}FNS_3Sb_2$ (1074.4): C 24.59, H 1.13, N 1.30, S 8.95; Found: C 28.38, H 1.25, N 1.54, S 10.47.

Conclusions

The electronic properties of oxidized species of all three anellative bis[1]benzothieno[1,4]thiazine (BBTT) isomers were exten-sively characterized by experimental and computational methods. All three regioisomeric *N-p*-fluorophenyl-BBTTs (*syn–syn*, *syn–anti* and *anti–anti*) were successfully transformed into the corresponding radical cation and dication salts by using antimony pentachloride as an oxidant on a preparative scale. By EPR spectroscopy the radical character was unambiguously proven and the predominant localization of the unpaired spin on the central 1,4-thiazine moiety was supported by the result-ing hyperfine coupling with the nitrogen nucleus. Moreover, experimentally determined hyperfine coupling constants and computationally determined spin density distributions and Wiberg bond orders underline that a stronger delocalization arises from *anti*-alignment of the benzo[*b*]-thiophene units. The diamagnetic singlet character as well as geometric struc-ture of the dications was assigned by NMR spectroscopy, especially supported by high-field shifted signals in diamag-netic anisotropy cones of *syn*-aligned wings due to planariza-tion of the BBTT core.

Spectroelectrochemical measurements by *in situ* generation of the oxidized species allowed characterization of the elec-tronic absorption bands. Finally, the experimentally observed absorption bands could be reproduced TDDFT calculations and assigned to the underlying molecular orbital transitions. The calculated electronic structure of the radical cations and dications underlines for all transitions a high polarizability of BBTT upon photonic excitation, as revealed by pronounced charge transfer from the benzo[*b*]thiophene moieties or the *N*-aryl unit to the central 1,4-thiazine core, stating a strong acceptor character of oxidized BBTT. Further studies on the synthesis of functionalized BBTTs focusing on enhancing their neutral antiaromatic character as well as addressing their elec-troactive and electro-optical applications are currently underway.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Notes and references

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