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Diaryldichalcogenide radical cations†‡

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One-electron oxidation of two series of diaryldichalcogenides (C_6F_5E)₂ (**13a–c**) and (2,6-Mes₂C₆H₃E)₂ (**16a–c**) was studied (E = S, Se, Te). The reaction of **13a** and **13b** with AsF₅ and SbF₅ gave rise to the formation of thermally unstable radical cations [(C₆F₅S)₂]^{•+} (**14a**) and [(C₆F₅Se)₂]^{•+} (**14b**) that were isolated as [Sb₂F₁₁][−] and [As₂F₁₁][−] salts, respectively. The reaction of **13c** with AsF₅ afforded only the product of a Te–C bond cleavage, namely the previously known dication [Te₄]²⁺ that was isolated as [AsF₆][−] salt. The reaction of (2,6-Mes₂C₆H₃E)₂ (**16a–c**) with [NO][SbF₆] provided the corresponding radical cations [(2,6-Mes₂C₆H₃E)₂]^{•+} (**17a–c**; E = S, Se, Te) in the form of thermally stable [SbF₆][−] salts in nearly quantitative yields. The electronic and structural properties of these radical cations were probed by X-ray diffraction analysis, EPR spectroscopy, and density functional theory calculations and other methods.

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Introduction

The landmark paper by Gomberg on the stable free triphenylmethyl radical initiated numerous investigations on molecules containing unpaired electrons.¹ Several kinds of persistent and stable radicals² have been described ever since, and several more general classes of (poly)radicals have been developed in recent times.³ As stated recently, “much of the

current interest in stable radicals probably arises [...] from the fundamental structure and bonding issues that naturally arise with this class of compounds”.⁴ Most often, light carbon-based or heteroatom radicals have been studied. Investigations on heavier main group radicals are fewer in number, although the last decades witnessed spectacular discoveries in the chemistry of the heavier main group elements.⁵ The design of novel synthetic strategies, particularly the use of very bulky substituents, has led to the isolation of a wide range of compounds, including main group radicals, an area which has been reviewed recently.⁶ The interest in the one-electron oxidation of diorganodichalcogenides dates back to 1868 when it was observed that (PhS)₂ (**1a**) dissolves in conc. H₂SO₄ to give intensively coloured solutions of radical cations (Scheme 1).⁷ Nowadays, it is understood that the formation of the latter begins with the one-electron oxidation of (PhS)₂ (**1a**) providing the intermediate radical cation [(PhS)₂]^{•+} (**2a**), which upon loss of another electron gives rise to the intermediate dication [(PhS)₂]²⁺ (**3**). Charge repulsion (“Coulomb explosion”) leads to the dissociation into two sulfenyl cations [PhS]⁺ (**4**), which undergo mutual electrophilic substitution of their phenyl rings

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‡ Dedicated to Professor Hubert Schmidbaur on the occasion of his 80th birthday.



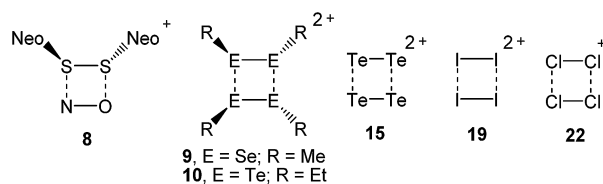
Scheme 1 Oxidation of diphenyldisulfide in conc. sulfuric acid.



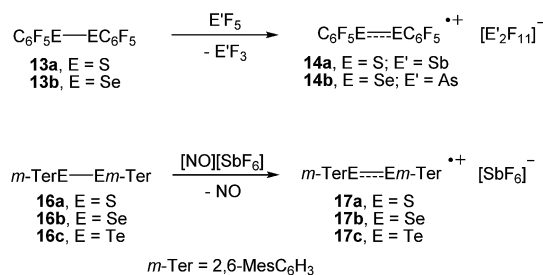
in *ortho*-position to produce thianthrene (5). Another one-electron oxidation eventually yields the thianthrene radical cation (6) and unaccounted products (Scheme 1).⁸ The one-electron oxidation of (PhS)₂ (1a) and its Se-congener (PhSe)₂ (1b) in the confined voids of the acidic pentasil zeolithe allowed the tentative characterization of the radical cations [(PhS)₂]^{•+} (2a) and [(PhSe)₂]^{•+} (2b) by EPR spectroscopy.⁹ Previous attempts to prepare a persistent dialkyldisulfide radical cation using (NeoS)₂ (7; Neo = neopentyl) and nitrosyl triflate [NO][O₃SCF₃] afforded a dialkyldisulfide nitrosonium adduct [(NeoS)₂·NO]⁺ (8) comprising a four-membered ring structure (Scheme 2).¹⁰ The related diamagnetic dicationic rings [(MeSe)₄]²⁺ (9) and [(EtTe)₄]²⁺ (10) were similarly obtained by the one-electron oxidation of heavier group 16 dialkyldichalcogenides (MeSe)₂ (11) and (EtTe)₂ (12) with [NO][O₃SCF₃].^{8b, 11} Compounds 9 and 10 can be regarded as dimers of persistent radical cations, which dimerise by π*–π* interactions.¹² The dissociation energies of these rings were estimated to be in the order of magnitude of 50 kcal mol⁻¹,¹⁰ which prompted us to investigate if persistent or even stable diaryldichalcogenide radical cations [(RE)₂]^{•+} can be prepared using fluorinated (R = C₆F₅) or bulky aromatic substituents (R = 2,6-Mes₂C₆H₃).

Results and discussion

In an initial foray, we investigated the one-electron oxidation of bis(pentafluorophenyl)dichalcogenides with antimony pentafluoride and arsenic pentafluoride. The reaction of (C₆F₅S)₂ (13a) and (C₆F₅Se)₂ (13b), respectively, with SbF₅ or AsF₅ gave an immediate colour change to dark blue and dark green upon contact, which suggested that paramagnetic species had formed (Scheme 3). Unfortunately, almost all attempts to isolate crystalline products by crystallisation from SO₂ClF at low temperatures using co-solvents such as aHF, F114 and CFCl₃ were impeded by decomposition. Only for the radical cations [(C₆F₅S)₂]^{•+} (14a; counterion [Sb₂F₁₁]⁻) and [(C₆F₅Se)₂]^{•+} (14b; counterion [As₂F₁₁]⁻) crystalline salts were obtained. However, even those crystals showed a limited thermal stability precluding any detailed spectroscopic characterisation. It is noted that the reaction of (C₆F₅S)₂ (13a) with AsF₅ in liquid SO₂ was studied previously at room temperature and provided [(C₆F₅S)₂SC₆F₅][AsF₆].¹³ The reaction of (C₆F₅Te)₂ (13c) with SbF₅ or AsF₅ under similar conditions gave no indication for the formation of radicals. The isolation of small amounts of the previously known [Te₄]²⁺ (15; counterion [AsF₆]⁻)¹⁴ provided evidence that Te–C bonds were cleaved.



Scheme 2 π*–π*-bonded four-membered rings.



Scheme 3 Synthesis of the radical cations 14a–b and 17a–c.

The oxidation of the bulky bis(*m*-terphenyl)dichalcogenides were studied by cyclic voltammetry first. At a stationary Pt electrode, electrochemical oxidation of 16a–c in CH₂Cl₂/0.1 M [*n*-Bu₄N][BF₄] within the potential sweep range 0 < *E* < 1.5 V is characterized by a one-electron quasi-reversible peak ($E_p^{Ox} - E_p^{Red} = 0.07 - 0.08$ V, $E_p^{Ox} - E_{p/2}^{Ox} = 0.06$ V, $J_p^{Red}/J_p^{Ox} \approx 0.8 - 0.9$). The observed quasi-reversibility of the peak indicates relative stability of the radical cations. Significant differences in the peak currents for 16a–c can be tentatively attributed to the differences in their diffusion coefficients. The anodic peak potentials with respect to a saturated calomel electrode of (2,6-Mes₂C₆H₃E)₂ decrease from $E_p^{Ox} = 1.22$ V (16a, E = S) over 1.09 V (16b, E = Se) to 0.79 V (16c, E = Te), respectively, which suggested nitrosonium salts to be suitable one-electron oxidizers.¹⁵ Indeed, the reaction of the bis(*m*-terphenyl)dichalcogenides (2,6-Mes₂C₆H₃E)₂ (16a, E = S; 16b, E = Se; 16c, E = Te) with [NO][SbF₆] in propionitrile provided the corresponding radical cations [(2,6-Mes₂C₆H₃E)₂]^{•+} (17a, E = S; 17b, E = Se; 17c, E = Te; counterion [SbF₆]⁻) as dark blue crystals in very high yields, which showed no signs of decomposition for several months when isolated from the mother liquor and stored under argon (Scheme 3).

The molecular structures of [(C₆F₅E)₂]^{•+} (14a, E = S; 14b, E = Se) and [(2,6-Mes₂C₆H₃E)₂]^{•+} (17a, E = S; 17b, E = Se; 17c, E = Te) are shown in Fig. 1. Selected bond parameters are collected in Table 1 together with those of the neutral parent compounds. The radical cations 14a, 14b, 17a and 17b containing S and Se atoms adopt nearly C_s symmetric conformations. The phenyl rings comprising C10–C15 are almost coplanar with the C10–E1–E2 plane pointing to delocalization of unpaired electron spin density across the aromatic π-system, whereas the phenyl rings including C20–C25 are perpendicular to the C20–E2–E1 plane (E = S, Se). The radical cation 16c containing Te atoms is centrosymmetric and possesses C₂ symmetry. Consequently, only one crystallographically independent Te atom is present. In the radical cations [(C₆F₅S)₂]^{•+} (14a) and [(2,6-Mes₂C₆H₃S)₂]^{•+} (17a) the delocalization is also reflected in the S–C bond lengths; S1–C10 (14a, 1.727(8) Å; 17a, 1.762(5) Å) is significantly shorter than S1–C20 (14a, 1.764(8); 17a, 1.799(5) Å) pointing to a quinoid structure of the coplanar phenyl ring (C10–C15). Indeed, the quinoid character of the coplanar phenyl rings of 14a (*Q* = 0.036 (26%)), 14b (*Q* = 0.012 (9%)), 17a (*Q* = 0.019 (14%)) and 17b (*Q* = 0.019 (14%)) is substantially higher than that of the perpendicular phenyl rings of 14a (*Q* = 0.024 (17%)), 14b (*Q* = 0.001 (<1%)), 17a (*Q* = 0.001 (<1%)) and 17b (*Q* = 0.002



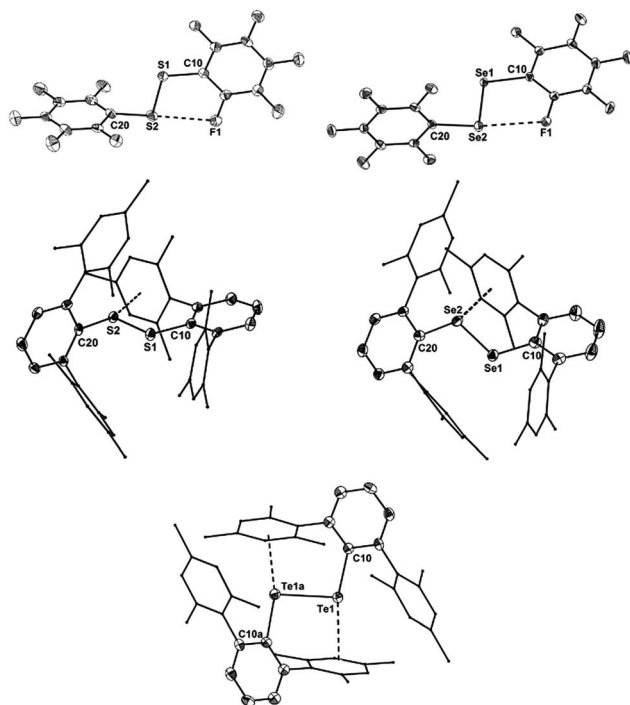


Fig. 1 Molecular structures of the radical cations **14a,b** and **17a-c**; thermal ellipsoids are set at 30% probability.

(1%).¹⁶ The quinoid character of **17c** ($Q = 0.037$ (27%)) adopts the highest value and nearly approaches that of the hexafluorobenzene radical cation $[\text{C}_6\text{F}_6]^{+\cdot}$ ($Q = 0.042$ (30%)).¹⁷

Comparison of the parent compounds with the corresponding radical cations reveals a shortening of the E–E bonds by 0.008 Å for **13a/14a**, 0.030 Å for **13b/14b**, 0.075 Å for **16a/17a**, 0.050 Å for **16b/17b** and 0.049 Å for **16c/17c** ($E = \text{Te}$), respectively. The E–E bond shortening unambiguously suggests that the bond order increased as electron density from π^* -orbitals of the chalcogens has been depleted upon oxidation. The increase of the E–E bond orders should be also reflected by an increase of the E–E stretching vibrations, however, all attempts to obtain reasonable Raman spectra failed due to the intense colour of the compounds. The most striking structural difference upon going from the parent compounds to the radical cations is the dramatic increase of the C–E–E–C torsion angles from 84.6(2)° to 175.8(4)° for **13a/14a**, 127.2(1) to 174.6(3)° for **16a/17a** ($E = \text{S}$), 75.3(1) to 178.1(1)° for **13b/14b**, 128.2(3) to 172.6(5)° for **16b/17b** ($E = \text{Se}$) and 123.1(1) to 155.5(3)° for **16c/17c** ($E = \text{Te}$), respectively. The positive charges of **17a-c** seem to be compensated by intramolecular Menshutkin interactions between chalcogen atoms and mesityl groups of the *m*-terphenyl substituents ($E\text{-Z}_\pi$ *ca.* 3 Å; Z_π = centroid of the phenyl ring).²¹ The neutral parent compounds exhibit interactions which are substantially longer ($E\text{-Z}_\pi$ *ca.* 3.4 Å). Presumably for the same reason, **14a** and **14b** possesses a short intramolecular S...F (2.712(6) Å) and Se...F (2.770(2) Å) contacts. These structural changes upon oxidation were satisfactorily reproduced by DFT calculations on two series of parent compounds, namely $(\text{PhE})_2$ (**1a-c**) and $(\text{C}_6\text{F}_5\text{E})_2$ (**13a-c**), and radical cations, namely $[(\text{PhE})_2]^{+\cdot}$ (**2a-c**) and $[(\text{C}_6\text{F}_5\text{E})_2]^{+\cdot}$ (**14a-c**) for $E = \text{S}, \text{Se}, \text{Te}$. The adiabatic ionization energies of **1a-c** (6.98–7.38 eV) are lower than those of **13a-c**

Table 1 Selected bond parameters [$^\circ$, Å] of the parent diaryldichalcogenides **13a-b**, **16a-c** and the corresponding radical cations **14a-b**, **17a-c** (italics)

$(\text{RE})_2$ $[(\text{RE})_2]^{+\cdot}$	13a (ref. 18)/ 14a , $E = \text{S}, R = \text{C}_6\text{F}_5$	13b (ref. 18)/ 14b , $E = \text{Se}, R = \text{C}_6\text{F}_5$	16a/17a , $E = \text{S}, R = m\text{-Ter}$	16b (ref. 19)/ 17b , $E = \text{Se}, R = m\text{-Ter}$	16c (ref. 20)/ 17c , $E = \text{Te}, R = m\text{-Ter}$
E–E	2.022(2) <i>2.014(3)</i>	2.319(4) <i>2.289(1)</i>	2.073(1) <i>1.998(2)</i>	2.339(2) <i>2.289(7)</i>	2.711(1) <i>2.662(1)</i>
C–E	1.785(6) <i>1.796(6)</i> <i>1.727(8)</i> <i>1.764(8)</i>	1.90(2) <i>1.92(1)</i> <i>1.876(2)</i> <i>1.904(2)</i>	1.790(2) <i>1.787(2)</i> <i>1.762(5)</i> <i>1.799(5)</i>	1.926(6) <i>1.926(7)</i> <i>1.94(1)</i> <i>1.947(8)</i>	2.144(3) <i>2.151(3)</i> <i>2.131(8)</i>
C–E–E	104.5(2) <i>106.2(2)</i> <i>96.2(3)</i> <i>107.5(3)</i>	98.7(5) <i>98.9(6)</i> <i>93.71(7)</i> <i>103.9(1)</i>	104.8(1) <i>104.4(1)</i> <i>95.4(2)</i> <i>110.4(2)</i>	102.2(3) <i>102.3(3)</i> <i>95.9(3)</i> <i>105.6(3)</i>	103.3(1) <i>103.0(1)</i> <i>98.5(2)</i>
C–E–E–C	84.6(2) <i>175.8(4)</i>	75.3(1) <i>178.1(1)</i>	127.2(1) <i>174.6(3)</i>	128.2(3) <i>172.6(5)</i>	123.1(1) <i>155.5(3)</i>
$Q_{(\text{C}10\text{-C}15)}^a$	0.036 <i>(26%)</i>	0.012 <i>(9%)</i>	0.019 <i>(14%)</i>	0.019 <i>(14%)</i>	0.037 <i>(27%)</i>
$Q_{(\text{C}20\text{-C}25)}^a$	0.024 <i>(17%)</i>	0.001 <i>(<1%)</i>	0.001 <i>(<1%)</i>	0.002 <i>(1%)</i>	—
$E\text{-Z}_\pi^b$	—	—	3.439(1) <i>2.975(2)</i>	3.452(2) <i>3.01(1)</i>	3.377(1) <i>3.481(1)</i> <i>3.161(2)</i>
$E\cdots\text{F}$	2.712(6)	2.770(2)	—	—	—

^a Quinoid character is defined as $Q_{(\text{C}10\text{-C}15)} = (d_{\text{C}10\text{-C}11} + d_{\text{C}12\text{-C}13} + d_{\text{C}13\text{-C}14} + d_{\text{C}15\text{-C}10})/4 - (d_{\text{C}11\text{-C}12} + d_{\text{C}14\text{-C}15})/2$ and $Q_{(\text{C}20\text{-C}25)} = (d_{\text{C}20\text{-C}21} + d_{\text{C}22\text{-C}23} + d_{\text{C}23\text{-C}24} + d_{\text{C}25\text{-C}20})/4 - (d_{\text{C}21\text{-C}22} + d_{\text{C}24\text{-C}25})/2$ and is 0 for a perfectly delocalized hexagonal benzene structure and 0.138 for a perfect quinoid structure where $d_{\text{C}10\text{-C}11} = d_{\text{C}12\text{-C}13} = d_{\text{C}13\text{-C}14} = d_{\text{C}15\text{-C}10} = 1.455$ Å and $d_{\text{C}20\text{-C}21} = d_{\text{C}22\text{-C}23} = 1.317$ Å. ^b Element distance to the centroid of the phenyl ring $E\text{-Z}_\pi$.



(7.64–8.16 eV) and follow the same trend as the cathodic peak potentials of series (2,6-Mes₂C₆H₃E)₂ (**16a–c**) for E = S, Se, Te. In contrast to the neutral (PhE)₂ (**1a–c**) that all have a C₂ (H₂O₂ type) structure with dihedral angles around 90°, all radical cations are calculated having much larger dihedral angles. The phenyl-substituted radical cations [(PhE)₂]^{•+} (**2a–c**), can be considered as essentially freely rotating around the E–E bonds, with a minimum energy at dihedral angles around 160° and a C₂-symmetric structure. The experimentally found C_s symmetric structure is less than 1 kcal mol⁻¹ higher in energy and corresponds to a calculated transition state (one imaginary frequency (>10 cm⁻¹)). The size of the butterfly shaped *m*-terphenyl groups easily explains why the radical cations [(2,6-Mes₂C₆H₃E)₂]^{•+} (**17a–b**; E = S, Se) have C_s symmetry, simply for reasons of sterical crowding. In the pentafluorophenyl substituted radical cations [(C₆F₅E)₂]^{•+} (**14a–c**) with E = S and Se, the experimentally observed C_s symmetry is the ground state, and the C₂ symmetric structure is about 1.5 kcal mol⁻¹ higher in energy. Interestingly, [(C₆F₅Te)₂]^{•+} (**14c**) behaves differently. Its ground state is C_{2h} symmetric, completely flat, with a dihedral angle of 180°. The C_s symmetric structure is again only <0.5 kcal mol⁻¹ higher in energy. It is unclear whether this peculiar structure is the reason why it has not been possible to isolate it. The HOMO of (C₆F₅Se)₂ (**13b**) is an admixture of components situated at the Se atoms and the π-system of the pentafluorophenyl groups, whereas in the SOMO of the corresponding radical cation [(C₆F₅Se)₂]^{•+} (**14b**), the unpaired electron is strongly distributed over the coplanar phenyl group (Fig. 2).

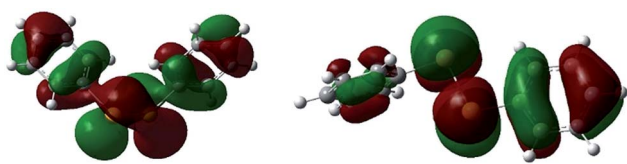


Fig. 2 HOMO (left) of C₂ symmetric (C₆F₅Se)₂ (**13b**) and SOMO (right) of C_s symmetric [(C₆F₅Se)₂]^{•+} (**14b**).

The Wiberg bond indices (WBIs) of the E–E bonds increase from about 1 for the parent compounds **1a–c** and **13a–c** to values between 1.216 and 1.315 for the [(PhE)₂]^{•+} (**2a–c**; E = S, Se, Te) series and between 1.176 and 1.282 for the [(C₆F₅E)₂]^{•+} (**14a–c**; E = S, Se, Te) series of radical cations. In the absence of electron delocalization across the coplanar phenyl rings the WBIs of the radical cations would have been expected to be close to 1.5.

The stability of **17a–c** in solution dramatically depends on the solvents and in CH₂Cl₂ on the nature of the chalcogen. Ink-blue solutions of **17a** and **17b** in CH₂Cl₂ are stable for months, whereas purple-blue **17c** decomposes within a few days. The blue colour arises from very broad absorptions in the near IR region, which stretch into the visible range. Only **17c** shows an absorption maximum (CH₂Cl₂) at λ_{max} = 583 nm in the visible range that is only slightly shifted compared to that of **16c** (553 nm) and responsible for the purple tinge. The absorption is

tentatively assigned to a n(Te) → σ* transition.²² Dark blue solutions of **17a–c** in acetonitrile and propionitrile are stable only for a few days. Notably, after some time the parent compounds **16a–c** slowly form back and eventually precipitate. Electrospray mass spectra (MeCN, positive mode) of **17a–c** show prominent mass clusters at *m/z* = 690.4, 786.3 and 882.2, which were unambiguously assigned to the radical cations on the basis of the correct isotopic patterns. The molecular conductivities (MeCN, *c* = 5 × 10⁻⁷ mol L⁻¹) of *A* = 1800, 600 and 540 Ω⁻¹ cm² mol⁻¹ confirm a high concentration of electrolytes in solutions of **17a–c**. The one-electron oxidation upon going from (2,6-Mes₂C₆H₃E)₂ (**16b**, E = Se; **16c**, E = Te) to the radical cation [(2,6-Mes₂C₆H₃E)₂]^{•+} (**17b**, E = Se; **17c**, E = Te) is well reflected by multinuclear NMR spectroscopy. The ⁷⁷Se NMR spectra show signals at δ = 426.2 (CDCl₃) for **16b** and at δ = 1362.3 (CDCl₃, 223 K) and 1362.4 (MeCN) for **17b**, respectively. The ¹²⁵Te NMR spectra exhibit signals at δ = 322.2 (CDCl₃) for **16c** and at δ = 1703.8 (CDCl₃) and 1698.7 (MeCN) for **17c**, respectively. These values are in good agreement with those calculated for (PhSe)₂ (**1b**, δ = 451.5), [(PhSe)₂]^{•+} (**2b**, δ = 1237.2 and 1493.0; average 1365.1), (PhTe)₂ (**1c**, δ = 185.1) and [(PhTe)₂]^{•+} (**1c**, δ = 1346.8 and 1778.9; average 1562.9). ¹H and ¹³C NMR spectra gave expectedly broad signals. The paramagnetism of the radical cations was unambiguously confirmed by EPR spectroscopy and SQUID magnetometry. X- and Q-band field-sweep EPR spectra measured in frozen CH₂Cl₂/THF solution (1 : 1) show that all three species have *g* values that deviate significantly from the free-electron value. All radicals exhibit a rhombic *g*-matrix with the same *g*-value ordering, inferring the electronic structure of the three radicals is very similar. The *g* anisotropy for the radical cations [(2,6-Mes₂C₆H₃E)₂]^{•+} increases in the order S (**17a**) < Se (**17b**) < Te (**17c**), as expected from the increasing spin-orbit coupling constant and for radicals where the spin density is predominately located on the central dichalcogen moiety. The EPR spectra of the [(2,6-Mes₂C₆H₃E)₂]^{•+} radical cations feature a single proton hyperfine coupling resolved at *g*₁ and *g*₂ and this coupling was fully characterized by ¹H Davies ENDOR spectra recorded at Q-band (Fig. 3, see ESI† for details). Simulation of the data yielded the hyperfine matrix (Table 2, see ESI† for details), which reveals a small but significant isotropic component of |*a*_{iso}| = 7.1 MHz (*a*_{iso} = (*a*₁ + *a*₂ + *a*₃)/3), proving a small amount of spin density delocalises onto one of the *m*-terphenyl ligands.²³ The findings were supported by spin density calculation (B3LYP/def2-TZVP) showing noticeable negative spin density on the *para*-H atom of one *m*-terphenyl ring (Fig. 4). The EPR spectra of the [(2,6-Mes₂C₆H₃Se)₂]^{•+} (**17b**) at X- (Fig. 3 top) and Q-band (Fig. 3 bottom) exhibit two distinct Se hyperfine couplings along *g*₁ (⁷⁷Se, *S* = 1/2, 7.6% abundance), consistent with a radical containing a Se–Se moiety with a small asymmetry in the electronic structure. Along *g*₂ and *g*₃, ⁷⁷Se hyperfine splittings were partially resolved and enabled an estimation of the remaining principal values of the two hyperfine interactions. Q-band Davies ENDOR was used to investigate the proton hyperfine interactions (see ESI† for details) which revealed (again) one largest resolved proton hyperfine interaction. The latter is, however, smaller than in the case of **17a**, with |*a*_{iso}| = 4.2 MHz. This result is consistent with the small





Fig. 3 Field-sweep EPR spectra for **17a** ($E = S$), **17b** ($E = Se$) and **17c** ($E = Te$) measured in frozen CH_2Cl_2/THF solution (1 : 1), along with the corresponding simulations.

Table 2 Experimental and DFT (in parentheses) EPR parameters for **17a–c**. Hyperfine values are in MHz

Parameter	Principal values
17a	
g	2.0014, 2.0115, 2.0285 (2.0022, 2.0107, 2.0213)
$A(^1H)$	-3.0, -7.2, -11.0 (-1.8, -5.8, -8.5)
17b	
g	1.9956, 2.0438, 2.1543 (2.0021, 2.0452, 2.1165)
$A(^{77}Se)$	-50, -100, 465 (-106, -116, 306)
$A(^{77}Se)$	-80, -115, 610 (-135, -140, 407)
$A(^1H)$	-1.8, -4.3, -6.4 (-1.2, -4.2, -6.1)
17c	
g	1.9542, 2.0411, 2.4566 (2.0021, 2.0876, 2.3688)
$A(^{127}Te)$	300, 350, -1000 (350, 370, -720) (352, 372, -724)
$A(^1H)$	-, -, -3.8 (-0.6, -2.6, -3.8)

asymmetry in the spin density inferred from the two inequivalent ^{77}Se hyperfine couplings. The EPR spectra of $[(2,6-Mes_2C_6H_3Te)_2]^+$ (**17c**) established principal values for the

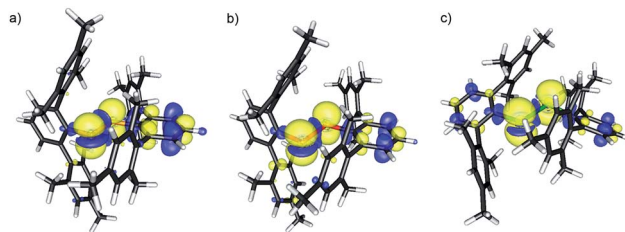


Fig. 4 Calculated spin densities for **17a–c** at contour levels of 0.0025 (yellow) and -0.0005 (blue). Note the small but noticeable negative spin density on the *para*-H atom of one terphenyl ring for **17a** and **17b**.

rhombic g -matrix (Fig. 3 and Table 2). Because of the low natural abundance of ^{125}Te ($S = 1/2$, 7% abundance), the broad EPR spectrum and small background signals of the data only allowed estimates for the hyperfine couplings and were not of sufficient quality to resolve a potential inequivalence in the two Te nuclei. Along g_1 , the hyperfine coupling is resolved with $|A(^{125}Te)| = 1000$ MHz; along g_2 and g_3 , the ^{125}Te hyperfine couplings were partially resolved and allowed the estimates shown in Table 2. Davies ENDOR was used to investigate the proton hyperfine interactions and showed that the largest proton hyperfine interaction has further decreased in comparison to **17a** and **17b**. Along g_2 , the coupling is $|A| = 3.8$ MHz, whereas for **17b**: $|A| = 6.4$ MHz, and for **17a**: $|A| = 11.0$ MHz. The experimental EPR data are very well modelled by DFT calculations (Fig. 4 and ESI† for details). The principal g -values are well reproduced as well as the two inequivalent hyperfine couplings for ^{77}Se . The DFT data are able to unambiguously assign the largest proton coupling to the *para*-H atom of the central phenyl group of one of the *m*-terphenyl substituents (Fig. 4). The trend to smaller 1H hyperfine values observed experimentally in the series **17a**, **17b**, and **17c** follows the change in the orientation of two *m*-terphenyl substituents with respect to the central dichalcogen moiety: for **17a**, the phenyl group π system of one *m*-terphenyl substituent is well orientated to allow overlap with the π -type orbitals carrying the unpaired electron on the S–S moiety and thus facilitates spin density delocalization. In contrast, the phenyl group of the second *m*-terphenyl substituent is approximately at 90° to this orientation and thus there is poor overlap with the S π -orbitals carrying the unpaired electron and less spin delocalisation. For **17c**, the π -orbitals of both phenyl rings of the *m*-terphenylsubstituent have essentially the same orientation with respect to the π -type orbitals of the Te–Te moiety carrying the unpaired electron but with relatively poor orbital overlap, resulting in an equivalent but relatively small delocalisation of the spin densities onto the two *m*-terphenyl substituents. As expected, **17b** has a structure intermediate between **17a** and **17c** and this is reflected in the experimental 1H hyperfine coupling assigned to one *para*-H of the *m*-terphenyl group. The summed chalcogen-based spin densities from DFT calculations are 0.698, 0.734 and 0.778 for **17a**, **17b**, and **17c**, respectively. These findings in conjunction with the EPR data leave no doubt that all three radical cations are characterized best as chalcogen-centred (*i.e. ca.* 70–80% spin density on the chalcogen atoms).



The effective magnetic moments (μ_{eff}) of **17a–c** at 300 K are 1.61, 1.48 and 1.41 μ_{B} , respectively, and reasonably close to the theoretically expected value of 1.744 μ_{B} , 1.7880 μ_{B} , 1.8625 μ_{B} , respectively, for a system of uncoupled paramagnetic centers with spins $S = 1/2$; the μ_{eff} decreases slightly with lowering temperature, that implies weak antiferromagnetic interactions in the solid state (see ESI† for details).

Conclusion

The stable group 16 radical cations $[(\text{RE})_2]^+$ (**14a–b**, **17a–c**; E = S, Se, Te) were prepared and fully characterized by various methods for the first time. Note that these radical cations are isoelectronic with the previously described group 15 radical anions $[(\text{RE})_2]^-$ (E = P, As, Sb; R = 2,6- $[(\text{Me}_3\text{Si})_2\text{CH}]_2$ -4- $[(\text{Me}_3\text{Si})_3\text{C}]-\text{C}_6\text{H}_2$).²⁴ Bearing in mind that the aryltellurenyl fragment RTe is isolobal with atomic iodine I, the paramagnetic $[(\text{RTe})_2]^+$ (**17c**) and the diamagnetic $[(\text{RTe})_4]^{2+}$ (**10**, R = Et)¹¹ also resemble the paramagnetic $[\text{I}_2]^+$ (**18**)²⁵ and the diamagnetic I_4^{2+} (**19**).²⁶ In the same notion, the radical cations $[(\text{RSe})_2]^+$ (**14b**, R = C_6F_5 ; **17b**, R = 2,6-Mes₂C₆H₃) and $[(\text{RS})_2]^+$ (**14a**, R = C_6F_5 ; **17a**, R = 2,6-Mes₂C₆H₃) are closely related to the known $[\text{Br}_2]^+$ (**20**)²⁷ and the elusive $[\text{Cl}_2]^+$ (**21**), which forms $[\text{Cl}_4]^+$ (**22**) with Cl₂ (Scheme 2).²⁸ The reported findings leave no doubt that the radical cations **14a–b**, **17a–c** contain chalcogen-centred odd-electron π -bonds.²⁹ It is noteworthy that recently also odd-electron S–S and Se–Se σ -bonded 1,8-bis(arylchalcogenyl)naphthalene were investigated.³⁰ In view of the fact that many odd-electron species exist as (reactive) intermediates in various chemical reactions and play an important role in bond formation and cleavage processes, alongside many applications that have been reported or envisaged for stable radicals, the successful isolation of the title compounds provide a suitable entry point for an in-depth exploration of these and related species. Accordingly, the isolation and characterization of other examples for odd-electron bonding, which is of both fundamental and practical interest, is currently under way in our laboratories.

Experimental

Synthesis of $(\text{C}_6\text{F}_5\text{S})_2\text{Sb}_2\text{F}_{11}$ (**[14a][Sb₂F₁₁]**)

SbF₅ (220 mg, 1.0 mmol) was added to solid **13a** (40 mg, 0.1 mmol) in a PFA tube. **13a** directly changed the colour to dark blue when SbF₅ reached the solid. The mixture has been kept 3 h at room temperature until the dissolving of **13a** in SbF₅ completed. The reaction mixture was then evacuated and SO₂ClF (1.0 mL) was condensed into the PFA tube at –196 °C. The mixture was allowed to slowly warm up to –80 °C using a dry ice/ethanol bath. After the melting of the solvent (1 h) completed, the reaction mixture was allowed to slowly warm up to room temperature (4 h) to ensure the complete dissolution of the reaction mixture. Afterwards the PFA tube was sealed under reduced temperature and pressure. Dark blue needles of **[14a][Sb₂F₁₁]** formed at –30 °C after several

careful cycles of cooling and warming between –50 °C and to –78 °C.

Synthesis of $(\text{C}_6\text{F}_5\text{Se})_2\text{As}_2\text{F}_{11}$ (**[14b][As₂F₁₁]**)

SO₂ClF (0.20 mL) was slowly condensed onto solid **13b** (25 mg, 0.05 mmol) in a PFA tube at –196 °C. The mixture was slowly warmed up to room temperature for complete dissolution. Then the mixture was cooled again to –196 °C and AsF₅ (0.04 mL, 0.5 mmol) was condensed into the PFA tube. The solidified mixture directly changed the colour to dark violet. Afterwards the mixture was slowly warmed up to –78 °C where the mixture melts and partial dissolution of the solid gave a deep green colour. The reaction was allowed to further warm up until reaching –30 °C after 3 h, when additional SO₂ClF (1.0 mL) was added. Afterwards the PFA tube was sealed under reduced temperature and pressure. Black needles of **[14b][As₂F₁₁]** formed after careful cooling to –78 °C over two days.

Attempts to oxidize $(\text{C}_6\text{F}_5\text{Te})_2$ (**13c**)

At –196 °C, SO₂ (0.20 mL) was condensed onto **13c** (30 mg, 0.05 mmol) in a PFA tube. After warming up to room temperature, the reaction mixture was cooled down to –196 °C a second time to add AsF₅ (0.04 mL, 0.5 mmol). The colour of the reaction changed to dark violet during the process of condensation. Afterwards the mixture was slowly warmed up to –78 °C, to remove the excess of AsF₅ under reduced pressure. After a third cooling process (–196 °C), further SO₂ (1.0 mL) was added and the reaction mixture was allowed to slowly warm up to –78 °C to give a deep violet solution. For a complete melting process of the solvent, the mixture was further warmed up to –30 °C over a period of 3 h. Afterwards F114 (0.3 mL) has been condensed onto the solidified reaction mixture at –196 °C and the PFA tube sealed afterwards. $[\text{Te}_4][\text{AsF}_6]_2$ crystallized as dark violet plates after a careful cooling process from room temperature to –30 °C over two days.

Synthesis of 2,6-Mes₂C₆H₃SH

A solution of 2,6-dimesityliodobenzene (6.50 g, 14.7 mmol) in *n*-hexane (150 mL) was treated with a 2.5 M solution of *n*-BuLi (6.60 mL, 14.7 mmol) at room temperature and stirred overnight. The volume of the suspension has been reduced to 20 mL under reduced pressure and the white solid has been filtered off. The product, 2,6-dimesitylphenyl lithium (3.33 g, 10.4 mmol), was dissolved in THF (50 mL), cooled to –78 °C and sulfur powder (0.40 g, 12.5 mmol) was cautiously added. The red suspension was stirred overnight and allowed to warm up to room temperature during that period. Hydrochloric acid (40 mL, 10%) was added to the suspension and the stirring has been extended for another 2 h. The organic phase was extracted with CHCl₃ (3 × 50 mL), filtrated and the solvent removed under reduced pressure. The crude product was recrystallized from CH₂Cl₂ to yield pale yellow crystals of 2,6-Mes₂C₆H₃SH (3.02 g, 8.72 mmol, 84%; Mp. 192–194 °C).

¹H-NMR (CDCl₃): δ = 7.27 (t, 1H, ³J = 8.1 Hz), 7.06 (d, 2H, ³J = 7.5 Hz), 7.02 (s, 4H), 3.07 (s, 1H, SH), 2.38 (s, 6H), 2.07 (s, 12H)



ppm. $^{13}\text{C}\{^1\text{H}\}$ -NMR (CDCl_3): $\delta = 138.7, 137.3, 136.2, 132.7, 128.4, 128.2, 128.0, 125.1, 21.2, 20.0$ ppm.

Synthesis of (2,6-Mes₂C₆H₃S)₂ (16a)

A solution of 2,6-Mes₂C₆H₃SH (2.00 g, 5.77 mmol) in toluene (80 mL) was treated with a solution of ethyl nitrite in ethanol (~15%, 25 mL, 40 mmol). The solution was kept at room temperature for 4 h before stirring it at 76 °C over a period of 72 h. The resulting yellow solid was filtered off and dried under vacuum. The crude product was recrystallized from CH_2Cl_2 to afford yellow crystals of **16a** (1.80 g, 2.60 mmol, 90%; Mp. >230 °C).

^1H -NMR (CDCl_3): $\delta = 7.19$ (t, 2H, $^3J = 7.4$ Hz), 6.88 (d, 4H, $^3J = 7.5$ Hz), 6.80 (8H), 2.35 (12H), 1.70 (24H) ppm. $^{13}\text{C}\{^1\text{H}\}$ -NMR (CDCl_3): $\delta = 142.9, 137.4, 136.6, 136.5, 136.1, 129.2, 127.7, 127.1, 21.1, 20.3$ ppm. Anal. calcd for C₄₈H₅₀S₂ (691.06): C, 83.43; H, 7.29. Found C, 83.29; H, 6.91.

Synthesis of 2,6-Mes₂C₆H₃SeH

A solution of 2,6-dimesityliodobenzene (6.50 g, 14.7 mmol) in *n*-hexane (150 mL) was treated with a 2.5 M solution of *n*-BuLi (6.60 mL, 14.7 mmol) at room temperature and stirred overnight. The volume of the suspension has been reduced to 20 mL under reduced pressure and the white solid has been filtered off. The product, 2,6-dimesitylphenyl lithium (3.33 g, 10.4 mmol), was dissolved in THF (50 mL), cooled to -78 °C and selenium powder (0.990 g, 12.3 mmol) was cautiously added. The black suspension was stirred overnight and allowed to warm up to room temperature during that period. Hydrochloric acid (40 mL, 5%) was added to the suspension and the stirring was extended for another 2 h. The organic phase has been extracted with CHCl_3 (3 × 50 mL), filtrated and the solvent removed under reduced pressure. The remaining solid was dissolved in THF (50 mL) and slowly added to a suspension of LiAlH₄ (0.79 g, 20.8 mmol) in THF (10 mL) at 0 °C. The suspension was stirred for 1 h, cautiously poured onto ice-water and extracted with CHCl_3 (3 × 50 mL). The crude product was recrystallized from *n*-hexane to afford yellow crystals of 2,6-Mes₂C₆H₃SeH (1.76 g, 4.47 mmol, 43%; Mp. = 222–224 °C).

^1H -NMR (CDCl_3): $\delta = 7.19$ (t, 1H, $^3J = 7.5$ Hz), 7.09 (d, 2H, $^3J = 7.4$ Hz), 7.04 (s, 4H), 2.41 (s, 6H), 2.10 (s, 12H), 1.11 (s, 1H, SeH), 1J ($^{77}\text{Se}-^1\text{H} = 63.4$ Hz) ppm. $^{13}\text{C}\{^1\text{H}\}$ -NMR (CDCl_3): $\delta = 141.1, 138.9, 137.3, 136.0, 129.1, 128.4, 128.0, 126.0, 21.2, 20.0$ ppm. ^{77}Se -NMR (CDCl_3): $\delta = 71.8$ (d, 1J ($^1\text{H}-^{77}\text{Se} = 63.4$ Hz) ppm.

Synthesis of (2,6-Mes₂C₆H₃Se)₂ (16b)

A solution of 2,6-Mes₂C₆H₃SeH (1.72 g, 4.37 mmol) was dissolved in toluene (100 mL) and treated with a solution of ethyl nitrite in ethanol (~15%, 16.2 g, 32.3 mmol). The solution was kept at room temperature for 4 h before stirring it at 82 °C over a period of 48 h. The remaining solvent was removed under reduced pressure. The product was crystallized from CHCl_3 to afford deep red prisms of **16b** (580 mg, 1.49 mmol, 34%; Mp. >230 °C).

^1H -NMR (CDCl_3): $\delta = 7.19$ (t, 2H, $^3J = 7.5$ Hz), 6.82 (d, 4H, $^3J = 7.5$ Hz), 6.74 (8H), 2.32 (12H), 1.72 (24H) ppm. $^{13}\text{C}\{^1\text{H}\}$ -NMR

(CDCl_3): $\delta = 144.1, 138.2, 136.4, 136.2, 132.8, 129.0, 127.9, 127.3, 20.9, 20.4$ ppm. ^{77}Se -NMR (CDCl_3): $\delta = 426.2$ ppm.

Synthesis of (2,6-Mes₂C₆H₃S)₂SbF₆ ([17a][SbF₆])

Solid **16a** (193 mg, 0.28 mmol) was added to a stirred solution of [NO][SbF₆] in propionitrile (15 mL). After 18 h of stirring, the solvent was removed under reduced pressure affording a deep blue solid that was recrystallized from $\text{CH}_2\text{Cl}_2/n$ -hexane (1 : 1) at room temperature to yield [17a][SbF₆] (250 mg, 0.027 mmol, 96%).

ESI MS (CH_3CN , positive mode): $m/z = 690.4$ [C₄₈H₅₀S₂]⁺ for **17a**. Molar conductivity (CH_3CN , $c = 5 \times 10^{-7}$ mol L⁻¹): $\Lambda = 1800$ Ω⁻¹ cm² mol⁻¹. SQUID: μ_{eff} (300 K) = 1.61 μ_B. Anal. calcd for C₄₈H₅₀F₆S₂Sb (926.80): C, 62.21; H, 5.44. Found C, 62.57; H, 5.62.

Synthesis of (2,6-Mes₂C₆H₃Se)₂SbF₆ ([17b][SbF₆])

Solid **16b** (220 mg, 0.28 mmol) was added to a stirred solution of [NO][SbF₆] in propionitrile (15 mL). After 12 h of stirring, the solvent was removed under reduced pressure affording a deep blue solid that was recrystallized from $\text{CH}_2\text{Cl}_2/n$ -hexane (1 : 1) at room temperature to yield [17b][SbF₆] (280 mg, 0.27 mmol, 98%).

^{77}Se -NMR (CDCl_3 , r.t.): $\delta =$ no signal. ^{77}Se -NMR (CDCl_3 , 223 K): $\delta = 1362.3$ ppm. ^{77}Se -NMR (CD_3CN): $\delta = 1362.4$ ppm. ESI MS (CH_3CN , positive mode): $m/z = 786.3$ [C₄₈H₅₀Se₂]⁺ for **17b**. UV-vis (CH_2Cl_2 , $c = 1 \times 10^{-3}$ mol L⁻¹): $\lambda_{\text{max}} = 710$ nm. Molar conductivity (CH_3CN , $c = 5 \times 10^{-7}$ mol L⁻¹): $\Lambda = 600$ Ω⁻¹ cm² mol⁻¹. SQUID: μ_{eff} (300 K) = 1.48 μ_B. Anal. calcd for C₄₈H₅₀F₆Se₂Sb (1105.59): C, 56.49; H, 4.94. Found C, 56.57; H, 5.12.

Synthesis of (2,6-Mes₂C₆H₃Te)₂SbF₆ ([17c][SbF₆])

Solid **16c** (247 mg, 0.28 mmol) was added to a stirred solution of [NO][SbF₆] in propionitrile (15 mL). After 8 h of stirring, the solvent was removed under reduced pressure affording a purple blue almost black solid that was recrystallized from propionitrile at room temperature to yield [17c][SbF₆] (300 mg, 0.27 mmol, 95%).

^{125}Te -NMR (CD_3CN): $\delta = 1698.7$ ppm. ^{125}Te -NMR (CDCl_3): $\delta = 1703.8$ ppm. ESI MS (CH_3CN , positive mode): $m/z = 882.2$ [C₄₈H₅₀Te₂]⁺ for **17c**. UV-vis (CH_2Cl_2 , $c = 1 \times 10^{-3}$ mol L⁻¹): $\lambda_{\text{max}} = 583$ nm. Molar conductivity (CH_3CN , $c = 5 \times 10^{-7}$ mol L⁻¹): $\Lambda = 540$ Ω⁻¹ cm² mol⁻¹. SQUID: μ_{eff} (300 K) = 1.41 μ_B. Anal. calcd for C₅₁H₅₅F₆NTe₂Sb (1172.94): C, 52.22; H, 4.73; N, 1.19. Found C, 52.38; H, 4.62; N, 1.24.

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