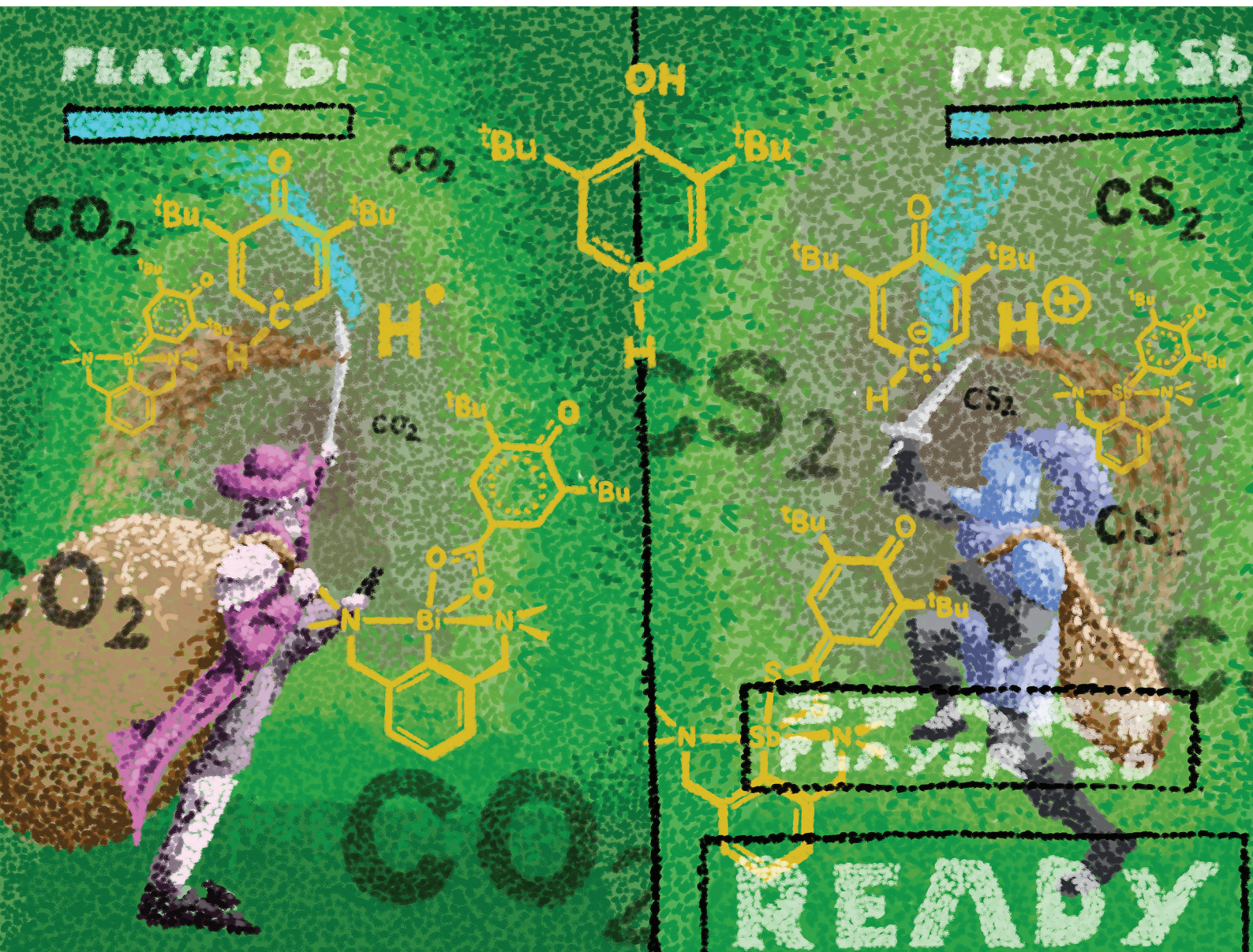


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C–H bond activation at antimony(III): synthesis and reactivity of Sb(III)–oxyaryl species†‡

Gabriel Duneş,^{a,b} Marie Cordier,^a Samia Kahlal,^a Alpar Pöllnitz,^b Jean-Yves Saillard,^{*a} Cristian Silvestru^{ID} ^{*b} and Yann Sarazin^{ID} ^{*a}

We report on the synthesis, structure and reactivity of $[(\text{NCN}^{\text{Me4}})\text{Sb}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-}3,5\text{-O-}4)]$ (**3**), an organoantimony(III)–oxyaryl species obtained upon $\text{C}_{\text{sp}^2}\text{-H}$ bond activation in a phenolate ligand and stabilised by the monoanionic pincer $(\text{NCN}^{\text{Me4}})^-$. The mechanism leading to the formation of **3** is highly sensitive to steric considerations. It was probed experimentally and by DFT calculations, and a number of intermediates and related complexes were identified. All data agree with successive heterolytic bond cleaving and bond forming processes involving charged species, rather than a pathway involving free radicals as previously exemplified with congeneric bismuth species. The nucleophilic behaviour of the oxyaryl ligand in **3**, a complex that features both zwitterionic and quinoidal attributes, was illustrated in derivatisation reactions. In particular, insertion of CS_2 in the $\text{Sb-C}_{\text{oxyaryl}}$ bond generates $[(\text{NCN}^{\text{Me4}})\text{Sb}(\text{S}_2\text{C-C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-}3,5\text{-O-}4)]$.

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Introduction

The activation of C–H bonds is essential in organic and organometallic synthesis, catalysis, and industrial processes. While this arena has been dominated by late transition metals, main group compounds have also entered the game in recent years.¹ Salient examples with metals from groups 2, 13 and 14 include C–H bond activation in aryl, allylic, and benzylic positions, in hydropyrimidine and in α -positions of cyclic thioethers.^{2–8} In group 15, bismuth compounds have also been investigated, as this metal is a component of the “ $n\text{MnO}_3/\text{Bi}_2\text{O}_3$ ” catalyst of the SOHIO process.⁹ BiCl_3 activated the *para*-C–H position in the phenolate $2,6\text{-}^t\text{Bu}_2\text{-C}_6\text{H}_3\text{-O}^-$, yielding C–C coupled quinoidal (**A**) and biphenolic (**B**) products *via* a free radical mechanism (eqn (1)).¹⁰

The organobismuth(III) $[(\text{NCN}^{\text{Me4}})\text{BiCl}_2]$, where NCN^{Me4} is the pincer $2,6\text{-}(\text{Me}_2\text{NCH}_2)_2\text{-C}_6\text{H}_3^-$, generated equimolar amounts of the Bi–oxyaryl $[(\text{NCN}^{\text{Me4}})\text{Bi}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-}3,5\text{-O-}4)]$ (Fig. 1, **I**) and of $2,6\text{-}^t\text{Bu}_2\text{-C}_6\text{H}_3\text{OH}$ upon reaction with two equivalents of $[(2,6\text{-}^t\text{Bu}_2\text{-C}_6\text{H}_3\text{O})\text{K}]$,^{11,12} based on Hanna's BiCl_3 precedent,¹⁰ the mechanism was suggested to involve radical species. A case of Bi-mediated intramolecular C–H activation in benzylic position was recently reported in the planar Bi(III) compound $[(\text{NCN}^{\text{DiPP2}})\text{Bi}]$ (**II**) supported by a trianionic pincer ligand, with concomitant extrusion of H_2 .¹³ Astounding work by Cornella has demonstrated the exquisite ability of Bi(III) to engage in controlled redox cycles with Bi(I) and Bi(V) and to catalyse a suite of organic reactions.^{14–18}

The chemistry of the lighter metalloid, antimony(III), is comparatively less developed.¹⁹ Recent breakthroughs in organoantimony(III) chemistry stem from the implementation of supporting ligands that stabilise highly reactive mononuclear species.^{20–24} Pincer ligands have been key to the developments of organoantimony compounds,^{19,20,25} and NCN^- pincers have a leading role among these. Although SbF_5 or SbF_6^- superacids are known to promote C–H bond activation and functionalisation processes,^{26–32} examples of metal-mediated reactions involving discrete organoantimony complexes for C–H bond activation are seldom. The azastibocine triflate $[(\text{C}_6\text{H}_{11}\text{N}(\text{CH}_2\text{C}_6\text{H}_4)_2)\text{SbOTf}]$ (Fig. 1, **III**) is an efficient Lewis acid catalyst for Mannich reactions; $[(\text{C}_6\text{H}_{11}\text{N}(\text{CH}_2\text{C}_6\text{H}_4)_2)\text{SbF}]$ triggers cyclisation-aromatisation from arylacetylenes, anilines and aldehydes.^{33,34} Antimony(V)-oxo porphyrins mediate C–H to C–C bond conversion in Michael acceptors under photoirradiation, without variation of the high-valent oxidation state.³⁵

^aUniversité de Rennes, CNRS, Institut des Sciences Chimiques de Rennes, UMR 6226, Campus de Beaulieu, 35042 Rennes, Cedex, France.E-mail: yann.sarazin@univ-rennes.fr, jean-yves.saillard@univ-rennes.fr^bDepartment of Chemistry, Supramolecular Organic and Organometallic Chemistry Centre (SOOMCC), Faculty of Chemistry and Chemical Engineering, Babeş-Bolyai University, 11 Arany Janos, 400028 Cluj-Napoca, Romania.E-mail: cristian.silvestru@ubbcluj.ro

† Dedicated to Prof. Hans J. Breunig, in recognition of his immense contribution to the chemistry of organopnictogens.

‡ Electronic supplementary information (ESI) available: Experimental details, spectroscopic characterisations, crystallographic and computational data. CCDC 2335992–2335998. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d4dt01400f>

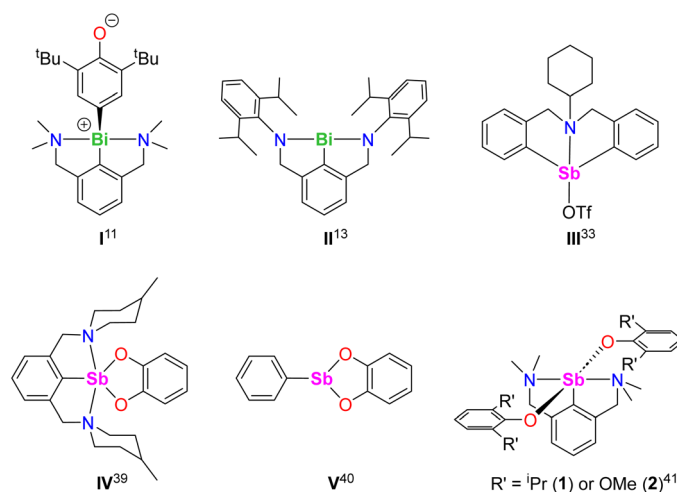
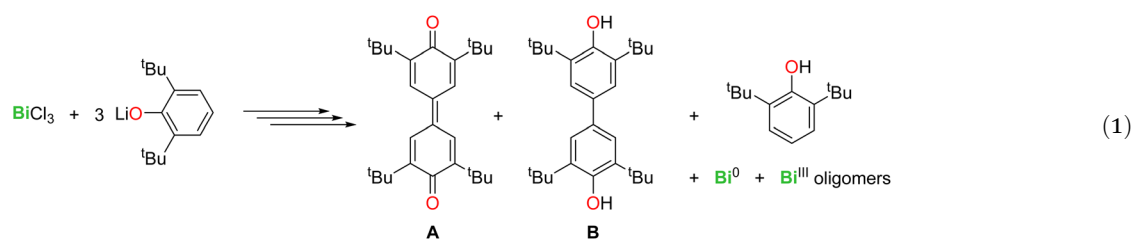


Fig. 1 Selected examples of known Sb(III) and Bi(III) relevant to the present work.

As part of our program targeting p-block complexes with alkoxides and other O-based derivatives,^{36–38} we have taken an interest in heteroleptic Sb(III)-phenolato complexes. Although their Sb(V) congeners are plethora, few examples of heteroleptic arylantimony(III) phenolates are known. They include the catecholates $[\{2,6\text{-}(\text{MeN}[\text{CH}_2\text{CH}_2)_2\text{NCH}_2)_2\text{C}_6\text{H}_3\}\text{Sb}(\text{O}_2\text{-}1,2\text{-C}_6\text{H}_4)]$ (IV) and $[\text{PhSb}(\text{O}_2\text{-}1,2\text{-C}_6\text{H}_4)]$ (V).^{39,40} Besides, we have recently disclosed the bis(phenolate)s $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{OC}_6\text{H}_3\text{-}^i\text{Pr}_2\text{-}2,6)_2]$ (1) and $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{OC}_6\text{H}_3\text{-(OMe)}_2\text{-}2,6)_2]$ (2).⁴¹ However, we are showing herein that under identical conditions, their derivative with *t*Bu substituents, $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{OC}_6\text{H}_3\text{-}^t\text{Bu}_2\text{-}2,6)_2]$, cannot be obtained, and instead the oxyaryl $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-}3,5\text{-O-}4)]$ (3) is isolated quantitatively. We are reporting here on the syntheses and structures of 3 and related Sb(III) complexes. Mechanisms, bonding features and reactivity patterns, including a case of insertion into a Sb(III)–C bond, are discussed. Investigations probing whether a radical-based mechanism could account for the formation of 3 provide a different perspective.

Results and discussion

General syntheses and characterisation

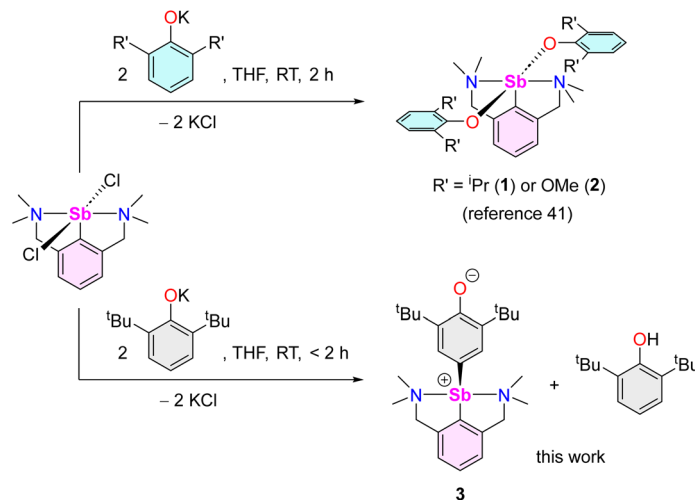
The reaction of $[\{\text{NCN}^{\text{Me}_4}\}\text{SbCl}_2]$ ⁴² with two equivalents of $[\{2,6\text{-}^t\text{Bu}_2\text{-C}_6\text{H}_3\text{O}\}\text{K}]$ affords clean formation of the Sb(III)-

oxyaryl complex $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-}3,5\text{-O-}4)]$ (3) upon release of $2,6\text{-}^t\text{Bu}_2\text{-C}_6\text{H}_3\text{OH}$ and KCl (Scheme 1). This outcome contrasts with the analogous reactions involving the less bulky phenolates $[\{2,6\text{-}^i\text{Pr}_2\text{-C}_6\text{H}_3\text{O}\}\text{K}]$ and $[\{2,6\text{-(OMe)}_2\text{-C}_6\text{H}_3\text{O}\}\text{K}]$, which were shown to afford the anticipated $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{OC}_6\text{H}_3\text{-}^i\text{Pr}_2\text{-}2,6)_2]$ (1) and $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{OC}_6\text{H}_3\text{-(OMe)}_2\text{-}2,6)_2]$ (2).⁴¹ The overall behaviour was reminiscent of that observed with bismuth, where a radical mechanism was suggested for the formation of the congeneric $[\{\text{NCN}^{\text{Me}_4}\}\text{Bi}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-}3,5\text{-O-}4)]$.^{11,12,43}

The air-sensitive 3 forms in yields over 85% within 2 h in THF at room temperature. The complex can be depicted as a zwitterion with a positively charged Sb(III) centre, or its quinoid-like, neutral-at-metal mesomer (Fig. 2). The presence of the oxidation products resulting from free radical couplings, *i.e.* 3,3',5,5'-tetra-*tert*-butyldiphenylquinone (A) and 3,3',5,5'-tetra-*tert*-butyl- $[1,1'\text{-biphenyl}]\text{-}4,4'\text{-diol}$ (B) as reported in the reaction of BiCl_3 with $[\{2,6\text{-}^t\text{Bu}_2\text{-C}_6\text{H}_3\text{O}\}\text{Li}]$ (see eqn (1)),¹⁰ was not detected in the synthesis of 3.

Measurements showed that as expected for zwitterions,⁴⁴ the conductivity of a THF solution of 3 ($\Lambda_m = 27.2 \text{ mS cm}^2 \text{ mol}^{-1}$) is relatively low. It is comparable to that for the neutral bis(phenolate) 1 ($21.2 \text{ mS cm}^2 \text{ mol}^{-1}$), and only marginally higher than that for $[\text{Sb}(\text{mesityl})_3]$ and $[\{\text{NCN}^{\text{Me}_4}\}\text{SbCl}_2]$ (resp. 0.0 and $9.1 \text{ mS cm}^2 \text{ mol}^{-1}$). By comparison, the





Scheme 1 Synthesis of $[(\text{NCN}^{\text{Me}_4})\text{Sb}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-3,5-O-4})]$ (**3**).

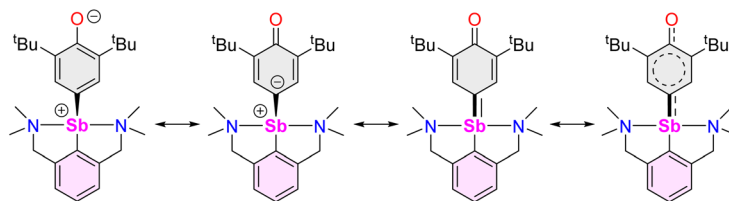


Fig. 2 Mesomeric forms of $[(\text{NCN}^{\text{Me}_4})\text{Sb}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-3,5-O-4})]$ (**3**).

dissociated ion pair $[\text{Na}(\text{OEt}_2)_4]^+ \cdot [\text{H}_2\text{N}\{\text{B}(\text{C}_6\text{F}_5)_3\}_2]^-$ (ref. 45) gives rise to much greater conductivity under otherwise identical conditions ($22\,000\text{ mS cm}^2\text{ mol}^{-1}$). Complex **3** is diamagnetic. The NMR data of **3** recorded in CD_3CN and CD_2Cl_2 were very different from those in $\text{THF-}d_8$ although they corroborated in all cases the identity and the purity of the complex (see ESI, Fig. S1–S6†). The VT ^1H NMR spectra in $\text{THF-}d_8$ were consistent with free rotation of the oxyaryl fragment around the $\text{Sb-C}_{\text{oxyaryl}}$ bond at high temperature (338 K),⁴⁶ and with a frozen edge-inversion processes⁴⁷ at 263 K (see ESI† for VT NMR). The FTIR spectrum of **3** (Fig. S7†) features a strong band at 1559 cm^{-1} for the stretching of the C–O bond. It is below the values for carbonyl stretching in quinones, *ca.* $1660\text{--}1690\text{ cm}^{-1}$, and lower even than the remarkably low out-of-phase carbonyl stretching in substituted diphenoquinones, *e.g.* 1602 cm^{-1} in **A**.⁴⁸ Such low $\nu_{\text{C=O}}$ stretching frequency is consistent with a partial C=O double bond order, and agrees with the dual zwitterionic and quinoid-like nature of **3**.

The molecular solid-state structure of **3** features a four-coordinate metalloid in a pseudotrigonal bipyramidal arrangement, with the $(pR_{\text{N1}}, pR_{\text{N2}})$ enantiomer⁴⁹ depicted in Fig. 3. The short Sb1–N1 and Sb1–N2 interatomic distances (2.4458(11) and 2.4436(11) Å) testify to two strong N→Sb dative

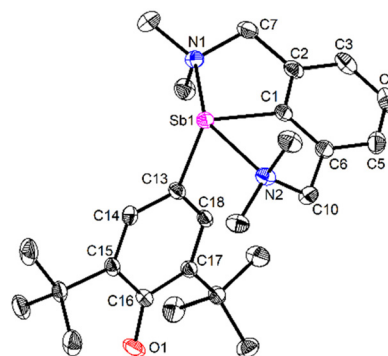


Fig. 3 View of the molecular structure of $[(\text{NCN}^{\text{Me}_4})\text{Sb}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-3,5-O-4})]$ (**3**). H atoms omitted for clarity. Ellipsoids at the 50% probability level. Only one of the two enantiomers ($pR_{\text{N1}}, pR_{\text{N2}}$)⁴⁹ in the asymmetric unit is shown. Selected distances (Å) and angles ($^\circ$): Sb1–C1 = 2.1134(13), Sb1–C13 = 2.0862(12), Sb1–N1 = 2.4458(11), Sb1–N2 = 2.4436(11), C13–C14 = 1.4063(17), C13–C18 = 1.4047(17), C14–C15 = 1.3803(17), C15–C16 = 1.4572(18), C16–C17 = 1.4559(17), C17–C18 = 1.3777(17), C16–O1 = 1.2708(16); C1–Sb1–C13 = 101.02(5), C1–Sb1–N1 = 73.94(4), C1–Sb1–N2 = 74.16(4), C13–Sb1–N1 = 96.54(4), C13–Sb1–N2 = 89.09(4), N1–Sb1–N2 = 148.10(4).

bonds,⁴² and the coordination geometry about Sb is best described as see-saw ($\tau_4 = 0.66$).⁵⁰ The Sb–C_{oxyaryl} bond length (Sb1–C13 = 2.0862(12) Å) is below the low end of Sb–C_{aryl} dis-



tances in four-coordinate arylantimony(III) complexes.^{51–53} The Sb–C_{oxyaryl} interatomic distance in **3** is comparable with the value found for the few reported Sb=C double bonds.^{54–56} The Sb–C13 bond in **3** is hence intermediary between single and double Sb(III)–carbon bonds. The C16–O1 distance in **3** (1.2708 (16) Å) also ranges between the values of single and double carbon–oxygen bonds, e.g. as in **B** (1.385(4) and 1.391(3) Å)⁵⁷ vs. 2,6-di-*tert*-butyl-1,4-benzoquinone (1.2456(6) and 1.2570(6) Å)⁵⁸ or 2,4,6-tri-*tert*-butylphenoxy radical (1.246(2) Å).⁵⁹ It hence feels legitimate to consider that this fragment exhibits partial quinoidal behaviour. A more detailed description of the structure of **3** is provided in ESI.†

Synthetic and NMR insight

Beyond the fact that it requires *tert*-butyl groups in positions 2 and 6 of the phenolate instead of smaller substituents (see Scheme 1), we investigated the mechanism leading to **3**. Monitoring of the reaction in THF-*d*₈ by ¹H NMR spectroscopy indicated clean and quantitative production of **3** and 2,6-*t*-Bu₂-C₆H₃OH, without detectable presence of **A** or **B** (Fig. S37 and S42†). Complex **3** is EPR silent (as a crystalline solid or in solution in THF); only minor signals corresponding to an organic fragment, most likely resulting from minute decomposition during sample preparation, were detected. The sequential NMR-EPR-NMR spectroscopic analysis of the same sample of crystalline **3** dissolved in THF-*d*₈ confirmed the sample remained intact in solution, within the accuracy allowed by NMR, over the course of 2 h (Fig. S11†). Addition of TEMPO or galvinoxyl to a mixture of [{NCN^{Me4}}SbCl₂] and [{2,6-*t*-Bu₂-C₆H₃O}K] in THF did not prevent formation of **3**, although TEMPO ultimately led to an oxo-bridged dimer (*vide infra*). Moreover, NMR monitoring showed that the rates of formation of **3** in the dark and in ambient light are near-identical, which further militates against the involvement of radicals. Based on these observations (also corroborated by DFT calculations, *vide infra*), we propose that unlike earlier suggestions for bismuth,¹¹ **3** does not result from homolytic bond cleavages and recombination of free radicals.

The identity of the precursors required to achieve the C–H activation leading to **3** was probed (Scheme 2). The complex can be obtained from [{NCN^{Me4}}SbCl₂], although the reaction is slow in this case (12 h in THF-*d*₈, Scheme 2a). The two –NMe₂ side-arms in the pincer ligand can be replaced by bulkier and stronger –N^{*t*}Pr₂ donors; the Sb(III)-oxyaryl resulting from C–H activation of the phenolate, **4** (Scheme 2b), is near-quantitatively isolated. However, with the dichloro precursor [{CN^{Me2}}SbCl₂],⁵³ where the CN^{Me2} bidentate ligand bears a single –NMe₂ tether, the reaction with two equivalents of [{2,6-*t*-Bu₂-C₆H₃O}K] gives the bis(phenolate) [{CN^{Me2}}Sb(OC₆H₃-*t*-Bu₂-2,6)] (**5**) akin to **1** and **2** (Scheme 2c). No C–H activation of the phenolate occurred, indicating that the presence of *two* Lewis bases is mandatory for the process to occur. Accordingly, addition of *N,N*-dimethyl-aminopyridine (DMAP) to a solution of pristine **5** affords the tetramer [{CN^{Me2}}Sb(C₆H₂-*t*-Bu₂-3,5-O-4)]₄ (**6**₄) as insoluble crystals.⁶⁰ ¹H NMR monitoring of this

reaction in THF-*d*₈ showed disappearance of all Sb-based species, with regeneration of DMAP and stoichiometric release of 2,6-*t*-Bu₂-C₆H₃OH (Fig. S19†). We assume the addition of DMAP to **5** (or, more generally, the presence of two stabilising Lewis bases) is required in order to stabilise the transient, positively charge antimony centre *en route* towards the formation of **6**₄ (or, in the seminal case with the {NCN^{Me4}} ligand, the production of **3**), as pointed out by our DFT calculations (*vide infra*).

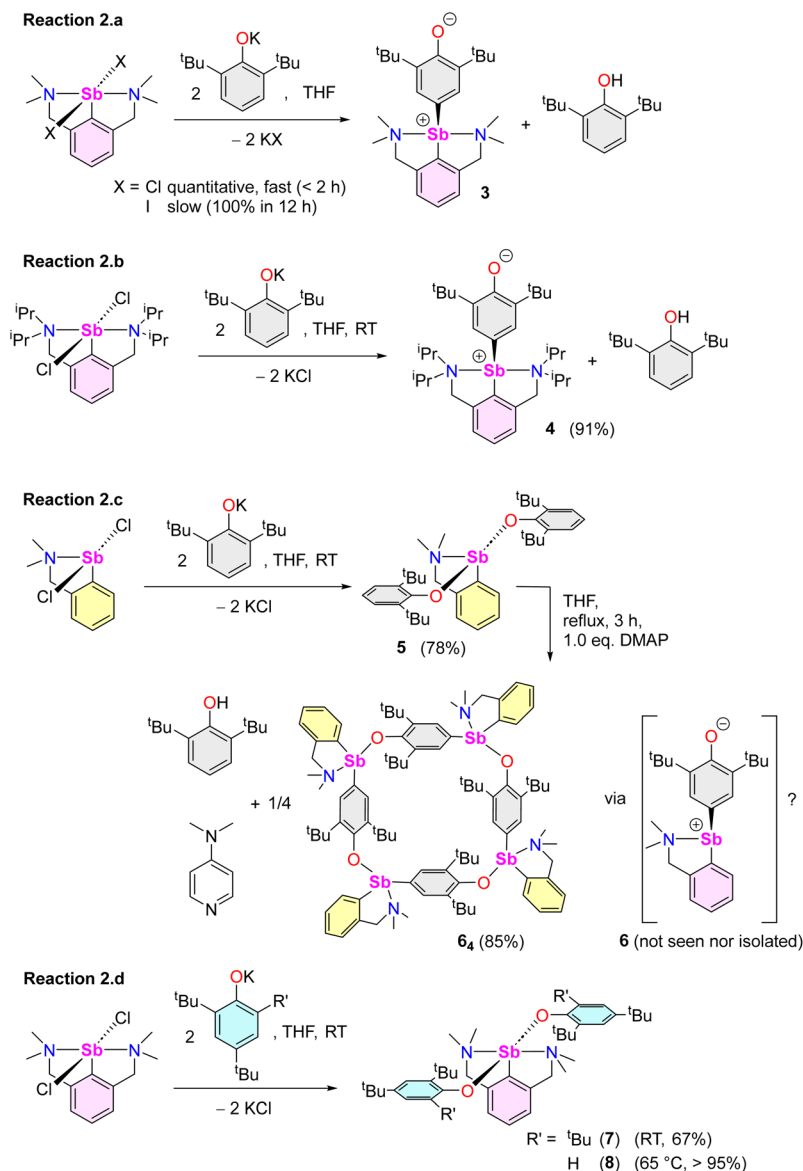
Compound **6**₄ can be seen as the tetrameric equivalent of zwitterionic **3** (Fig. 4). Each Sb centre is tetracoordinated. The O- and N-atoms are in *trans* position (N1–Sb1–O1 = 157.25 (12)°), whereas the two carbon atoms are in *cis* (C4'–Sb1–C15 = 98.36(16)°). The Sb–C_{oxyaryl} distance is noticeably larger than that in **3** (2.121(5) vs. 2.0862(12) Å). The Sb1–O1 bond in **6**₄ (2.088(4) Å) is much shorter than in the four-coordinate Sb-bis(phenolate) **1** (2.187(1) Å).⁴¹ However, the amino side-arm in **6**₄ is binding with antimony considerably more weakly (Sb1–N1 = 2.574(4) Å) than in both **3** and **1** (Sb–N = 2.4436(11)–2.4458(11) Å and 2.426(1) Å,⁴¹ respectively). Complex **6**₄ is the utmost expression of the zwitterionic form of **6** (a complex that we could not isolate nor detect spectroscopically), yielding a charge-neutral tetramer through binding of the negatively-charged O atoms to Sb cations.

The reaction of [{NCN^{Me4}}SbCl₂] with two equivalents of [{2,4,6-*t*-Bu₃-C₆H₂O}K] affords [{NCN^{Me4}}Sb(OC₆H₂-*t*-Bu₃-2,4,6)]₂ (**7**, Scheme 2d), demonstrating that the presence of *t*-Bu groups in positions 2 and 6 of the phenolate is compatible with the formation of regular organoantimony(III) bis(phenolate)s. Finally, the bis(phenolate) [{NCN^{Me4}}Sb(OC₆H₂-*t*-Bu₂-2,4)] (**8**) was obtained with [{2,4-*t*-Bu₂-C₆H₃O}K], even at 65 °C (Scheme 2d); hence, C–H activation only occurs in position 4 of di-*tert*-butylphenolate. This last finding also confirms that steric hindrance in the direct vicinity of the O_{phenolate} atom is not an impeding factor for the C–H activation step, as inferred from the opposite reactivity observed during the formations of **1** and **2** vs. **3** from differently substituted 2,6-dialkylphenolates (Scheme 1).

Overall, it transpires from these experimental data that the behaviour of Sb(III) cannot be extrapolated from that of its heavier congener, Bi(III). Besides, in our hands, the reaction of SbCl₃ with three equivalents of [{2,6-*t*-Bu₂-C₆H₃O}K] in THF gave [{2,6-*t*-Bu₂-C₆H₃O}Sb] quantitatively; no traces of **A** or **B** were detected.^{10,61}

Several model reactions were performed in THF-*d*₈ and monitored by NMR to probe the behaviour of the system (Scheme 3). First, although potassium phenolate was fully consumed, the reaction of [{NCN^{Me4}}SbCl₂] with an *equimolar* amount of [{2,6-*t*-Bu₂-C₆H₃O}K] did *not* produce [{NCN^{Me4}}Sb(Cl)(OC₆H₃-*t*-Bu₂-2,6)] (**9**). Instead, it yielded half equivalents of **3**, 2,6-*t*-Bu₂-C₆H₃OH and unreacted [{NCN^{Me4}}SbCl₂] within the first point of analysis (30 min; Scheme 3e and Fig. S38†). This suggests that **9** is either never formed, or that it is an intermediate in the formation of **3**. By contrast, in a two-step process (Scheme 3f), the reaction of [{NCN^{Me4}}SbCl₂] with equimolar [{2,4,6-*t*-Bu₃-C₆H₂O}K] first generated a mixture of





Scheme 2 Reactions of Sb(III) dihalides with potassium phenolates.

unreacted dichloride, bisphenolate 7 and a small amount of, at least, one other phenolate thought to be $[\{NCN^{Me4}\}Sb(Cl)](OC_6H_2-^tBu_3-2,4,6)]$ (**10**). This intractable mixture, assumed to be an equilibrium between **10** vs. $[\{NCN^{Me4}\}SbCl_2]$ and 7, was treated with one equivalent of $[\{2,6-^tBu_2-C_6H_3O\}K]$, generating **3** and 2,4,6-^tBu₃-C₆H₂OH (Fig. S43†). We hence reasoned that **3** may form through reaction of the monochloro, monophenolato intermediate **10**, with one equivalent of $[\{2,6-^tBu_2-C_6H_3O\}K]$. Consumption of **10** would drive the reaction towards complete formation of **3** and release of 2,4,6-^tBu₃-C₆H₂OH, with concomitant disappearance of $[\{NCN^{Me4}\}SbCl_2]$ and 7. In agreement with this hypothesis, in a separate experiment, the combination of $[\{NCN^{Me4}\}SbCl_2]$ with equimolar 7 in THF-*d*₈ at RT led to an equilibrated mixture with an NMR signature (Fig. S44†) matching that of the first step of reaction (3.f); the same reaction at −35 °C gave a single, new species, assumed

to correspond to pure **10**. However, it is also possible that **3** results from direct action of $[\{2,6-^tBu_2-C_6H_3O\}K]$ on 7; indeed, a native sample of 7 reacts with one equivalent of $[\{2,6-^tBu_2-C_6H_3O\}K]$ to give **3**, 2,4,6-^tBu₃-C₆H₂OH and $[\{2,4,6-^tBu_3-C_6H_2O\}K]$ (Fig. S46†). Both mechanisms may well be simultaneously at play; yet, regardless of the prevailing manifold, production of **3** and release of 2,4,6-^tBu₃-C₆H₂OH according to reaction (3.f) are clean and quantitative.

The synthesis of a Bi(III)-oxyaryl species by reaction of $[\{NCN^{Me4}\}BiCl_2]$ with a 1 : 1 combination of $[\{2,6-^tBu_2-C_6H_3O\}K]$ and $[\{2,6-^iPr_2-C_6H_3O\}K]$ was reported to have failed.¹¹ In this light, and in view of the outcome of reaction (3.f), it was surprising to observe that in a two-step process, $[\{NCN^{Me4}\}SbCl_2]$ reacts with $[\{2,6-^iPr_2-C_6H_3O\}K]$ and then $[\{2,6-^tBu_2-C_6H_3O\}K]$ to afford half-equivalents of **3**, $[\{NCN^{Me4}\}Sb](OC_6H_2-^iPr_2-2,6)_2]$ (**1**) and 2,6-^tBu₂-C₆H₃OH (Scheme 3g and



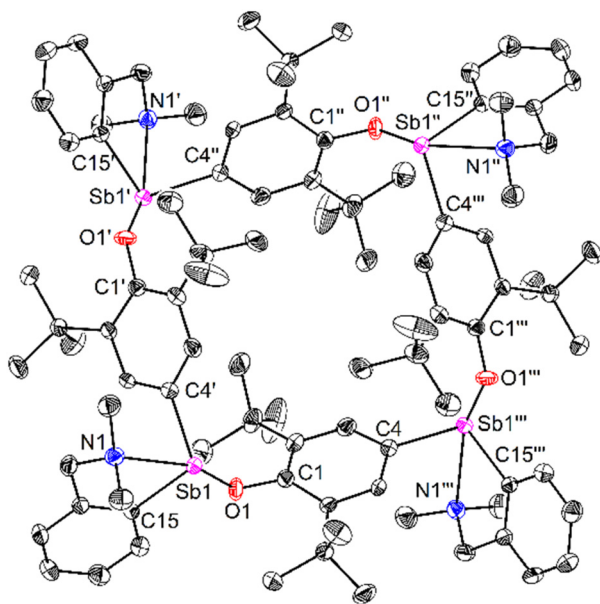


Fig. 4 Representation of the molecular structure of $[(\text{CN}^{\text{Me}_2})\text{Sb}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-}3,5\text{-O-}4)]_4$ (**64**). H atoms omitted for clarity. Ellipsoids at the 50% probability level. Selected distances (Å) and angles (°): Sb1–C4' = 2.121(5), Sb1–C15 = 2.157(4), Sb1–N1 = 2.574(4), Sb1–O1 = 2.088(4), C1–O1 = 1.346(5), C4'–Sb1–O1 = 99.36(16), C4'–Sb1–C15 = 98.36(16), C4'–Sb1–N1 = 82.47(15), C15–Sb1–N1 = 72.87(15), C15–Sb1–O1 = 84.46(15), N1–Sb1–O1 = 157.25(12).

Fig. S47†). Besides, the reaction of $[\{\text{NCN}^{\text{Me}_4}\}\text{SbCl}_2]$ with equimolar $[\{2,6\text{-}^i\text{Pr}_2\text{-C}_6\text{H}_3\text{O}\}\text{K}]$ was consistent with the formation of **1** along unreacted dichloride. The reason behind the different behaviour of $[\{2,4,6\text{-}^t\text{Bu}_3\text{-C}_6\text{H}_2\text{O}\}\text{K}]$ and $[\{2,6\text{-}^i\text{Pr}_2\text{-C}_6\text{H}_3\text{O}\}\text{K}]$ towards $[\{\text{NCN}^{\text{Me}_4}\}\text{SbCl}_2]$ originates from the fact that unlike **7** (*vide supra*), complex **1** is inert toward $[\{\text{NCN}^{\text{Me}_4}\}\text{SbCl}_2]$; the 1 : 1 mixture of the two compounds in THF-*d*₈ remains essentially unchanged after 24 h at 65 °C. Unexpectedly, treatment of **3** with one equivalent of 2,6-ⁱPr₂-C₆H₃OH generated the exact same mixture of **1**, **3** and 2,6-^tBu₂-C₆H₃OH (half equivalents of each; Scheme 3h and Fig. S49†) as that obtained in reaction (3.g), in a process postulated to evolve through the intermediate $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-}3,5\text{-OH-}4)(\text{OC}_6\text{H}_2\text{-}^i\text{Pr}_2\text{-}2,6)]$ (**11**). Note also that **3** and 2,6-^tBu₂-C₆H₃OH engage in an equilibrium in solution; the ¹H and ²H monitoring of a 1 : 1 mixture of **3** and 2,6-^tBu₂-C₆H₃OD returned a solution that contained **3** along 2,6-^tBu₂-4-^D-C₆H₂OH, 2,6-^tBu₂-4-^D-C₆H₂OD and 2,6-^tBu₂-C₆H₃OH (Fig. S50 and S51†).

We then attempted to prepare a complex amenable to nucleophilic substitution. Crystals of $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-}3,5\text{-OH-}4)]\text{Cl}$ (**12**) were obtained in 75% yield by reaction of $[\text{Me}_3\text{NH}]^+\text{Cl}^-$ with **3** (Scheme 4), a reaction that illustrates the zwitterionic character of **3**. The Sb(III)-hydroxyaryl **12**, formally the product of C–H activation on 2,6-di-*tert*-butylphenol, is, to our knowledge, unique; there is no related antimony complex in the CCDC database.⁶² Similarly, treatment of **3** with Bochmann's acid⁴⁵ afforded the loose ion pair $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-}3,5\text{-OH-}4)]^+[\text{H}_2\text{N}\{\text{B}(\text{C}_6\text{F}_5)_3\}_2]^-$ (**12'**).

The geometry about the Sb atom in **12** (Fig. 5) is *pseudo*-trigonal bipyramidal (see ESI† for details). Complex **12** forms a loose ion pair with a long Sb1–Cl1' distance (3.6761(14) Å). The Cl[–] anion is engaged in H-bonding with the OH group (OH...Cl = 2.1538(341) Å).⁶³ Complexes **12** and **3** present noticeable structural differences. In particular, the C–O_{hydroxyl} distance in **12** (C16–O1 = 1.373(4) Å) is longer than the corresponding C–O_{oxy} one in **3** (1.2708(16) Å). The discrepancies in the metric parameters from **3** to **12** indicate protonation of the O atom, re-aromatisation of the C13–C14–C15–C16–C17–C18 ring, and loss of quinoidal character. The molecular solid-state structure of **12'** was also elucidated (Fig. S62†). Characteristically, there is no interaction between the cation and $[\text{H}_2\text{N}\{\text{B}(\text{C}_6\text{F}_5)_3\}_2]^-$.^{64,65}

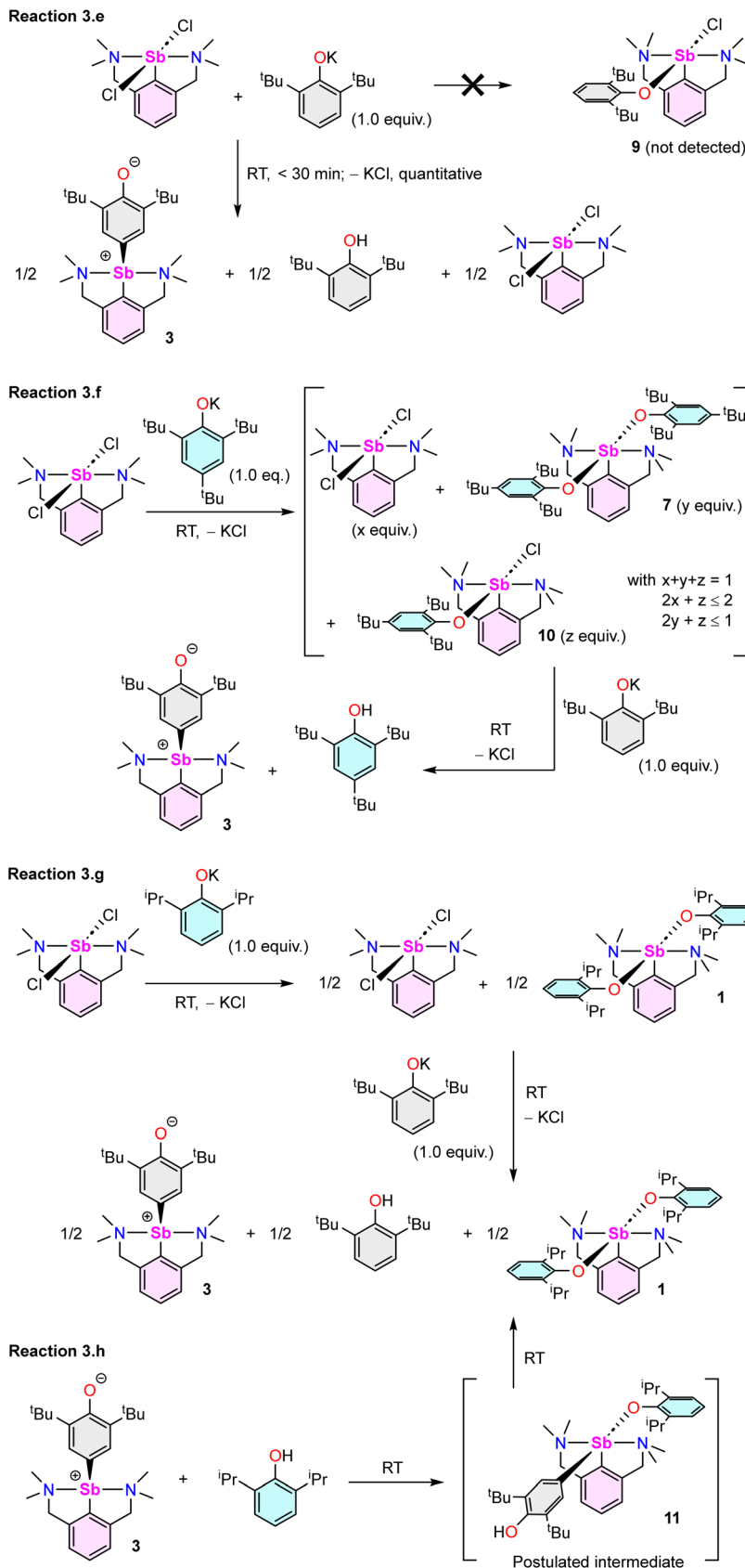
Complex **12** reacts with equimolar $[\{2,6\text{-}^t\text{Bu}_2\text{-C}_6\text{H}_3\text{O}\}\text{K}]$ in THF-*d*₈ to afford a 1 : 1 mixture of **3** and 2,6-^tBu₂-C₆H₃OH (Fig. S52†). Two pathways can be envisaged. The reaction may occur *via* salt metathesis that implicates an intermediate $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-}3,5\text{-OH-}4)(\text{OC}_6\text{H}_3\text{-}^t\text{Bu}_2\text{-}2,6)]$ akin to putative **11**. Alternatively, since this intermediate could not be isolated nor seen spectroscopically, deprotonation of **12** by $[\{2,6\text{-}^t\text{Bu}_2\text{-C}_6\text{H}_3\text{O}\}\text{K}]$ with subsequent extrusion of KCl could also take place. To distinguish between these two pathways, the reaction of **12** with one equivalent of $[\{2,6\text{-}^i\text{Pr}_2\text{-C}_6\text{H}_3\text{O}\}\text{K}]$ was carried out. It afforded *half* equivalents of **3**, **1** and 2,6-^tBu₂-C₆H₃OH (Scheme 5; Fig. S53†).

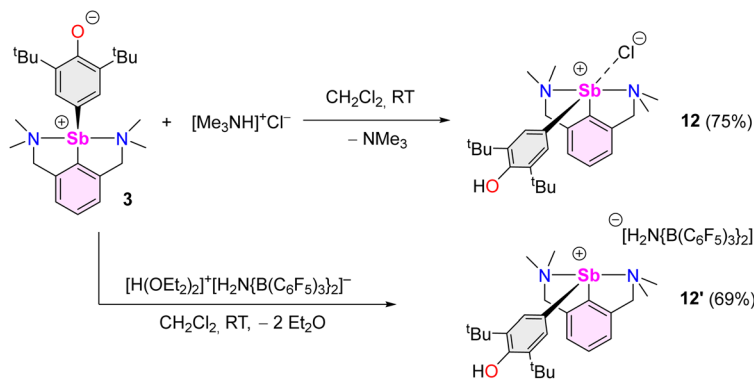
Again, initial steps consisting of salt metathesis progressing through **11** or deprotonation of **12** by $[\{2,6\text{-}^i\text{Pr}_2\text{-C}_6\text{H}_3\text{O}\}\text{K}]$ could both account for this outcome. If generated upon deprotonation of **12**, complex **3** and 2,6-ⁱPr₂-C₆H₃OH⁶⁶ would indeed further react to give half equivalents of **1**, **3** and 2,6-^tBu₂-C₆H₃OH, as learnt from reaction (3.h). Low temperature NMR in THF-*d*₈ provided conclusive evidence. The equimolar reaction of **12** with $[\{2,6\text{-}^i\text{Pr}_2\text{-C}_6\text{H}_3\text{O}\}\text{K}]$ at –40 °C led to a *ca.* 1 : 1 mixture of **3** and 2,6-ⁱPr₂-C₆H₃OH (Fig. S54†). Upon warming to 20 °C, this mixture evolved toward the final production of **3**, **1** and 2,6-^tBu₂-C₆H₃OH in equal concentrations. In addition, the DOSY spectrum recorded at –40 °C in THF-*d*₈ indicates the presence of two major species, assigned as **3** and 2,6-ⁱPr₂-C₆H₃OH; it was independently confirmed by cross-referencing with the ¹H NMR spectrum of a 1 : 1 mixture of these two compounds (THF-*d*₈, –40 °C). At any rate, the fact that *two* species are detected in substantial concentrations rules out a mechanism involving the formation of a *single* intermediate **11**. Hence, the reaction of **12** with $[\{2,6\text{-}^i\text{Pr}_2\text{-C}_6\text{H}_3\text{O}\}\text{K}]$ proceeds *via* initial deprotonation, followed by reaction of the resulting 2,6-ⁱPr₂-C₆H₃OH with **3** (Scheme 5).⁶⁷

Mechanistic proposal

DFT calculations at the PBE0/Def2TZVP-D3(BJ) level were performed to provide insights into the stability of the complexes and intermediates discussed therein and to shed some light on the mechanism leading to **3**. Calculations on the isolated bisphenolates $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{OC}_6\text{H}_3\text{-}^i\text{Pr}_2\text{-}2,6)_2]$ (**1**), $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{OC}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-}2,4,6)_2]$ (**7**) and $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{OC}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-}2,4)_2]$ (**8**), as well as the non-observed $[\{\text{NCN}^{\text{Me}_4}\}\text{Sb}(\text{OC}_6\text{H}_3\text{-}^t\text{Bu}_2\text{-}2,6)_2]$



Scheme 3 Reactions monitored by ^1H NMR spectroscopy in $\text{THF}-d_8$.



Scheme 4 Synthesis of $[(\text{NCN}^{\text{Me}_4})\text{Sb}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-3,5-OH-4})][\text{X}]$ ($\text{X} = \text{Cl}$, **12**; $\text{H}_2\text{N}(\text{B}(\text{C}_6\text{F}_5)_3)_2$, **12'**).

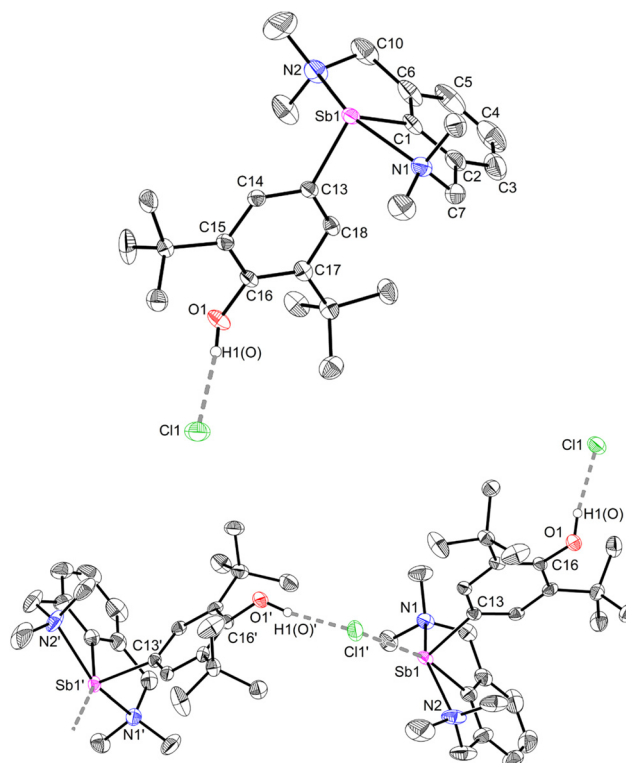
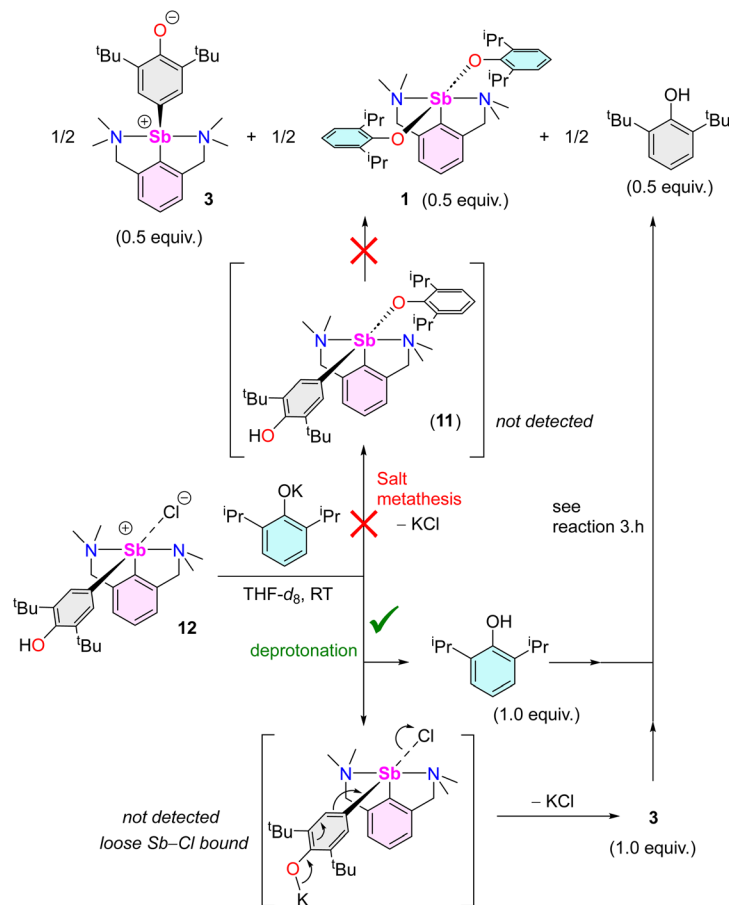


Fig. 5 Top: representation of the molecular structure of $[(\text{NCN}^{\text{Me}_4})\text{Sb}(\text{C}_6\text{H}_2\text{-}^t\text{Bu}_2\text{-3,5-OH-4})][\text{Cl}]$ (**12**). Bottom: arrangement of two adjacent units in the lattice. H atoms (except that on O atoms) and non-interacting CH_2Cl_2 molecules omitted for clarity. Ellipsoids at the 50% probability level. Selected distances (Å) and angles ($^\circ$): Sb1–C1 = 2.106(4), Sb1–C13 = 2.141(4), Sb1–N1 = 2.398(4), Sb1–N2 = 2.453(4), Sb1–Cl1 = 3.6761(14), C13–C14 = 1.387(5), C13–C18 = 1.389(5), C14–C15 = 1.404(5), C15–C16 = 1.410(5), C16–C17 = 1.422(5), C17–C18 = 1.387(5), O1–C16 = 1.373(4), O1–H1(O) = 0.909(6), Cl1–H1(O) = 2.154 (34); C1–Sb1–C13 = 95.52(15), C1–Sb1–N1 = 74.59(15), C1–Sb1–N2 = 74.41(18), C13–Sb1–N1 = 90.80(13), C13–Sb1–N2 = 90.55(14), N1–Sb1–N2 = 148.95(14).

analogue and the non-sterically hindered $[(\text{NCN}^{\text{Me}_4})\text{Sb}(\text{OC}_6\text{H}_3\text{-Me}_2\text{-2,6})_2]$ species, indicate rather similar structural and electronic properties for the five complexes (Table S3†). The Sb–O distance in $[(\text{NCN}^{\text{Me}_4})\text{Sb}(\text{OC}_6\text{H}_3\text{-}^t\text{Bu}_2\text{-2,6})_2]$ is only 0.060 and 0.056 Å longer than in $[(\text{NCN}^{\text{Me}_4})\text{Sb}(\text{OC}_6\text{H}_3\text{-Me}_2\text{-2,6})_2]$ and **1**, respectively. However, the corresponding Wiberg bond indices suggest somehow weaker covalency in the case of the first, more encumbered complex. Complex **7** exhibits similar bonding features as $[(\text{NCN}^{\text{Me}_4})\text{Sb}(\text{OC}_6\text{H}_3\text{-}^t\text{Bu}_2\text{-2,6})_2]$, whereas

the less sterically hindered **8** presents stronger Sb-phenolate bonding. The atomic charges are indicative of a non-negligible ionic bonding component, especially for $[(\text{NCN}^{\text{Me}_4})\text{Sb}(\text{OC}_6\text{H}_3\text{-}^t\text{Bu}_2\text{-2,6})_2]$ and **7**. The related single -NMe_2 tethered $[(\text{CN}^{\text{Me}_2})\text{Sb}(\text{OC}_6\text{H}_3\text{-}^t\text{Bu}_2\text{-2,6})_2]$ (**5**) exhibits similar bonding features, but with a somewhat lower HOMO–LUMO gap, owing to its unsaturation. To summarise, all computed bisphenolato species have similar stability and bonding mode, the two more sterically hindered species $[(\text{NCN}^{\text{Me}_4})\text{Sb}(\text{OC}_6\text{H}_3\text{-}^t\text{Bu}_2\text{-2,6})_2]$ and





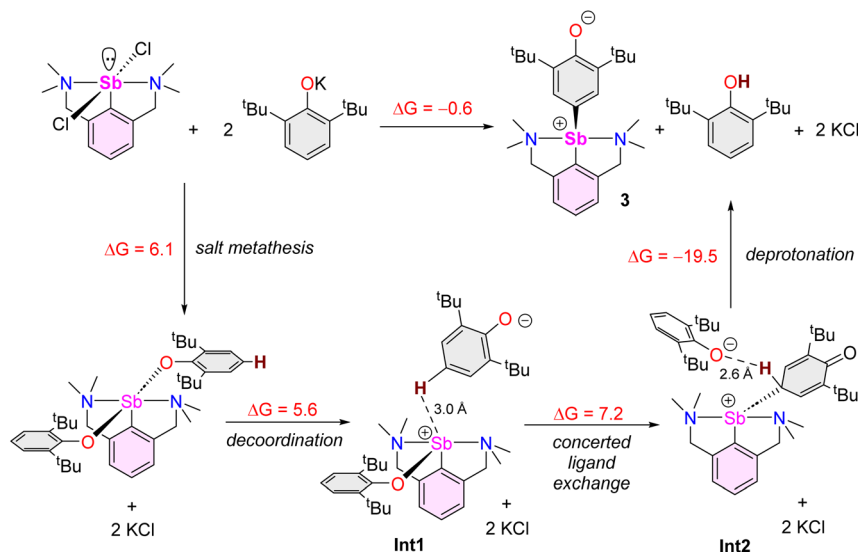
Scheme 5 Reaction of $[\{NCN^{Me_4}\}Sb(C_6H_2-tBu_2-3,5-OH-4)(Cl)]$ (**12**) with $[(2,6-iPr_2-C_6H_3O)K]$, highlighting the two potential mechanistic pathways.

7 having moderately weaker Sb–ligand bonds. Similarly, calculations on **3** and its hypothetical oxyaryl analogues $[\{NCN^{Me_4}\}Sb(C_6H_2-R_2-3,5-O-4)]$ ($R = Me$ and iPr) did not reveal any noticeable stability or structural differences between them (Table S4†). In particular, they all exhibit partial double bond character of the Sb–C(aryl) bond (compare the Sb–C(aryl) and Sb–C(NCN^{Me_4}) Wiberg bond indices in Table S3†), together with non-negligible zwitterionic character (see the Sb and O NBO charges on Table S4†). The single $-NMe_2$ tethered elusive species **6** also presents similar bonding features. Its lower HOMO–LUMO gap is related to its LUMO lower energy, an accepting orbital whose main lobe points in a direction approximately *trans* to N (Fig. S64†), thus explaining the oligomerisation of **6**. Accordingly, formation of **6** from **6** is estimated to be exothermic by 27 kcal mol^{-1} in THF (total energy).

On the basis of our computed results and all the above-reported experimental findings, we suggest that **3** is the outcome of a succession of heterolytic bond cleaving and forming processes (Scheme 6), in a mechanism that bears some resemblance with that described for the double C–H activation of bismuth-bound diphenyl amide.⁶⁸ A process of formation of **3** by reaction of $[\{NCN^{Me_4}\}SbCl_2]$ with $[(2,6-tBu_2-C_6H_3O)K]$ involving free radicals, similar to what was seen with $BiCl_3$ ¹⁰ or proposed in the case of $[\{NCN^{Me_4}\}BiCl_2]$,¹¹ should

be excluded. Instead, our mechanistic proposal involves partially charged species. Complex **3** itself has a pronounced ionic character, as indicated by the conductivity measurements. In this context, it is worth noting that related bismuth(III) bis(phenolate)s bearing NCN tridentate ligands have been shown to present unusually long Bi–OAr interatomic distances in the solid state and, in fact, that they dissociate in THF solution to generate $[\{NCN\}Bi(OAr)]^+[OAr]^-$ separated ion pairs;^{43,69} similar phenomena can be considered for the lighter Sb(III) derivatives. Since the Sb-bis(phenolate)s **5** and **7** can be synthesised and calculations indicate identical stability for **7** and $[\{NCN^{Me_4}\}Sb(OC_6H_3-tBu_2-2,6)_2]$, we propose this later species can be obtained by salt metathesis and acts as a transient, non-observable intermediate (step 1). Our computational search for a subsequent intramolecular rearrangement leading to the formation of an intermediate in which one of the phenolate ligand has undergone dearomatisation and is now bonded through $C_{oxyaryl}$ was unsuccessful. Rather, our calculations indicate that decooordination of one phenolate should occur, followed by its head-to-tail reversal and the formation of a Sb– $C_{oxyaryl}(sp^3)$ bond accompanied by the decooordination of the second phenolate ligand. Two ion pairs have been successively identified during this process, namely **Int1** and **Int2** in Scheme 6. Even if one cannot be certain that the last ion pair





Scheme 6 Proposed pathway for the formation of **3** based on both experimental and computational data. The free energies at 298 K are given in kcal mol⁻¹. They were calculated assuming solvent (THF) effects (see Computational details).

structure is its global energy minimum, it suggests that the leaving phenolate tears out the not-too-remote proton attached to *C_{para}*, to produce 2,6-*t*Bu₂-C₆H₃OH and **3**. Owing to the complexity of the potential energy surface associated with the ion pair ground state and that of the subsequent C–H activation reaction, it was not possible to evaluate the energy barrier of the latter. However, the almost thermoneutral computed thermodynamic balance of the whole process consisting in the formation of **3** agrees with its observation (Scheme 6). Since no important electronic differences were found in our various computed species between those bearing *i*Pr and *t*Bu substituents, the fact that this reaction is not observed in the case of *i*Pr (and OMe) suggests that the formation of **3** is to some extent favoured by the weaker Sb–O bonds in the corresponding bisphenolate intermediate. Yet, we propose that it is chiefly dictated by the specific shape and volume of the *t*Bu groups. They control the specific topology of the **Int2** ion pair (Scheme 6; note that this intermediate is reminiscent of Sb-promoted Friedel–Crafts reactions^{70,71}) in a way that favours, among several types of non-covalent interactions, the proper C_{oxyaryl}–H...OC₆H₃-*t*Bu₂,6 one, thus preparing the C–H activation step. It should be noted that the reaction free energies given in Scheme 6 were computed with solvent (THF) corrections. The computed thermodynamic balance of the whole reaction in vacuum is much higher (23.7 kcal mol⁻¹). This result stresses the importance of solvent effect, which could not be considered explicitly in the calculations; note also that KCl could only be treated, once formed, as a non-interacting spectator. Both factors may play a role in the critical discrimination between the reaction product formations in Scheme 1. Our mechanistic proposal involves ionic intermediates rather than neutral radicals. Consistent with this, the formation of **3** is much slower in toluene than in the more polar THF: where complete formation of the complex is observed within the

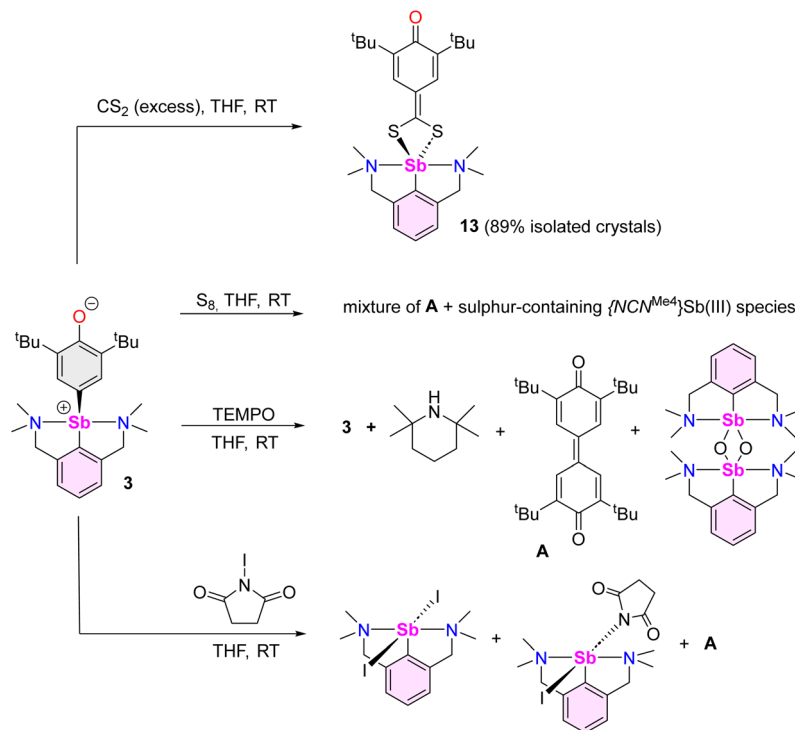
first point of analysis in THF-*d*₈ (*ca.* 10 min), the reaction is not yet over after 24 h in toluene-*d*₈.

Further reactivity studies

Beyond the reactions leading to **12** and **12'**, the reactivity of complex **3** towards various reagents was examined (Scheme 7, Fig. S58–S60†). The nucleophilic nature of the C_{oxyaryl} atom was evidenced by reaction of **3** with excess CS₂. It proceeded to return [{NCN^{Me4}}Sb(S₂C-C₆H₂-*t*Bu₂-3,5-O-4)] (**13**), the product of insertion of CS₂ in the Sb–C_{oxyaryl} bond. By contrast, no reaction takes place between **12** and CS₂, hence highlighting that the greater nucleophilic character of the C_{oxyaryl} atom in zwitterion-like **3** is required for insertion to occur. The synthesis of **13** is a rare example of insertion chemistry into a Sb(III)–C bond.⁷² Our attempts to assess potential radical behaviour in **3** gave mitigated results. The reaction with S₈ gave a mixture of the diphenoquinone **A**, [{NCN^{Me4}}Sb(μ²-S)]₂ and crystalline [{[NCN^{Me4}}Sb(μ-S₅)]₂],⁷³ but we were unable to delineate the details of this reaction. Treatment of **3** with excess TEMPO gave mixtures of unreacted **3**, tetramethylpiperidine (TMP) and crystals of **A** and of [{NCN^{Me4}}Sb(μ²-O)]₂,⁷⁴ the latter being the outcome of abstraction of the oxygen in TEMPO and formation of TMP. Upon reaction with *N*-iodosuccinimide, **3** generated a mixture of [{NCN^{Me4}}Sb(I)(succinimide)] (*ca.* 80%), crystalline [{NCN^{Me4}}SbI₂] (*ca.* 20%) and **A**.

Compound **13** recrystallised as a five-coordinate hemidirected complex (Fig. 6). The metric parameters around Sb(III) match those in [{NCN^{Me4}}Sb(S₃C)]⁷². The C17–O1 bond length (1.248(2) Å) in **13** is marginally shorter than the corresponding one in **3** (1.2708(16) Å). The substantial variations of the C_{*i*}–C_{*i*+1} bond lengths in the S₂C-C₆H₂-*t*Bu₂-3,5-O-4 ligand (compare for instance C17–C18, 1.475(2) Å, and C18–C19, 1.355(2) Å) indicate dearomatisation of the cyclic core, although the atoms S1, S2, C13 to C19 and O1 remain copla-





Scheme 7 Probing the reactivity of 3.

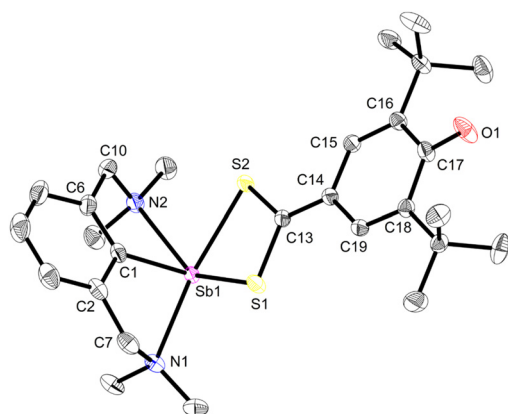


Fig. 6 Representation of the molecular structure of $[\{NCN^{Me_4}\}Sb(S_2C-C_6H_2-tBu_2-3,5-O-4)]$ (**13**). H atoms and non-interacting pyridine molecules omitted for clarity. Ellipsoids at the 50% probability level. Selected distances (Å) and angles (°): Sb1–C1 = 2.1429(17), Sb1–N1 = 2.5492(15), Sb1–N2 = 2.6003(15), Sb1–S1 = 2.5505(5), Sb1–S2 = 2.5723(5), C17–O1 = 1.248(2), S1–C13 = 1.7369(17), S2–C13 = 1.7364(17), C13–C14 = 1.395(2), C14–C15 = 1.429(2), C14–C19 = 1.431(2), C15–C16 = 1.358(2), C16–C17 = 1.474(2), C17–C18 = 1.475(2), C18–C19 = 1.355(2); C1–Sb1–N1 = 72.36(6), C1–Sb1–N2 = 71.69(6), C1–Sb1–S1 = 96.31(5), C1–Sb1–S2 = 96.96(5), N1–Sb1–N2 = 126.95(5), N1–Sb1–S1 = 78.32(4), N1–Sb1–S2 = 144.59(4), S1–Sb1–N2 = 143.01(4), S1–Sb1–S2 = 69.240(15), S2–Sb1–N2 = 77.48(3).

nar. Compound **13** hence adopts a quinoid-like arrangement. By comparison, the C_i-C_{i+1} interatomic distances in $[(p\text{-tol-CS}_2)_3Sb]$, in the range 1.3594(91)–1.4233(88) Å,⁷⁵ agree better with those expected in an aromatic ring. The C13–C14 inter-

atomic distance matches that for a double bond, e.g. it is identical to that in a Pd(methylene-1,1'-dithiolato)-cyclohexanone complex (1.395(6) Å),⁷⁶ and hence the bidentate ligand in **13** must be regarded as a dithiolate.

Conclusion

The mechanism of C–H bond activation leading to the selective formation of $[\{NCN^{Me_4}\}Sb(C_6H_2-tBu_2-3,5-O-4)]$ (**3**) starting from $[\{NCN^{Me_4}\}SbCl_2]$ ⁴² and $[2,6-tBu_2-C_6H_3O]K$ has been probed. Our experimental and DFT data indicate it proceeds through heterolytic bond breaking and bond forming steps involving charged species; no evidence for a radical-based pathway was detected by EPR, and **3** is formed even in the presence of galvinoxyl or TEMPO. This contrasts with the pertaining chemistry of the larger congener, Bi(III).^{10,11} Complex **3** reacts with CS_2 to afford $[\{NCN^{Me_4}\}Sb(S_2C-C_6H_2-tBu_2-3,5-O-4)]$ (**13**), following a migratory insertion step that highlights the highly nucleophilic character of the $C_{oxyaryl}$ atom in **3**. By contrast, $[\{NCN^{Me_4}\}Sb(C_6H_2-tBu_2-3,5-OH-4)]^+[Cl]^-$ (**12**) is inert towards CS_2 .

Representative interatomic distances for **3**, **12**, **13** and 3,3',5,5'-tetra-*tert*-butyldiphenylquinone (**A**) are summarised in Fig. 7.⁷⁷ Key comparative structural features include the followings: (i) the $\{C_6H_2-tBu_2-3,5-OH\}^-$ ligand in **12** features an aromatic ring, with C–O distances as expected for a phenol, (ii) the $S_2C-C_6H_2-tBu_2-3,5-O-4$ ligand in **13** adopts a quinoid-like pattern, with large variations between C–C bond lengths, and (iii) the main bond distances in **3** are intermediate between



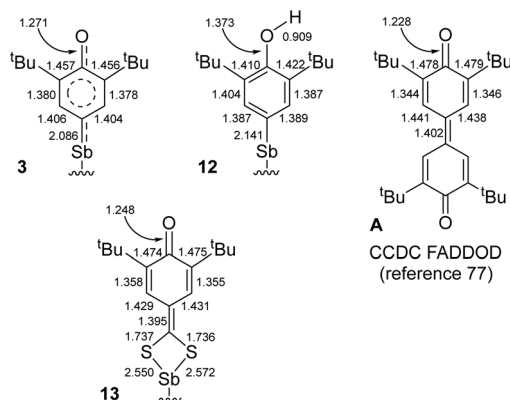


Fig. 7 Selected interatomic distances in **3**, **12**, **13** and **A**. Supporting ligand framework {NCN^{Me4}}[−] and ESD's omitted for clarity.

those in **A** or **13** and those in **12**. The oxyaryl fragment in **3** is hence intermediate between quinoidal and aromatic, and the Sb–C_{oxyaryl} bond features some double bond character. We note that the length of the Sb^V=C(R)(R') double bond reported for a triphenylstibonium(v) ylide, 2.049(4) Å,⁷⁸ is only slightly shorter than that in **3**; however, the different oxidation states between the compounds limits the usefulness of this comparison.

Driven by the impressively successful implementation of bismuth in molecular catalysis in recent years,^{14–18} and by the intriguingly different behaviours of antimony(III) and bismuth(III) towards phenolates disclosed herein, we will next assess how **3** and related phenolato complexes can be exploited in efficient catalytic manifolds, and whether further specificities of antimony with respect to its heavier congener, bismuth, can be highlighted. The discovery and the understanding of elementary reactions such as those presented herein, associated to the mechanistic changeover observed between congeneric Sb and Bi complexes, constitute a necessary first step in developing applications in fields such as catalysis.

Data availability

The experimental and theoretical data that supports the findings and conclusions of this study are available in the ESI† of this article.

Author contributions

G. D. carried out the synthetic and spectroscopic experimental work and participated to the data analysis and their processing, and to the preparation of the manuscript. M. C. solved the crystallographic structures. A. P. participated to the synthetic experimental work. S. Kahlal and J.-Y. Saillard performed the theoretical calculations and their analysis, and participated to the preparation of the manuscript. C. Silvestru par-

ticipated to data analysis, planning of experimental work and manuscript preparation. Y. Sarazin was the lead scientist; he secured the funding, participated to experiment planning, to data analysis, and to the writing of the manuscript.

Conflicts of interest

There are no conflicts to declare.

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- 66 We have checked by NMR monitoring in THF-*d*₈ that complex **1** does not react with 2,6-*t*Bu₂-C₆H₃OH (3 equivalents), even after overnight reflux.
- 67 Consistent with this conclusion, the reaction of **12** with KN(SiMe₃)₂ in THF-*d*₈ gives **3**, KCl and HN(SiMe₃)₂, without formation of [{NCN^{Me4}}Sb{N(SiMe₃)₂}(C₆H₂-*t*Bu₂-3,5-O-4)K], see Fig. S56.†
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