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Synthesis, properties and structural features of molybdenum(v) oxide trichloride complexes with neutral chalcogenoether ligands†

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Complexes of oxotrichloromolybdenum(v) with neutral group 16 donor ligands, $[\text{MoOCl}_3(\text{L-L})]$ ($\text{L-L} = \text{RS}(\text{CH}_2)_2\text{SR}$, $\text{R} = {}^i\text{Pr}$, Ph ; $\text{MeS}(\text{CH}_2)_3\text{SMe}$; $\text{MeSe}(\text{CH}_2)_2\text{SeMe}$; $\text{MeSe}(\text{CH}_2)_3\text{SeMe}$), $\{[\text{MoOCl}_2(\text{EMe}_2)]_2(\mu\text{-Cl})_2\}$ ($\text{E} = \text{S}$, Se , Te), $\{[\text{MoOCl}_3]_2(o\text{-C}_6\text{H}_4(\text{EMe}_2)_2)\}_n$ ($\text{E} = \text{Se}$ or Te) and $\{[\text{MoOCl}_3]_2(\text{MeTe}(\text{CH}_2)_3\text{TeMe})\}_n$, have been obtained by reaction of the ligands with $[\text{MoOCl}_3(\text{thf})_2]$ or MoOCl_3 in either CH_2Cl_2 or toluene, and characterised by microanalysis, IR and UV-visible spectroscopy and magnetic measurements. The telluroethers are the first examples containing Mo in a positive oxidation state. X-ray crystal structures are reported for the six-coordinate *fac*- $[\text{MoOCl}_3(\text{MeS}(\text{CH}_2)_3\text{SMe})]$, *mer*- $[\text{MoOCl}_3({}^i\text{PrS}(\text{CH}_2)_2\text{S}^i\text{Pr})]$ and *mer*- $[\text{MoOCl}_3(\text{MeSe}(\text{CH}_2)_2\text{SeMe})]$, as well as the six-coordinate chloride-bridged dimers, $\{[\text{MoOCl}_2(\text{SMe}_2)]_2(\mu\text{-Cl})_2\}$ and $\{[\text{MoOCl}_2(\text{SeMe}_2)]_2(\mu\text{-Cl})_2\}$. The structure of the mixed-valence decomposition product, $[\text{Mo}^{\text{IV}}\text{Cl}(o\text{-C}_6\text{H}_4(\text{TeMe})_2)_2(\mu\text{-O})\text{Mo}^{\text{VO}}\text{OCl}_4]$, was also determined. In toluene solution MoOCl_4 is reduced by $\text{MeS}(\text{CH}_2)_3\text{SMe}$ to produce the Mo(v) complex, $[\text{MoOCl}_3(\text{MeS}(\text{CH}_2)_3\text{SMe})]$. Crystal structures of the previously unknown diphosphine analogue, $[\text{MoOCl}_3(\text{Me}_2\text{P}(\text{CH}_2)_2\text{PMe}_2)]$, and the mixed-valence derivative $[\text{Mo}^{\text{IV}}\text{Cl}(\text{Me}_2\text{P}(\text{CH}_2)_2\text{PMe}_2)_2(\mu\text{-O})\text{Mo}^{\text{VO}}\text{OCl}_4]$ are also reported for comparison and help to clarify earlier contradictory literature reports. In contrast to the dimeric EMe_2 complexes, $\{[\text{MoOCl}_2(\text{EMe}_2)]_2(\mu\text{-Cl})_2\}$, PMe_3 forms the monomeric complex, *fac*- $[\text{MoOCl}_3(\text{PMe}_3)_2]$.

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Introduction

The coordination chemistry of high oxidation state molybdenum halides and oxide halides was first explored in some detail in the 1970's, with the emphasis on neutral N- and O-donor ligands^{1–4} and with much of the impetus coming from modelling of the metal sites in molybdenum enzymes and applications in catalysis.^{4–6} Interest in high oxidation molybdenum complexes bearing sulfur donor ligands stems in part from the presence of Mo–S coordination in the molybdenum-containing enzymes, nitrate reductase, sulfite oxidase and Fe–Mo nitrogenases which involve (anionic) cysteine or sulfide ligands.^{4–6} The chemistry with neutral P- and As-donor ligands with Mo(v) has also been investigated,^{7–11} but sulfur-

based ligands were mostly represented by charged thiolate and dithiocarbamate ligands.^{2–4} More recent work has reported a series of extremely moisture sensitive Mo(vi) complexes $[\text{MoO}_2\text{X}_2(\text{dithioether})]$ ($\text{X} = \text{Cl}$ or Br ; dithioether = $\text{RS}(\text{CH}_2)_2\text{SR}$, $\text{R} = \text{Me}$, Et , ${}^i\text{Pr}$), which have distorted octahedral structures with the sulfur donor atoms *trans* to $\text{Mo}=\text{O}$;^{12,13} there are also some thia-macrocyclic analogues.^{14,15} Complexes of the type $[\text{MoOCl}_3(\text{dithioether})]$ were briefly described in the 1970's, characterised only by microanalysis and IR spectroscopy, but the structures and isomer(s) present were not established.^{15,16} There is a single preliminary report of a selenoether complex of MoOCl_3 ,¹⁷ but no known telluroether complexes.

We have recently examined the complexes of WOCl_4 , WOCl_3 , WSCl_4 and WSCl_3 with mono- and di-thio- and -seleno-ethers, and found that W(vi) or W(v) complexes could be isolated depending upon the reaction conditions. We also showed that selected dithioether complexes, for example $[(\text{WSCl}_4)_2(\mu\text{-}{}^i\text{PrSCH}_2\text{CH}_2\text{S}^i\text{Pr})]$, can function as single source LPCVD (low pressure chemical vapour deposition) reagents for the growth of thin films of WS_2 , an important semiconducting material.¹⁸ In contrast, very little data on the molybdenum chalcogenide halides or their coordination complexes exists.¹⁹ The crystal structures of two forms of MoSCl_3 obtained from crystals grown at high temperature found that both contain

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† Electronic supplementary information (ESI) available: Crystallographic parameters (Table S1), IR and UV/visible spectra for the new complexes. CCDC 2050667: $\{[\text{MoOCl}_2(\text{SeMe}_2)]_2(\mu\text{-Cl})_2\}$, 2050668: $[\text{MoOCl}_3(\text{Me}_2\text{P}(\text{CH}_2)_2\text{PMe}_2)]$, 2050669: $[\text{MoOCl}_3(\text{PhS}(\text{CH}_2)_2\text{SPh})]$, 2050670: $[\text{MoOCl}_3(\text{MeSe}(\text{CH}_2)_2\text{SeMe})]$, 2050671: $[\text{MoOCl}_3({}^i\text{PrS}(\text{CH}_2)_2\text{S}^i\text{Pr})]$, 2050672: $\{[\text{MoOCl}_2(\text{SMe}_2)]_2(\mu\text{-Cl})_2\}$, 2050673: $[\text{MoCl}(\text{Me}_2\text{P}(\text{CH}_2)_2\text{PMe}_2)_2(\mu\text{-O})(\text{MoOCl}_4)]$, 2050674: $[\text{MoOCl}_3(\text{MeS}(\text{CH}_2)_3\text{SMe})]$, 2050891: $[\text{MoCl}(o\text{-C}_6\text{H}_4(\text{TeMe})_2)_2(\mu\text{-O})\text{MoOCl}_4]\cdot\text{CH}_2\text{Cl}_2$. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/d1dt00038a



Mo(IV) as Mo₂ units and disulfide groups, and not Mo(V).²⁰ It is unclear if MoSCl₃ prepared at low temperatures from MoCl₅ and S(SiMe₃)₂ or Sb₂S₃ contains Mo(V),^{21,22} while MoSCl₄ is unknown.¹⁹

In order to allow comparisons with the WOCl₄, WOCl₃, WSCL₄ and WSCL₃ chemistry, we have examined the chemistry of MoOCl₃ with neutral chalcogenoethers and report here complexes of mono- and bi-dentate thio-, seleno- and telluroethers. Data on diphosphine analogues, which clarifies some of the (inconsistent) earlier studies,^{7–9} is also presented.

Results and discussion

Scheme 1 shows the range of chalcogenoether complexes of Mo(V) prepared in this study and the different structure types observed.

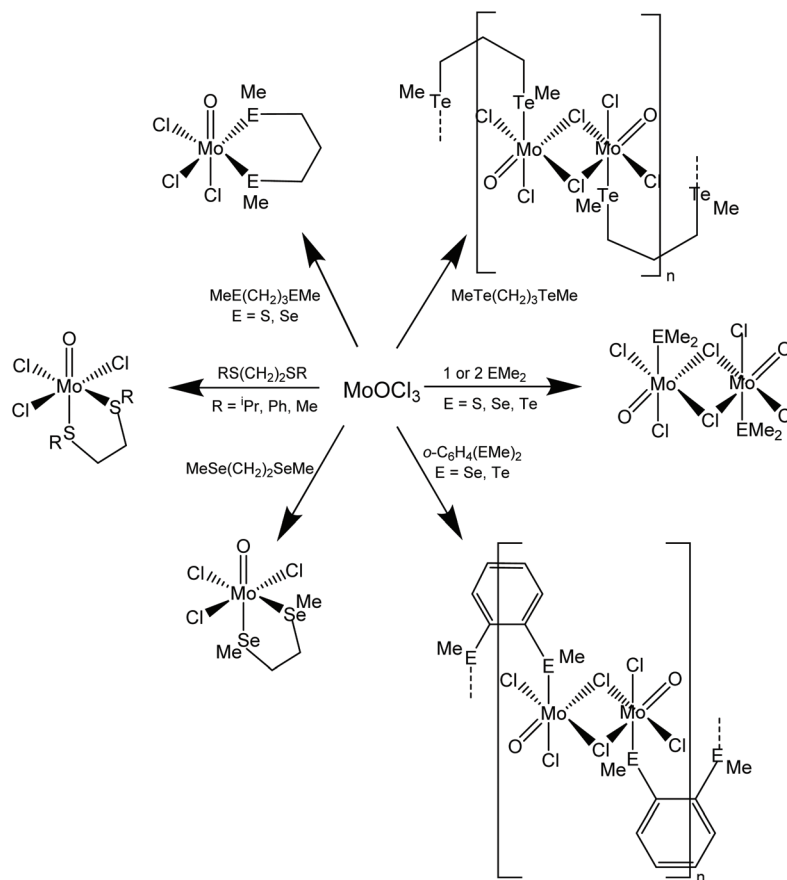
Dithio- and diseleno-ether complexes

The reaction of [MoOCl₃(thf)₂] with MeS(CH₂)₃SMe or ⁱPrS(CH₂)₂SⁱPr in dry CH₂Cl₂ produced moisture sensitive, green [MoOCl₃(dithioether)] complexes. Structures of both species were determined and revealed that [MoOCl₃{MeS(CH₂)₃SMe}] (six-membered chelate ring) was the *fac* isomer, whilst

[MoOCl₃{ⁱPrS(CH₂)₂SⁱPr}] (five-membered chelate ring) was the *mer*-isomer (Fig. 1). The reason for the different isomers with the five- and six-membered rings is uncertain, although the difference in the S–Mo–S chelate angles of ~20° is notable. The behaviour replicates that found with the tungsten(V) analogues, *fac*-[WOCl₃{MeS(CH₂)₃SMe}] and *mer*-[WOCl₃{MeS(CH₂)₂SMe}].¹⁸ The bond lengths within the two structures show the expected short Mo=O of ~1.67 Å and that the Mo–Cl and Mo–S *trans* to Mo=O are longer than the other bonds of each type, indicating the high *trans*-influence of the Mo=O bond.

The reaction of MoOCl₄ with MeS(CH₂)₃SMe in dry toluene gave a green product with an identical IR spectrum to that of *mer*-[MoOCl₃{MeS(CH₂)₃SMe}] and the X-ray structure determination of a crystal obtained from the MoOCl₄ synthesis route (Method 2) indeed confirmed it to be the Mo(V) complex. The structural data were identical to that in Table S1,† and hence are not reported, but confirm that thioether ligands reduce MoOCl₄ to MoOCl₃ complexes, similar to the behaviour reported with some O- and N-donor ligands.²³

The weaker σ-donor PhS(CH₂)₂SPh failed to displace the thf from [MoOCl₃(thf)₂], but it reacted with a suspension of MoOCl₃ in CH₂Cl₂ to form brown [MoOCl₃{PhS(CH₂)₂SPh}]. The crystal structure of this complex showed it to be the *mer*-



Scheme 1 Methods for the synthesis of the Mo(V) chalcogenoether complexes obtained from MoOCl₃. Note that for some of the alkyl-substituted dithioether and diselenoether complexes [MoOCl₃(thf)₂] was used as the Mo(V) source – see discussion below and Experimental.



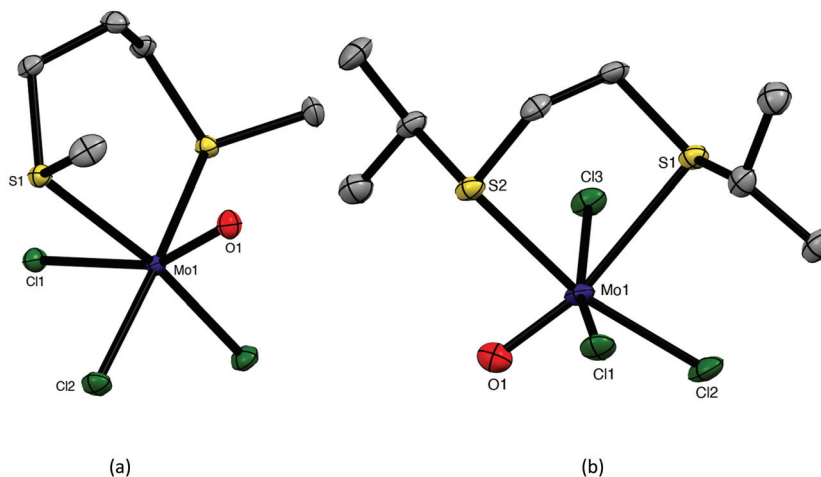


Fig. 1 Crystal structures of *fac*-[MoOCl₃(MeS(CH₂)₃SMe)] (a) and *mer*-[MoOCl₃(^tPrS(CH₂)₂S^tPr)] (b) showing the atom numbering scheme. Ellipsoids are shown at 50% probability, hydrogen atoms omitted for clarity. Selected bond lengths (Å) and angles (°): (a) Mo1–Cl1 = 2.4540(2), Mo1–Cl2 = 2.3451(2), Mo1–Cl3 = 2.3451(4), Mo1–O1 = 1.674(1), Mo1–S1 = 2.5388(3), Cl1–Mo1–Cl2 = 94.52(2), Cl2–Mo1–Cl3 = 92.55(3), Cl2–Mo1–O1 = 101.87(4), Cl3–Mo1–O1 = 101.87(4), S1–Mo1–S1 = 97.95(2); (b) Mo1–Cl1 = 2.3578(8), Mo1–Cl2 = 2.3378(8), Mo1–Cl3 = 2.3618(7), Mo1–O1 = 1.671(2), Mo1–S1 = 2.8298(8), Mo1–S2 = 2.5665(7), Cl1–Mo1–Cl2 = 91.25(3), Cl1–Mo1–O1 = 98.99(8), Cl1–Mo1–Cl3 = 89.69(3), Cl2–Mo1–O1 = 98.99(8), S1–Mo1–S2 = 78.83(2).

isomer (Fig. 2), which suggests that the ability to form a five-membered chelate ring with a smaller chelate angle (S1–Mo1–S2 = 78.55(3)°) may be an important factor influencing the isomer formed. The structure also reveals a very markedly longer Mo–S_{trans}O = 2.911(1) Å, which compares with Mo–S_{trans}Cl = 2.531(1) Å, showing the high *trans*-influence of the Mo=O bond on the weaker aryl thioether donor ligand.

Brownish diselenoether complexes, [MoOCl₃(diselenoether)] (diselenoether = MeSe(CH₂)₂SeMe, MeSe(CH₂)₃SeMe), were obtained from reaction of the ligands with MoOCl₃ or

[MoOCl₃(thf)₂] in a 1 : 1 molar ratio, but *o*-C₆H₄(SeMe)₂ did not displace thf from [MoOCl₃(thf)₂]. The reaction of MeSeCH₂SeMe with MoOCl₃ produced a black oily decomposition product. However, the 1 : 1 reaction of *o*-C₆H₄(SeMe)₂ with MoOCl₃ in CH₂Cl₂ gave a brown product for which the microanalytical data indicated a 2 : 1 MoOCl₃ : diselenoether stoichiometry. This is discussed along with the similar ditelluroether complexes below. The X-ray crystal structure of *mer*-[MoOCl₃{MeSe(CH₂)₂SeMe}] was obtained (Fig. 3).

The five complexes described have room temperature magnetic moments of ~1.7 B.M., similar to other MoOCl₃ complexes,^{1,2,8,9} and close to the spin-only value expected for a

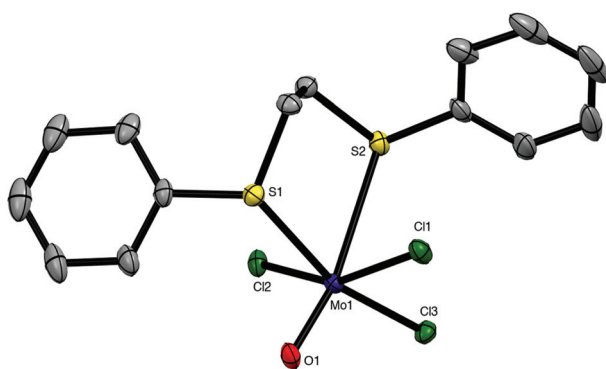


Fig. 2 Crystal structure of *mer*-[MoOCl₃(PhS(CH₂)₂SPh)] showing the atom numbering scheme. Ellipsoids are shown at 50% probability and hydrogen atoms are omitted for clarity. Note that the O/Cl exhibited disorder, which was modelled with split atom sites, refined to occupancies of 0.53 : 0.47. Only the major form is shown. Selected bond lengths (Å) and angles (°): Mo1–Cl1 = 2.324(1), Mo1–Cl2 = 2.394(1), Mo1–Cl3 = 2.311(3), Mo1–O1 = 1.706(2), Mo1–S1 = 2.531(1), Mo1–S2 = 2.911(1), Cl1–Mo1–Cl3 = 89.93(7), Cl1–Mo1–O1 = 102.2(5), O1–Mo1–Cl2 = 101.1(5), O1–Mo1–Cl3 = 106.6(3), Cl2–Mo1–Cl3 = 91.28(7), Cl2–Mo1–S1 = 88.39(4), Cl1–Mo1–S2 = 81.49(4), S1–Mo1–S2 = 78.55(3).

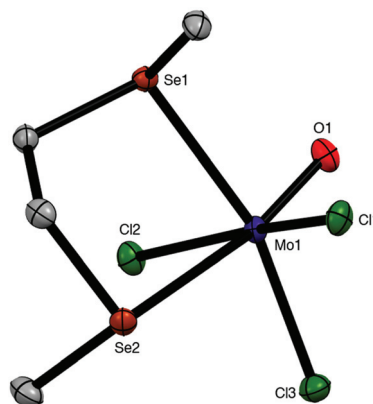


Fig. 3 Crystal structure of *mer*-[MoOCl₃{MeSe(CH₂)₂SeMe}] showing the atom numbering scheme. Ellipsoids are shown at 50% probability and hydrogen atoms are omitted for clarity. Selected bond lengths (Å) and angles (°): Mo1–Cl1 = 2.3553(5), Mo1–Cl2 = 2.3517(5), Mo1–Cl3 = 2.3453(5), Mo1–O1 = 1.673(1), Mo1–Se1 = 2.6564(2), Mo1–Se2 = 2.8937(3), Cl1–Mo1–Cl3 = 90.35(2), Cl1–Mo1–O1 = 98.98(5), Cl2–Mo1–Cl3 = 90.65(2), Cl3–Mo1–O1 = 107.36(5), Se1–Mo1–Se2 = 79.76(1).



d^1 complex. This indicates that any orbital contribution is quenched by the very asymmetric field of the molybdenum environment.²⁴ The IR spectra show very strong single bands due to $\nu(\text{Mo}=\text{O})$ in the range 950–980 cm^{-1} , as well as strong overlapping bands at 355–300 cm^{-1} assigned as Mo–Cl modes, but do not appear to readily distinguish the isomer present. The UV/visible spectra of the solids show a clear band at 13 000–14 000 cm^{-1} and a second band or shoulder at ~19 000–21 000 cm^{-1} . Assuming C_{4v} symmetry (the actual metal centre symmetry is lower) and placing Mo=O as the dominant contribution along the four-fold axis, leads to the assignment as the d–d bands as $^2B_2 \rightarrow ^2E$ and $^2B_2 \rightarrow ^2B_1$, respectively.²⁵ The intense absorptions $>20\,000\text{ cm}^{-1}$, assigned as charge transfer bands, are less clearly resolved, but based upon the usual ligand electronegativities,²⁵ we assign the first intense feature (~21 000–22 000 cm^{-1}) as S/Se(π) \rightarrow Mo(d) and the broad overlapping features at ~25 000–30 000 cm^{-1} as Cl(π) \rightarrow Mo(d). The complexity of the electronic spectra in compounds of this type is shown by a combined UV/visible absorption, MCD and DFT study of $[\text{MoOCl}_3\{\text{Ph}_2\text{P}(\text{CH}_2)_2\text{PPh}_2\}]$;²⁶ here we are using the spectra to confirm the presence of Mo(v) in the isolated complexes.

Dimethylchalcogenides (EMe₂, E = S, Se, Te)

Neither SMe₂ or SeMe₂ was found to displace thf from $[\text{MoOCl}_3(\text{thf})_2]$. However, reaction of a suspension of MoOCl₃ in dry CH₂Cl₂ with 2 equivalents of EMe₂ produced complexes with a 1 : 1 Mo : EMe₂ empirical composition, MoOCl₃(EMe₂) (E = S, Se). There was no evidence for the formation of the 1 : 2 $[\text{MoOCl}_3(\text{EMe}_2)_2]$ complexes. Crystals of both MoOCl₃(EMe₂) (E = S, Se) complexes were obtained and the structures, which are isomorphous (Fig. 4), showed them to be chloride-bridged dimers, with six-coordinate Mo(v) centres, *i.e.* $[\{\text{MoOCl}_2(\text{EMe}_2)\}_2(\mu\text{-Cl}_2)]$ (E = S, Se).

The Mo=O bonds (~1.65 Å) are trans to asymmetrically bound (by ~0.4 Å) bridging chlorides, with the EMe₂ groups arranged anti and perpendicular to the Mo₂Cl₄O₂ plane. The geometries are very similar to those found in $[(\text{MoOCl}_2\text{L})_2(\mu\text{-Cl})_2]$ (L = O=C(H)OME, thf, O = CEt₂).^{27–29}

The reaction of MoOCl₃ with TeMe₂ in toluene produced brown $[\{\text{MoOCl}_2(\text{TeMe}_2)\}_2(\mu\text{-Cl})_2]$, which is the first Mo(v) complex with a neutral tellurium donor ligand. Crystals were not obtained from this complex due to poor solubility and limited stability in solution, but spectroscopically it is very similar to the other EMe₂ complexes. The failure to produce the six-coordinate monomers, $[\text{MoOCl}_3(\text{EMe}_2)_2]$, even in the presence of excess EMe₂, shows that the molybdenum(v) prefers to bind a chloride from another molecule, creating the bridged dimer structure, and is consistent with the weak donor properties of the EMe₂. The dimers are clearly distinguished from the $[\text{MoOCl}_3(\text{dichalcogenoether})]$ monomers by their IR spectra, with the dimers showing a strong $\nu(\text{Mo}=\text{O})$ vibration in the range at 985–1005 cm^{-1} (higher frequency than in the monomeric $[\text{MoOCl}_3(\text{dichalcogenoether})]$ type) and terminal Mo–Cl modes 360–310 cm^{-1} ; weaker bands in the region ~270–250 cm^{-1} and absent in the spectra of the $[\text{MoOCl}_3(\text{dichalcogenoether})]$ monomers, may be due to the chloride bridges. The magnetic moments of ~1.7 B.M./Mo confirm the Mo(v) assignment and the absence of any magnetic interactions between the molybdenum centres.

Ditelluroethers

The reaction of *o*-C₆H₄(SeMe)₂, *o*-C₆H₄(TeMe)₂ and MeTe(CH₂)₃TeMe (L–L) with MoOCl₃ in a 1 : 1 molar ratio in CH₂Cl₂ failed to produce the expected $[\text{MoOCl}_3(\text{L-L})]$ type complexes. Instead, dark brown complexes, identified by microanalysis as $[(\text{MoOCl}_2)_2(\text{L-L})]$, were obtained. Once isolated the compounds are very poorly soluble in CH₂Cl₂ and many attempts to

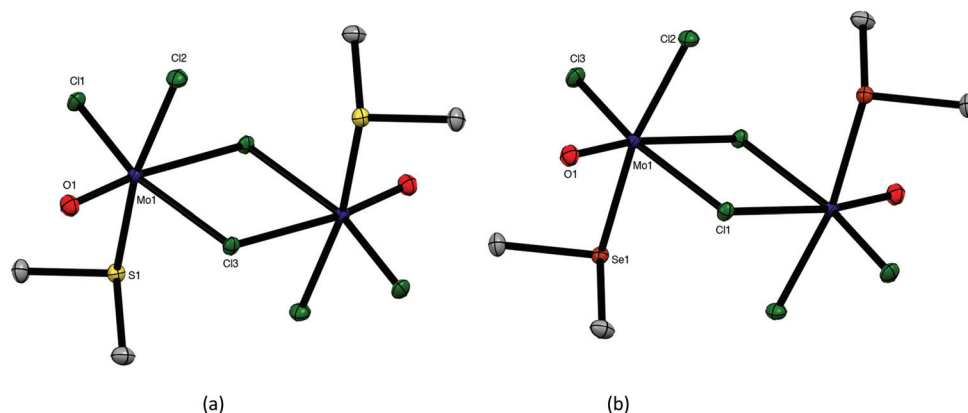


Fig. 4 Crystal structures of $[\{\text{MoOCl}_2(\text{SMe}_2)\}_2(\mu\text{-Cl})_2]$ (a) and $[\{\text{MoOCl}_2(\text{SeMe}_2)\}_2(\mu\text{-Cl})_2]$ (b) showing the atom numbering scheme. Ellipsoids are shown at 50% probability and hydrogen atoms are omitted for clarity. Selected bond lengths (Å) and angles (°): (a) Mo1–Cl1 = 2.3264(2), Mo1–Cl2 = 2.3341(3), Mo1–Cl3 = 2.3953(3), Mo1–Cl3' = 2.7942(2), Mo1–O1 = 1.6515(8), Mo1–S1 = 2.5537(3), Cl1–Mo1–O1 = 102.48(1), Cl2–Mo1–O1 = 102.08(3), Cl3–Mo1–O1 = 99.55(3), Cl2–Mo1–Cl3 = 85.77(1), Cl1–Mo1–Cl3 = 92.340(9), Cl1–Mo1–S1 = 85.233(9), O1–Mo1–S1 = 92.67(3), Cl3–Mo1–Cl3 = 77.520(8); (b) Mo1–Cl1 = 2.4024(4), Mo1–Cl2 = 2.3385(4), Mo1–Cl3 = 2.3299(4), Mo1–Cl1' = 2.7927(4), Mo1–O1 = 1.653(1), Mo1–Se1 = 2.6647(3), Cl1–Mo1–O1 = 98.89(4), Cl2–Mo1–O1 = 103.10(4), Cl3–Mo1–O1 = 102.38(4), Cl2–Mo1–Cl3 = 92.32(2), Se1–Mo1–Cl1 = 87.438(10), O1–Mo1–Se1 = 92.89(4).



produce crystals for an X-ray structure determination have been unsuccessful. However, the magnetic moments of ~ 1.7 B. M./Mo and the UV-visible spectra of these solids are consistent with their formulation as six-coordinate oxo-molybdenum(v) complexes.

The UV-visible spectra of the ditelluroether complexes show a d-d band at $\sim 14\,000\text{ cm}^{-1}$ (${}^2B_2 \rightarrow {}^2E$); a second more intense feature $18\,000\text{--}20\,000\text{ cm}^{-1}$ may be the second d-d band (${}^2B_2 \rightarrow {}^2B_1$), but given the lower electronegativity of Te²⁴ is probably the Te(π) \rightarrow Mo(d) charge transfer transition, which obscures the d-d band.

The IR spectra are significantly different to those of $[\text{MoOCl}_3(\text{L-L})]$ (L-L = dithioether or diphosphine),^{8,9} but are similar to those of $[\{\text{MoOCl}_2(\text{EMe}_2)\}_2(\mu\text{-Cl})_2]$. In particular, the $\nu(\text{Mo}=\text{O})$ vibrations are at higher frequency ($985\text{--}1000\text{ cm}^{-1}$), and in addition to several terminal $\nu(\text{Mo}-\text{Cl})$ modes $\sim 320\text{--}300\text{ cm}^{-1}$, also show a peak $\sim 250\text{ cm}^{-1}$, probably due to a chloride bridge. In the absence of a crystal structure, the geometries cannot be established unequivocally, but the spectroscopic data (and insolubility) are consistent with a structure type similar to those in $[\{\text{MoOCl}_2(\text{EMe}_2)\}_2(\mu\text{-Cl})_2]$, with the EMe_2 ligands replaced by bridging ditelluroethers, leading to the formulation as an oligomer, $[(\text{MoOCl}_2)_2(\mu\text{-Cl})_2(\mu\text{-ditelluroether})]_n$. There are several literature examples of Group 16 ligands with *o*-phenylene backbones adopting a bridging mode, authenticated by X-ray crystal structures.^{30–32}

The brown solution from the preparation of $[\{\text{MoOCl}_3\}_2\{o\text{-C}_6\text{H}_4(\text{TeMe})_2\}_n]$ also deposited a few dark green crystals, which were shown by X-ray crystallographic analysis to be the mixed valence complex, $[\text{Mo}^{\text{IV}}\text{Cl}\{o\text{-C}_6\text{H}_4(\text{TeMe})_2\}_2(\mu\text{-O})\text{Mo}^{\text{V}}\text{OCl}_4]$ (Fig. 5). This complex contains a Mo(IV) centre coordinated to

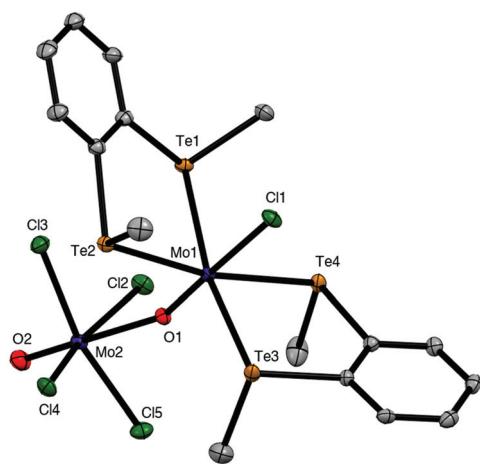


Fig. 5 Crystal structure of $[\text{MoCl}(o\text{-C}_6\text{H}_4(\text{TeMe})_2)_2(\mu\text{-O})\text{MoOCl}_4]\cdot\text{CH}_2\text{Cl}_2$ showing the atom numbering scheme. Ellipsoids are shown at 50% probability and hydrogen atoms and solvent are omitted for clarity. Selected bond lengths (Å) and angles ($^\circ$): Mo1–Cl1 = 2.4486(7), Mo2–Cl2 = 2.3640(7) – 2.3930(7), Mo1–Te1 = 2.7432(3) – 2.7822(3), Mo1–O1 = 1.704(2), Mo2–O1 = 2.370(2), Mo2–O2 = 1.655(2), O2–Mo2–Cl2 = 78.4(1) – 82.2(1), Te–Mo1–Te_(chelate) = 85.316(8), 86.185(8), Te1 – 4–Mo1–Cl1 = 79.647(18)–89.939(18), Cl1–Mo1–O1 = 177.91(7), Mo1–O1–Mo2 = 159.63.1(11).

two chelating ditelluroethers, a terminal chloride and an Mo=O group, which forms a very asymmetric bridge to a square pyramidal MoOCl_4^- anion, with Mo1–O1 = 1.705(4) Å and Mo2–O1 = 2.368(4) Å. These bond distances may be compared with the terminal Mo=O bond distance (Mo2–O2 = 1.659(5) Å) in the latter. This complex appears to be the first structurally characterised molybdenum-ditelluroether complex in a positive formal oxidation state of the metal; all previously reported complexes are substituted carbonyls.^{33,34} Analogous complexes with some diphosphine and diarsine ligands have been reported,^{8,9} and the structure of (the previously unknown) $[\text{MoCl}\{\text{Me}_2\text{P}(\text{CH}_2)_2\text{PMe}_2\}_2(\mu\text{-O})(\text{MoOCl}_4)]$ is discussed below. The crystals of $[\text{Mo}^{\text{IV}}\text{Cl}\{o\text{-C}_6\text{H}_4(\text{TeMe})_2\}_2(\mu\text{-O})\text{Mo}^{\text{V}}\text{OCl}_4]$ result from a redox reaction, and its structure is not consistent with the spectroscopic data on the bulk $[\{\text{MoOCl}_3\}_2\{o\text{-C}_6\text{H}_4(\text{TeMe})_2\}_n]$. The failure to isolate mononuclear $[\text{MoOCl}_3(\text{L-L})]$ complexes with chelating *o*-C₆H₄(SeMe)₂, *o*-C₆H₄(TeMe)₂ and MeTe(CH₂)₃TeMe, seems analogous to the case of $[\{\text{MoOCl}_2(\text{EMe}_2)\}_2(\mu\text{-Cl})_2]$, where the Mo(v) centre prefers to form chloride bridges rather than coordinate to a second, weakly donating chalcogenoether.

Phosphine complexes

The coordination behaviour of the chalcogenoether ligands to MoOCl_3 has both significant analogies and differences to that of some phosphine ligands, making for informative comparisons. Pink or red complexes $[\text{MoOCl}_3(\text{diphosphine})]$ (diphosphine = $\text{Ph}_2\text{P}(\text{CH}_2)_2\text{PPh}_2$, *cis*- $\text{Ph}_2\text{PCH}=\text{CHPPh}_2$, *o*-C₆H₄(PPh₂)₂) were reported in the 1970s and confirmed by IR, UV/visible spectroscopy and magnetic measurements as Mo(v) compounds.^{7–9} No structures were obtained, but EPR spectra supported *fac* octahedral isomers.^{8,9} A second (brown) form with $\text{Ph}_2\text{P}(\text{CH}_2)_2\text{PPh}_2$ and *cis*- $\text{Ph}_2\text{PCH}=\text{CHPPh}_2$ obtained by refluxing the red form in alcohol for several hours, had similar, but not identical, spectroscopic properties; Isovitsch *et al.*,¹⁰ confirmed the crystal structure of the red form of the $\text{Ph}_2\text{P}(\text{CH}_2)_2\text{PPh}_2$ complex as the *fac* isomer. In the present work we prepared the new complex $[\text{MoOCl}_3\{\text{Me}_2\text{P}(\text{CH}_2)_2\text{PMe}_2\}]$ from $[\text{MoOCl}_3(\text{thf})_2]$ and confirmed the *fac* geometry by a crystal structure (Fig. 6). The spectroscopic data on this complex (Experimental section) are in good agreement with that of the red isomers with other diphosphines.^{7–9} Notably, the five-membered chelate ring diphosphine complexes are *fac* isomers, contrasting with the *mer*- $[\text{MoOCl}_3(\text{dichalcogenoether})]$ described above.

The nature of the brown “isomers” is not entirely clear, but the original study⁸ of the red and brown forms of $[\text{MoOCl}_3\{\text{cis-Ph}_2\text{PCH}=\text{CHPPh}_2\}]$ showed they had identical EPR spectra with coupling to equivalent phosphorus donors, *i.e.* were both *fac* forms. Hence the brown form seems likely to be the red isomer co-crystallised with a second complex, probably an EPR silent Mo(IV) species. The presence of varying amounts of a co-crystallised second species would account for the various (small) differences reported by other workers.^{7,8,10} Similar problems, including X-ray structures with a variety of bond lengths for apparently the same complex, led to the proposal



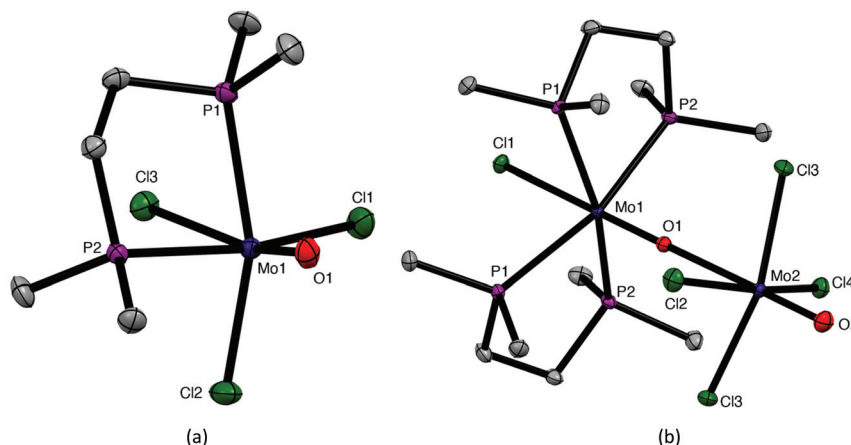


Fig. 6 Crystal structures of $[\text{MoOCl}_3(\text{Me}_2\text{PCH}_2\text{CH}_2\text{PMe}_2)]$ (a) and $[\text{MoCl}(\text{Me}_2\text{PCH}_2\text{CH}_2\text{PMe}_2)_2(\mu\text{-O})\text{MoOCl}_4]$ (b) showing the atom numbering scheme. Ellipsoids are shown at 50% probability and hydrogen atoms are omitted for clarity. Selected bond lengths (Å) and angles ($^\circ$): (a) Mo1–Cl1 = 2.3824(9), Mo1–Cl2 = 2.383(1), Mo1–Cl3 = 2.5011(8), Mo1–O1 = 1.680(2), Mo1–P1 = 2.5260(2), Mo1–P2 = 2.5250(8), Cl1–Mo1–Cl2 = 96.98(3), Cl1–Mo1–O1 = 100.19(8), Cl2–Mo1–Cl3 = 88.76(3), Cl2–Mo1–O1 = 104.66(8), P1–Mo1–P2 = 78.84(3); (b) Mo1–Cl1 = 2.5186(2), Mo1–O1 = 1.703(4), Mo1–P1 = 2.5131(2), Mo1–P2 = 2.5139(3), Mo2–O1 = 2.394(2), Mo2–O2 = 1.657(8), Mo2–Cl2 = 2.3573(4) – 2.3824(3), Cl1–Mo1–P1 = 78.92(2), Cl1–Mo1–P2 = 83.26(4), O2–Mo2–Cl2 = 97.57(2) – 98.89(4), P1–Mo1–P2 = 80.16(4), Mo1–O1–Mo2 = 178.05(8).

of *bond-stretch or distortional isomerism* in some other early d-block complexes, a concept subsequently considered to be erroneous.³⁵

Pink or purple complexes with microanalyses indicating a $[\text{MoCl}_{2.5}\text{O}(\text{diphosphine})]$ were isolated in some systems^{7–9} and were formulated as the ionic Mo(IV)–Mo(V) species $[\text{Mo}^{\text{IV}}\text{OCl}(\text{diphosphine})_2][\text{Mo}^{\text{V}}\text{OCl}_4]$, based upon spectroscopic data, and the observation that metathesis with NaBPh_4 gave $[\text{Mo}^{\text{IV}}\text{OCl}(\text{diphosphine})_2][\text{BPh}_4]$.

During attempts to grow crystals of orange-yellow $[\text{MoOCl}_3\{\text{Me}_2\text{P}(\text{CH}_2)_2\text{PMe}_2\}]$, a few deep purple crystals were also isolated that were confirmed by an X-ray structure (Fig. 6) to be $[\text{MoCl}\{\text{Me}_2\text{P}(\text{CH}_2)_2\text{PMe}_2\}_2(\mu\text{-O})(\text{MoOCl}_4)]$, analogous to $[\text{MoCl}\{o\text{-C}_6\text{H}_4(\text{TeMe})_2\}_2(\mu\text{-O})(\text{MoOCl}_4)]$ described above. Both molybdenum centres are in a distorted octahedral geometry and linked by a very asymmetric oxide bridge, Mo1–O1 = 1.703(4), Mo2–O1 = 2.394(2) Å, which may be compared with Mo2–O2 = 1.657(8) Å for the terminal Mo=O unit. The original formulation^{7,8} was as ionic salts, $[\text{Mo}^{\text{IV}}\text{OCl}(\text{diphosphine})_2][\text{Mo}^{\text{V}}\text{OCl}_4]$. The reformulation as neutral μ -oxido dimers in the solid state is likely to apply to all the reported examples, with the long Mo–O bond easily cleaved to give ions in solution.

Red *fac*- $[\text{MoOCl}_3(\text{PMe}_3)_2]$ was obtained by Limberg *et al.*¹¹ as one product from reaction of the alkoxide complex, $[\text{Cl}_2\text{OMo}(\mu\text{-OEt})_2(\mu\text{-EtOH})\text{MoOCl}_2]$ with PMe_3 ; we obtained the same complex directly from $[\text{MoOCl}_3(\text{thf})_2]$ and PMe_3 in CH_2Cl_2 . Our X-ray structure and the spectroscopy (Experimental section) are in good agreement with published data,⁷ and are not discussed further here. The interest lies in the formation of a discrete *pseudo*-octahedral 1:2 Mo:PMe₃ monomer with the strong σ -donor alkyl phosphine, which contrasts with the formation of chloride-bridged dimers,

$[\{\text{MoOCl}_2(\text{EMe}_2)_2(\mu\text{-Cl})_2\}]$ (E = S, Se, Te), with the weaker donor chalcogenoethers discussed above.

Experimental

Syntheses were performed using standard Schlenk and glovebox techniques under a dry N_2 atmosphere. Solvents were dried by distillation from CaH_2 (CH_2Cl_2) or Na/benzophenone ketyl (toluene, n-hexane, diethyl ether). MoCl_5 and $\text{O}(\text{SiMe}_3)_2$ were obtained from Sigma-Aldrich. The monodentate ligands (SMe_2 , PMe_3 , SeMe_2) were obtained from Sigma-Aldrich or Strem and dried over molecular sieves. TeMe_2 was made by the method of Kuhn *et al.*³⁶ The dithioethers,³⁷ diselenoethers^{38,39} and ditelluroethers^{40,41} were prepared as described or by minor modifications thereof. MoOCl_3 was prepared from MoCl_5 and $\text{O}(\text{SiMe}_3)_2$.⁴² and MoOCl_4 obtained from Climax Molybdenum.

Infrared spectra were recorded on a PerkinElmer Spectrum 100 spectrometer in the range 4000–200 cm^{-1} , with samples prepared as Nujol mulls between CsI plates. UV/visible spectra were recorded on powdered solids using the diffuse reflectance attachment of a PerkinElmer 750S spectrometer. Magnetic measurements were made using a Johnson Matthey magnetic balance. Microanalyses on new compounds were undertaken by London Metropolitan University or Medac Ltd.

mer- $[\text{MoOCl}_3(\text{thf})_2]$

Prepared following the literature method.⁴³ Yield: 87%. IR spectrum (Nujol, ν/cm^{-1}): 982 s Mo=O, 1117 s, 833 s br thf, 342 s, 315 m Mo–Cl. UV/Vis spectrum (diffuse reflectance) ν/cm^{-1} : 32 550, 26 200, 22 000, 13 250. μ_{eff} : 1.71 B.M.



fac-[MoOCl₃{MeS(CH₂)₃SMe}]

Method 1. [MoOCl₃(thf)₂] (0.150 g, 0.41 mmol) was suspended in CH₂Cl₂ (3 mL) and a solution of MeS(CH₂)₃SMe (0.056 g, 0.41 mmol) in CH₂Cl₂ (2 mL) was added slowly and the solution left to stir for 1 h. The brown solution formed was then concentrated to 3 mL *in vacuo* and the green solid which precipitated, was filtered off and dried *in vacuo*. Yield: 0.070 g, 40%. Required for C₅H₁₂Cl₃MoOS₂ (354.58): C, 16.94; H, 3.41. Found: C, 17.02; H, 3.39%. IR spectrum (Nujol, ν/cm⁻¹): 955 s Mo=O, 348 s, 327 s, 306 m Mo-Cl. UV/Vis spectrum (diffuse reflectance) ν/cm⁻¹: 27 400, 26 000, 21 150, 18 350, 13 700. μ_{eff}: 1.71 B.M.

Method 2. MoOCl₄ (0.150 g, 0.59 mmol) was suspended in toluene (5 mL) and a solution of MeS(CH₂)₃SMe (0.081 g, 0.59 mmol) in toluene (2 mL) was slowly added and the green solution left to stir for 1 h. The green solution was concentrated to 3 mL *in vacuo* to produce a green precipitate that was filtered off and dried *in vacuo*. The green solid was washed in hexane (3 × 5 mL) and dried. Yield: 0.153 g, 70%. Required for C₅H₁₂Cl₃MoOS₂·0.2C₇H₈ (373.00): C, 20.61; H, 3.67. Found: C, 20.82; H, 3.77%. The complex was spectroscopically identical to that made by Method 1. Green crystals suitable for X-ray crystallography were grown from CH₂Cl₂.

mer-[MoOCl₃{ⁱPrS(CH₂)₂SⁱPr}]

[MoOCl₃{ⁱPrS(CH₂)₂SⁱPr}] was prepared similarly to Method 1 above, and isolated as a pale green solid. Yield: 62%. Required for C₈H₁₈Cl₃MoOS₂ (396.66): C, 24.22; H, 4.57. Found: C, 24.45; H, 4.15%. IR spectrum (Nujol, ν/cm⁻¹): 979 s Mo=O, 349 s, 312 m Mo-Cl. UV/Vis spectrum (diffuse reflectance) ν/cm⁻¹: 32 300, 30 400, 27 700, 23 000 sh, 21 500 sh, 13 600. μ_{eff}: 1.69 B.M.

mer-[MoOCl₃{PhS(CH₂)₂SPh}]

MoOCl₃ (0.150 g, 0.69 mmol) was suspended in CH₂Cl₂ (3 mL) and a solution of PhS(CH₂)₂SPh (0.170 g, 0.69 mmol) in CH₂Cl₂ (2 mL) was added slowly and the resulting green solution left to stir for 1 h. The resulting brown solution was concentrated to 3 mL *in vacuo* and filtered, and the orange-brown solid dried *in vacuo*. Yield: 0.244 g, 76%. Required for C₁₄H₁₄Cl₃MoOS₂ (464.69): C, 36.19; H, 3.04. Found: C, 35.97; H, 3.18%. IR spectrum (Nujol, ν/cm⁻¹): 966 s Mo=O, 354 s, 319 m Mo-Cl. UV/Vis spectrum (diffuse reflectance) ν/cm⁻¹: 32 200, 26 900, 22 600, 21 300, 18 500 sh, 13 000. μ_{eff}: 1.71 B.M.

[{MoOCl₂(SMe₂)₂}(μ-Cl)₂]

MoOCl₃ (0.150 g, 0.69 mmol) was suspended in CH₂Cl₂ (3 mL) and a solution of SMe₂ (0.085 g, 1.38 mmol) in CH₂Cl₂ (2 mL) was added slowly and the green solution left to stir for 1 h. The clear green solution was then concentrated to 3 mL *in vacuo* and layered with hexane (3 mL). The green crystals formed were isolated *via* filtration and dried *in vacuo*. Yield: 0.73 g, 38%. Required for C₄H₁₂Cl₆Mo₂O₂S₂ (560.86): C, 8.57; H, 2.16. Found: C, 8.98; H, 2.37%. IR spectrum (Nujol, ν/cm⁻¹): 1004 s Mo=O, 356 s, 319 s, 268 m Mo-Cl. UV/Vis spec-

trum (diffuse reflectance) ν/cm⁻¹: 26 000, 22 300, 19 600 sh, 13 800. μ_{eff}: 1.72 B.M./Mo.

[{MoOCl₂(SeMe₂)₂}(μ-Cl)₂]

MoOCl₃ (0.150 g, 0.69 mmol) was suspended in CH₂Cl₂ (3 mL) and a solution of SeMe₂ (0.150 g, 1.38 mmol) in CH₂Cl₂ (2 mL) was slowly added and the green solution left to stir for 1 h. The red solution formed was concentrated to 3 mL *in vacuo* and layered with hexane (3 mL). The dark brown crystals were isolated *via* filtration, and dried *in vacuo*. Yield: 0.154 g, 68%. Required for C₄H₁₂Cl₆Mo₂O₂Se₂ (654.65): C, 7.34; H, 1.85%. Found: C, 7.43; H, 1.93%. IR spectrum (Nujol, ν/cm⁻¹): 1004 s Mo=O, 368 sh, 351 s, 313 m Mo-Cl. UV/Vis spectrum (diffuse reflectance) ν/cm⁻¹: 32 500, 26 500, 20 700, 14 100. μ_{eff}: 1.68 B.M./Mo.

mer-[MoOCl₃{MeSe(CH₂)₂SeMe}]

[MoOCl₃(thf)₂] (0.150 g, 0.41 mmol) was suspended in CH₂Cl₂ (3 mL) and a solution of MeSe(CH₂)₂SeMe (0.089 g, 0.41 mmol) in CH₂Cl₂ (2 mL) was added slowly and the green solution left to stir for 1 h. The resulting brown solution was concentrated to 3 mL *in vacuo* and filtered, then the solid dried *in vacuo*, isolating a dark brown solid. Crystals grown from CH₂Cl₂ were dark green. Yield: 0.160 g, 90%. Required for C₄H₁₀Cl₃MoOSe₂ (434.34): C, 11.06; H, 2.32. Found: C, 11.60; H, 2.50%. IR spectrum (Nujol, ν/cm⁻¹): 960 s Mo=O, 342 s, 310 m Mo-Cl. UV/Vis spectrum (diffuse reflectance) ν/cm⁻¹: 25 800, 21 500, 19 300, 14 600. μ_{eff}: 1.71 B.M.

[MoOCl₃{MeSe(CH₂)₃SeMe}]

MoOCl₃ (0.150 g, 0.69 mmol) was suspended in CH₂Cl₂ (3 mL) and a solution of MeSe(CH₂)₃SeMe (0.158 g, 0.69 mmol) in CH₂Cl₂ (2 mL) was slowly added and the red/brown solution left to stir for 1 h. The brown solution was concentrated to 3 mL *in vacuo* and filtered and the dark brown solid isolated was dried *in vacuo*. A deep orange-brown crystalline solid was obtained from CH₂Cl₂ solution. Yield: 0.178 g, 58%. Required for C₅H₁₂Cl₃MoOSe₂·CH₂Cl₂ (533.30): C, 13.51; H, 2.65. Found: C, 13.96; H, 2.95%. IR spectrum (Nujol, ν/cm⁻¹): 954 s Mo=O, 346 s vbr, Mo-Cl. UV/Vis spectrum (diffuse reflectance) ν/cm⁻¹: 32 000 sh, 2700 br, 21 400, 19 500, 14 000. μ_{eff}: 1.70 B.M.

[(MoOCl₃)₂{*o*-C₆H₄(SeMe)₂}]_n

MoOCl₃ (0.150 g, 0.69 mmol) was suspended in dichloromethane (3 mL) and a solution of *o*-C₆H₄(SeMe)₂ (0.226 g, 0.69 mmol) in dichloromethane (2 mL) was added slowly and the dark red/brown solution left to stir for 1 h. The brown solution was then concentrated to 3 mL *in vacuo*, producing a brown precipitate, which was washed with OEt₂ (3 × 5 mL), then the brown-pink solid was dried *in vacuo*. Yield: 0.170 g, 53%. Required for C₈H₁₀Cl₆Mo₂O₂Se₂ (700.68): C, 13.71; H, 1.44. Found: C, 13.43; H, 1.53%. IR spectrum (Nujol, ν/cm⁻¹): 999 s br Mo=O, 351 w, 302 s, 292 sh, 256 m Mo-Cl. UV/Vis spectrum (diffuse reflectance) ν/cm⁻¹: 29 500 sh, 24 900, 20 900, 14 300. μ_{eff}: 1.69 B.M./Mo.



[{MoOCl₂(TeMe₂)₂(μ-Cl)₂}

MoOCl₃ (0.150 g, 0.69 mmol) was suspended in toluene (3 mL) and a solution of TeMe₂ (0.217 g, 1.38 mmol) in toluene (2 mL) was added slowly and the purple solution left to stir for 1 h. The deep purple solution was concentrated to 3 mL *in vacuo* and filtered, then the dark brown solid was dried *in vacuo*. Yield: 0.203 g, 78%. Required for C₄H₁₂Cl₆Mo₂O₂Te₂ (751.93): C, 6.39; H, 1.61. Found: C, 6.76; H, 2.06%. IR spectrum (Nujol, ν/cm⁻¹): 985 s br Mo=O, 327, 302 s br, 256 m Mo-Cl. UV/Vis spectrum (diffuse reflectance) ν/cm⁻¹: 34 500, 29 700 sh, 27 150, 26 100, 20 900, 19 500, 14 400. μ_{eff}: 1.68 B.M./Mo.

[(MoOCl₃)₂{MeTe(CH₂)₃TeMe}]_n

MoOCl₃ (0.150 g, 0.69 mmol) was suspended in dichloromethane (3 mL) and a solution of MeTe(CH₂)₃TeMe (0.217 g, 0.69 mmol) in dichloromethane (2 mL) was added slowly and the brown solution left to stir for 1 h. The brown solution was concentrated to 3 mL *in vacuo*, producing a brown precipitate which was washed with OEt₂ (3 × 5 mL), then the dark brown solid was dried *in vacuo*. Yield: 0.322 g, 61%. Required for C₅H₁₂Cl₆Mo₂O₂Te₂ (763.95): C, 7.86; H, 1.58. Found: C, 7.20; H, 1.38%. IR spectrum (Nujol, ν/cm⁻¹): 988 m Mo=O, 303 s, 292 m, 249 m Mo-Cl. UV/Vis spectrum (diffuse reflectance) ν/cm⁻¹: 26 500, 21 700, 18 600, ~13 000. μ_{eff}: 1.68 B.M./Mo.

[(MoOCl₃)₂{o-C₆H₄(TeMe)₂}]_n

MoOCl₃ (0.150 g, 0.69 mmol) was suspended in dichloromethane (3 mL) and a solution of o-C₆H₄(TeMe)₂ (0.249 g, 0.69 mmol) in dichloromethane (2 mL) was added slowly and the dark brown solution left to stir for 1 h. The brown solution was concentrated to 3 mL *in vacuo*, producing a brown precipitate which was washed with OEt₂ (3 × 5 mL), and dried *in vacuo*. Yield: 0.285 g, 52%. Required for C₈H₁₀Cl₆Mo₂O₂Te₂ (797.96): C, 12.04; H, 1.26. Found: C, 12.27; H, 1.43%. IR spectrum (Nujol, ν/cm⁻¹): 992 s br Mo=O, 343 m, 328 m, 302 s, 254 m Mo-Cl. UV/Vis spectrum (diffuse reflectance) ν/cm⁻¹: 32 500, 25 000 sh, 21 000, 19 200, 14 500. μ_{eff}: 1.70 B.M./Mo.

fac-[MoOCl₃(PMe₃)₂]

[MoOCl₃(thf)₂] (0.150 g, 0.41 mmol) was suspended in CH₂Cl₂ (3 mL) and a solution of PMe₃ (0.063 g, 0.82 mmol) in CH₂Cl₂ (3 mL) was added slowly and the dark green solution left to stir for 1 h. The red solution produced was then concentrated to 3 mL *in vacuo* and filtered, and the red solid dried *in vacuo*. Yield: 0.047 g, 31%. Required for C₆H₁₈Cl₃MoOP₂ (370.45): C, 19.45; H, 4.90. Found: C, 19.28; H, 4.74%. IR spectrum (Nujol, ν/cm⁻¹): 957 s Mo=O, 352 sh, 324 s, 305 m Mo-Cl. UV/Vis spectrum (diffuse reflectance) ν/cm⁻¹: 29 600, 26 500, 21 600, 20 500 sh, 15 500.

fac-[MoOCl₃{Me₂P(CH₂)₂PMe₂}]

[MoOCl₃(thf)₂] (0.150 g, 0.41 mmol) was suspended in CH₂Cl₂ (3 mL) and a solution of Me₂PCH₂CH₂PMe₂ (0.165 g, 0.41 mmol) in CH₂Cl₂ (3 mL) was added slowly and the solu-

tion left to stir for 1 h. The was concentrated to 3 mL *in vacuo*, filtered and then the solid was dried *in vacuo*. Yield: 0.131 g, 81%. Required for C₆H₁₆Cl₃MoOP₂ (368.44): C, 19.56; H, 4.38. Found: C, 19.83; H, 4.26%. IR spectrum (Nujol, ν/cm⁻¹): 951 s Mo=O, 362 m, 325 s, 306 s Mo-Cl. UV/Vis spectrum (diffuse reflectance) ν/cm⁻¹: 29 600, 26 500 sh, 21 600, 20 000, 15 500. μ_{eff}: 1.72 B.M.

X-ray experimental

Crystals were grown from slow evaporation of saturated solutions in CH₂Cl₂ or by liquid-liquid diffusion using CH₂Cl₂ and hexane. Data collections used a Rigaku AFC12 goniometer equipped with an enhanced sensitivity (HG) Saturn724+ detector mounted at the window of an FR-E+ SuperBright molybdenum (λ = 0.71073 Å) rotating anode generator with VHF Varimax optics (70 micron focus) with the crystal held at 100 K (N₂ cryostream). Crystallographic parameters are presented in Table S1.† Structure solution and refinement were performed using SHELX(T)-2018/2, SHELX-2018/3 through Olex2⁴⁴ and were mostly straightforward. H atoms were added and refined with a riding model. Where additional restraints were required, details are provided in the cif file for each structure found on CCDC.

Conclusions

A range of MoOCl₃ complexes with thio- and seleno-ethers have been prepared from [MoOCl₃(thf)₂] and the ligands in anhydrous CH₂Cl₂ solution. The more weakly coordinating PhS(CH₂)₂SPh, SME₂ and SeMe₂ fail to displace the thf, but complexes of these can be obtained using a suspension of MoOCl₃ in CH₂Cl₂. The reaction of MoOCl₄ with dithioethers results in reduction to Mo(v) as [MoOCl₃(dithioether)], behaviour which contrasts with that of WOCl₄ or WSCL₄, where either W(vi) or W(v) complexes can be obtained depending upon the reaction conditions.¹⁸ The stabilising effect of two oxido-groups on molybdenum(vi) is shown by the successful isolation of [MoO₂X₂(dithioether)] (X = Cl or Br).^{12,13} The limited affinity of the hard MoOCl₃ for the weaker donor monochalcogenoethers is reflected in the formation of 1:1 adducts, which achieve six-coordination by forming chloride bridges, as in [{MoOCl₂(E'Me₂)₂(μ-Cl)₂] (E' = S, Se), rather than by coordinating a second neutral donor ligand. The same explanation accounts for the formation of oligomeric complexes, [(MoOCl₃)₂(L-L)]_n with o-C₆H₄(SeMe)₂, o-C₆H₄(TeMe)₂ and MeTe(CH₂)₃TeMe, postulated to have a structure with only one chalcogen donor atom on each molybdenum, and where six-coordination is achieved *via* bridging chlorides and bridging dichalcogenoethers (Scheme 1). Although bridging behaviour might seem unexpected for chelates with o-C₆H₄-backbones, the presence of aryl groups makes these ligands weaker donors to hard metal centres – compare PhS(CH₂)₂SPh and ⁱPrS(CH₂)₂SⁱPr. There are several literature examples of o-phe-



nylene-based dichalcogenoethers adopting a bridging coordination mode.^{30–32} The behaviour contrasts with that of *o*-C₆H₄-based group 15 ligands, where *o*-C₆H₄(PMe₂)₂ or *o*-C₆H₄(AsMe₂)₂ can produce seven- or eight-coordination in tungsten(vi) complexes, such as [WOCl₄{*o*-C₆H₄(PMe₂)₂}] or [WF₄{*o*-C₆H₄(PMe₂)₂}]²⁺.^{45,46} The present work has also reported the first examples of Mo(v) telluroether complexes. Although the large soft tellurium centres are not usually thought to be good ligands for high valent d-block metals, a range of compounds has been reported in the last few years, including examples with NbCl₄,⁴⁷ NbCl₅⁴⁸ and TaCl₅,⁴⁸ although the complexes reported here are the first examples in Group 6. Also notable is the X-ray structural characterisation of the mixed valence [Mo^{IV}Cl{o-C₆H₄(TeMe)₂}₂(μ-O)Mo^{VO}Cl₄] and of the diphosphine analogue [Mo^{IV}Cl{Me₂P(CH₂)₂PMe₂}₂(μ-O)(Mo^{VO}Cl₄)] complexes of the latter type were reported in the 1970s^{7,8} but this is the first structural authentication.

The work has provided detailed characterisation of MoOCl₃-chalcogenoether complexes, and comparison with the W(vi) and W(v) analogues, and lays the basis for exploration of corresponding molybdenum sulfide chloride complexes,¹⁹ which may provide single source LPCVD reagents for deposition of MoS₂ thin films. The sulfide chloride systems will form the basis of future work.

Conflicts of interest

The authors have no conflicts to declare.

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