

Synthesis, structure and reactivity of Ni site models of [NiFeSe] hydrogenases†

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A series of structural models of the Ni centre in [NiFeSe] hydrogenases has been developed which exhibits key structural features of the Ni site in the H₂ cycling enzyme. Specifically, two complexes with a hydrogenase-analogous four-coordinate 'NiS₃Se' primary coordination sphere and complexes with a 'NiS₂Se₂' and a 'NiS₄' core are reported. The reactivity of the complexes towards oxygen and protons shows some relevance to the chemistry of [NiFeSe] hydrogenases. Exposure of a 'NiS₃Se' complex to atmospheric oxygen results in the oxidation of the selenolate group in the complex to a diselenide, which is released from the nickel site. Oxidation of the selenolate ligand on Ni occurs approximately four times faster than oxidation with the analogous sulfur complex. Reaction of the complexes with one equivalent of HBF₄ results in protonation of the monodentate chalcogenolate and the release of this ligand from the metal centre as a thiol or selenol. Unrelated to their biomimetic nature, the complexes serve also as molecular precursors to modify electrodes with Ni–S–Se containing particles by electrochemical deposition. The activated electrodes evolve H₂ in pH neutral water with an electrocatalytic onset potential of –0.6 V and a current density of 15 μA cm⁻² at –0.75 V vs. NHE.

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Introduction

The sustainable generation of the energy vector H₂ would provide a possible solution to a fossil-free economy,¹ and the improvement of H₂ evolution catalysts is therefore attracting much current attention.² In nature, the ubiquitous metals nickel and iron are employed in the active sites of [NiFe] and [FeFe] hydrogenases to reversibly catalyse the interconversion of protons and electrons to H₂ and operate at high rates at modest over-potentials.³ [NiFeSe] hydrogenases are a subclass of the [NiFe] hydrogenases with a selenocysteine (Sec) in place of a cysteine (Cys) residue terminally bound to the Ni centre (Fig. 1).⁴ These selenium containing enzymes have been shown to display a number of advantageous properties in comparison with the conventional [NiFe] hydrogenases which make them good candidates as H₂ evolution biocatalysts: they show higher H₂ evolution activities than [NiFe] hydrogenases⁵ and have fast reactivation times from O₂ damage at a low redox potential.^{5b,6} They were also previously employed in photocatalytic H₂

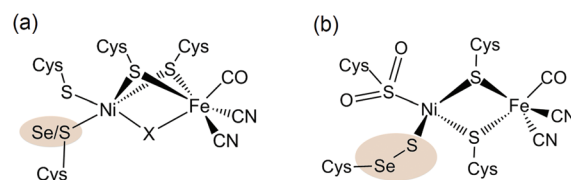


Fig. 1 (a) Molecular structure of the [NiFeSe] hydrogenase active site. The highlighted atom (Se/S) is S in [NiFe] hydrogenases and Se in [NiFeSe] hydrogenases. A hydride or oxo-group can serve as the bridging moiety X (the latter was only observed in oxidised [NiFe] hydrogenases) and an active reduced state does not contain a bridging ligand X. (b) Composition of a structural conformer in the oxidised [NiFeSe] hydrogenase from *Desulfovibrio vulgaris* Hildenborough.^{9f,g,10}

generation schemes^{5d,7} and even display catalytic H₂ production in the presence of O₂.^{5b,7} Therefore [NiFeSe] hydrogenases serve as an attractive blueprint and inspiration for the development of synthetic H₂ evolution catalysts.

The active site structure of the [NiFe(Se)] hydrogenases has been well studied using spectroscopic^{5a,6a,8} and crystallographic^{4,9} techniques (Fig. 1). It consists of a dinuclear [NiFe] centre bridged by two cysteine ligands. There is also a Cys and a Sec or Cys residue terminally bound to Ni.

The Ni ion serves as the active redox centre: its oxidation state varies between Ni(I) and Ni(III),^{8a,d-h} whereas Fe remains low spin Fe(II)^{8h} in all active site states. A catalytically active form of the enzyme has been shown to exist as Ni(III) with a bridging hydride between the Ni and Fe centres (position X in

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† Electronic supplementary information (ESI) available: Supporting table, figures and X-ray structure refinement details. CCDC 975187 (*n*-Bu₄N)[Ni(L^{Se})(Mes^S)], 939413 (*n*-Bu₄N)[Ni(L^S)(Mes^S)], 939414 (*n*-Bu₄N)[Ni(L^{Se})(Mes^{Se})], and 939415 (*n*-Bu₄N)[Ni(L^S)(Mes^{Se})]. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c3dt52967c



Fig. 1a) and a reduced Ni(II) form does not contain a bridging ligand X.^{8f,g,i} The Cys or Sec residue highlighted in Fig. 1a is proposed to act as a proton relay during H₂ cycling.^{4,9a-c} The different properties of Sec compared with Cys, such as its higher nucleophilicity and lower pK_a,¹¹ may cause the increased H₂ production activity of the [NiFeSe] subclass.⁵ We also note that there are other differences apart from the active site Sec, which could affect the chemistry of the enzyme. For example a medial [3Fe4S] cluster is present in *D. baculatum* [NiFeSe] hydrogenase rather than a [4Fe4S] cluster in conventional [NiFe] hydrogenases in the electron transfer chain between the active site and the protein surface.⁴

Despite a large number of structural mimics of the [NiFe] hydrogenase active site,¹² no systematic series of Ni subsite models with a stoichiometrically accurate 'NiS₃Se' core has been previously reported. Noteworthy approaches to mimic selenium in hydrogenases are complexes with a 'NiS₃SeP' centre such as penta-coordinate [Ni(SePh)(P(*o*-C₆H₄SH))₂-(*o*-C₆H₄SH)]⁻,^{12j,k} a number of Ni complexes with nitrogen and selenium-donor ligands,^{12m,n} and a series of diiron complexes containing mixed sulfur/selenium bridging ligands such as [Fe₂(μ-SC₃H₆Se-μ)(CO)₆].¹³

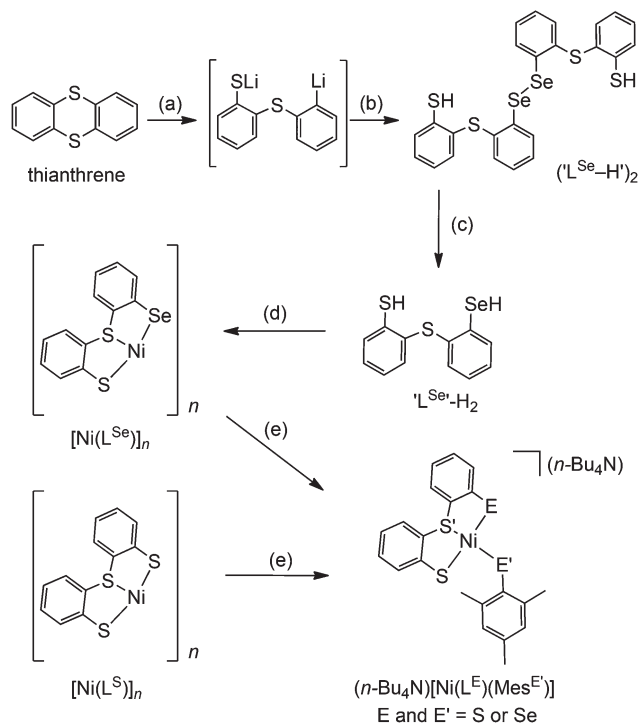
Herein we report on the synthesis, characterisation, reactivity towards O₂ and protons, and electrochemical study of four-coordinate Ni complexes with thiolate and selenolate ligands as structural models of the Ni site of reduced [NiFeSe] hydrogenases. The mononuclear model compounds display protonation and O₂ reactivity relevant to the hydrogenases. Electrochemical studies reveal that the Ni complexes do not operate as molecular H₂ production catalysts. However, they are useful precursors to catalytic Ni-containing particles for H₂ production in pH neutral H₂O.

Results and discussion

Synthetic pathway to hydrogenase inspired Ni complexes

The synthetic route to structural models of the Ni site in [NiFeSe] hydrogenase is outlined in Scheme 1. The novel tridentate mixed donor ligand 'L^{Se}-H₂' was synthesised from thianthrene. A catalytic amount of 4,4'-di-*tert*-butylbiphenyl was used with two equivalents of lithium powder in tetrahydrofuran at -90 °C to give a radical anion which is strong enough to cleave one of the thianthrene carbon-sulfur bonds.¹⁴ Introduction of elemental selenium into the lithiated product and acidic workup gave the diselenide ('L^{Se}-H')₂, which can be subsequently reduced with LiAlH₄ to give the selenol 'L^{Se}-H₂' after acidic workup. Refluxing a solution of 'L^{Se}-H₂' in tetrahydrofuran with one equivalent of Ni(OCOCH₃)₂·4H₂O in methanol yields the complex [Ni(L^{Se})_n] as an insoluble black precipitate in 87% yield.

[Ni(L^S)_n] (Scheme 1) was previously characterised by Sellmann and co-workers using single crystal X-ray analysis and it showed dimeric and trimeric forms, depending on crystal growth conditions (Fig. 2).^{12i,l} This composition is therefore also likely for [Ni(L^{Se})_n]. [Ni(L^S)_n] has been prepared following a previously



Scheme 1 Synthetic pathway to models of the Ni-site in [NiFe(Se)] hydrogenases: (a) Li, 4,4'-di-*tert*-butylbiphenyl, tetrahydrofuran, -90 °C. (b) (i) Se, -90 °C to -50 °C; (ii) H₃O⁺. (c) (i) NaBH₄; (ii) H₃O⁺. (d) Ni(OCOCH₃)₂·4H₂O, tetrahydrofuran-methanol, reflux. (e) NaMes^{E'} (E' = S or Se), *n*-Bu₄NOH, tetrahydrofuran-methanol.

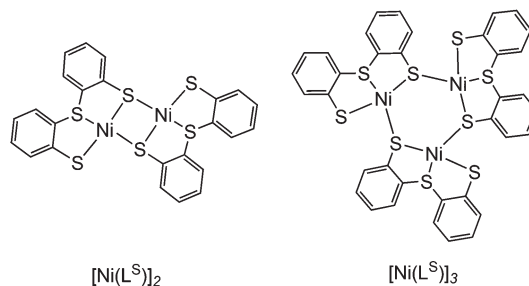


Fig. 2 Chemical structure of [Ni(L^S)_n] in the dimeric ($n = 2$)^{12l} and trimeric ($n = 3$)¹²ⁱ forms.

reported procedure.¹²ⁱ The oligomeric structure in [Ni(L^{Se})_n] and [Ni(L^S)_n] can be broken upon reaction with one equivalent of sodium mesityl selenolate or sodium mesityl thiolate in tetrahydrofuran-methanol (4:1) at room temperature. Addition of one equivalent of *n*-Bu₄NOH yields the corresponding four mononuclear Ni complexes (*n*-Bu₄N)[Ni(L^E)(Mes^{E'})] (E = S or Se; E' = S or Se, Scheme 1). Brown crystals of the Ni complexes were grown from a saturated solution in tetrahydrofuran-hexane giving the pure product in good yields (68 to 79%). The compounds have been characterised by ¹H NMR (Fig. S1-S8†), mass spectrometry (Fig. S9-S16†), ATR-IR spectroscopy (Fig. S17†) and elemental analysis. The tetraphenyl phosphonium salts of [Ni(L^S)(Mes^S)⁻] and [Ni(L^S-



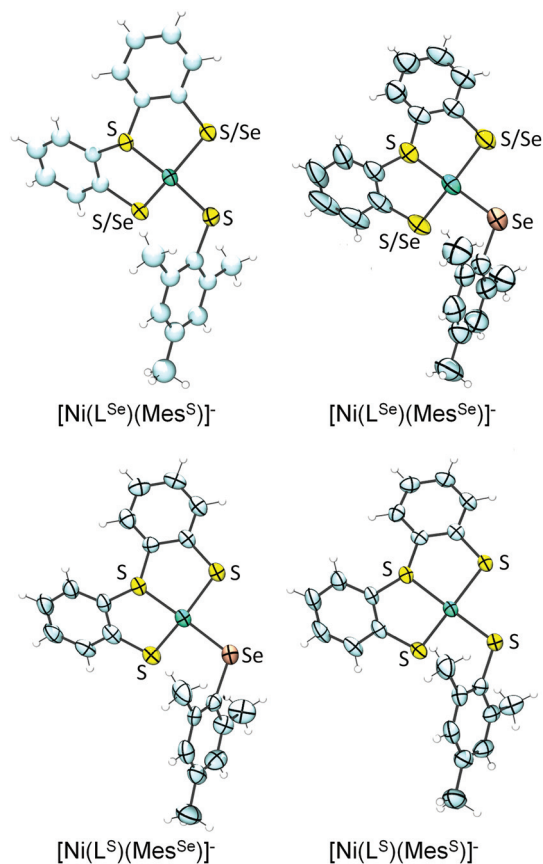


Fig. 3 X-ray single crystal structures of complexes (*n*-Bu₄N)[Ni(L^E)(Mes^{E'})] showing 50% probability ellipsoids (counteraction not shown; the asymmetric units of [Ni(L^{Se})(Mes^S)]⁻, [Ni(L^S)(Mes^{Se})]⁻ and [Ni(L^S)(Mes^S)]⁻ contain two independent anions; images created using Ortep 3 for Windows¹⁵). S/Se indicates disorder of S and Se over two positions.

(Mes^{Se})]⁻ have also been prepared and characterised by ¹H, ¹³C, ⁷⁷Se NMR spectroscopy, ESI-MS and elemental analysis.

Structural analyses of Ni complexes

The single crystal X-ray structures of all four complexes were obtained as tetrabutylammonium salts (Fig. 3; selected bond

lengths and angles are summarised in Table 1). The complex anions [Ni(L^{Se})(Mes^S)]⁻ and [Ni(L^S)(Mes^{Se})]⁻ contain a Ni²⁺ cation bound to three S and one Se donors and reflect the primary ligand sphere of the Ni site in [NiFeSe] hydrogenases.⁴

The complexes display a distorted square planar arrangement around the metal centre. The average *trans* S–Ni–SMes and S–Ni–SeMes bond angle is 160.2 ± 0.6°. The [NiFeSe] hydrogenase displays angles of 168.0° and 106.4° between Se–Ni–(μ-S) and S_{terminal}–Ni–(μ-S), respectively (PDB 1CC1).⁴ Crystallographic disorder of the thiolate and selenolate donors in 'L^{Se}', occurs in [Ni(L^{Se})(Mes^{Se})]⁻ (indicated as S/Se in Fig. 3) and [Ni(L^{Se})(Mes^S)]⁻ and we therefore only consider the bond lengths from the Ni to the Mes^S and Mes^{Se} ligand in our discussion. The average Ni–Se(Mes) bond length at 2.306 ± 0.003 Å is longer than the Ni–S(Mes) distance at 2.189 ± 0.003 Å. A Ni–Se bond length of 2.46 Å and average Ni–S distance of 2.2 Å (at 2.15 Å resolution) was reported for the reduced [NiFeSe] hydrogenase from *Desulfomicrobium baculatum*.⁴

Reactivity of the Ni complexes with atmospheric O₂

A distinct difference between the [NiFe] and [NiFeSe] hydrogenases is the reactivity of their active sites with O₂. EPR studies showed that upon reactivity with atmospheric O₂ the Ni in the conventional [NiFe] hydrogenases is oxidised to Ni(III), in one of two states known as the 'ready' (Ni-B) and 'unready' (Ni-A) states.^{8a,d,e,h} Crystal structures of the oxidised enzyme show an oxygen containing ligand bridging the Ni and Fe centre (position X, Fig. 1) in both oxidised states.^{9b,d,e} The oxidised active site in [NiFeSe] hydrogenase is quite different; the Ni centre is not oxidised and it remains diamagnetic Ni(II).^{5a} Single crystal X-ray structures of the oxidised [NiFeSe] hydrogenase in *D. vulgaris* and *D. baculatum* confirm that there is no bridging oxo ligand between the metal centres. In fact, selenium and/or sulfur are oxidised instead.^{9f-h} This different active site chemistry might explain the different rates for inactivation and re-activation between [NiFe] and [NiFeSe] hydrogenases.^{5b}

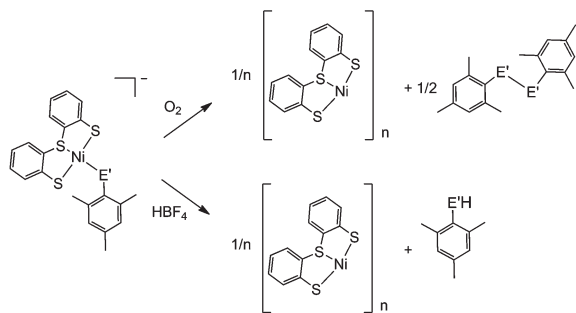
We therefore investigated the reactivity of the Ni complexes with O₂. As expected, all complexes are air sensitive and

Table 1 Selected bond lengths (Å) and angles (°) for complexes (*n*-Bu₄N)[Ni(L^E)(Mes^{E'})] (see Scheme 1 for labeling)

	<i>(n</i> -Bu ₄ N)[Ni(L ^{Se})(Mes ^S)] (E = Se, E' = S)		<i>(n</i> -Bu ₄ N)[Ni(L ^{Se})(Mes ^{Se})] (E = Se, E' = Se)		<i>(n</i> -Bu ₄ N)[Ni(L ^S)(Mes ^{Se})] (E = S, E' = Se)		<i>(n</i> -Bu ₄ N)[Ni(L ^S)(Mes ^S)] (E = S, E' = S)	
<i>Bond lengths</i>								
Ni–S	n.d. ^a	n.d. ^a	n.d. ^a	2.158(2)	2.155(2)	2.1630(12)	2.19642(14)	
Ni–S'	2.1257(17)	2.1263(18)	2.124(1)	2.124(2)	2.131(3)	2.1254(13)	2.1218(14)	
Ni–E	n.d. ^a	n.d. ^a	n.d. ^a	2.197(2)	2.190(2)	2.1987(12)	2.1960(14)	
Ni–E'	2.1930(18)	2.183(2)	2.3102(9)	2.3009(14)	2.3066(17)	2.1947(14)	2.1856(16)	
<i>Bond angles</i>								
S–Ni–E	153.2 ^a	157.0 ^a	155.2 ^a	153.63(9)	156.26(10)	152.78(5)	156.14(5)	
S–Ni–S'	89.6 ^a	90.5 ^a	90.0 ^a	92.12(9)	91.39(10)	91.76(5)	91.15(5)	
S–Ni–E'	90.9 ^a	89.7 ^a	90.6 ^a	96.51(8)	95.39(9)	97.07(5)	96.05(6)	
E–Ni–S'	92.2 ^a	91.3 ^a	91.8 ^a	89.47(9)	89.78(10)	89.24(5)	89.58(5)	
E–Ni–E'	96.9 ^a	95.8 ^a	95.7 ^a	91.36(7)	90.67(8)	91.64(5)	90.71(6)	
S'–Ni–E'	158.74(7)	161.21(8)	160.79(5)	158.64(7)	161.97(8)	158.64(5)	161.33(6)	

^a n.d. = not accurately determined due to crystallographic disorder.





Scheme 2 Reactivity of $[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{Se}})]^-$ ($\text{E}' = \text{Se}$) and $[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{S}})]^-$ ($\text{E}' = \text{S}$) with O_2 and HBF_4 .

convert completely to oxidised products over a maximum of five days. $(\text{PPh}_4)[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{Se}})]$ and $(\text{PPh}_4)[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{S}})]$ convert cleanly to $[\text{Ni}(\text{L}^{\text{S}})]_n$ and the dichalcogenide $(\text{Mes}^{\text{Se}})_2$ or $(\text{Mes}^{\text{S}})_2$ (Scheme 2). The oxidation products were isolated and characterised after exposing the complexes to atmospheric O_2 in dichloromethane. $[\text{Ni}(\text{L}^{\text{S}})]_n$ was isolated as a black precipitate and characterised by ATR-IR spectroscopy (Fig. S18†) and elemental analysis. The dichalcogenide was isolated from the supernatant and characterised by ^1H NMR and ESI-MS (Fig. S19–S21†).

The oxidation of the Se containing model emulates the reactivity of the enzyme. Oxidation of Se from the -2 to -1 oxidation state, and the formation of a Se–chalcogen bond concomitant with the loss of Se from the metal centre are all observed in the oxidised $[\text{NiFeSe}]$ hydrogenase from *Desulfovibrio vulgaris* Hildenborough.^{9f,g} Crystallographic analysis of this oxidised enzyme shows that the selenium is bound to a sulfur atom extrinsic to the active site. The source of the S atom is proposed to be H_2S in this sulfate reducing bacterium. In one oxidised conformer the selenium is not bound to the Ni centre and the extrinsic sulfur binds to Ni in its place (Fig. 1b).^{9f,g}

The oxidation of $(\text{PPh}_4)[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{Se}})]$ and $(\text{PPh}_4)[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{S}})]$ was also followed by ^1H NMR spectroscopy in deuterated dichloromethane (Fig. S22†). Over the course of oxidation the signals of the ‘ L^{S} ’ ligand at 6.8 to 7.5 ppm decrease as $[\text{Ni}(\text{L}^{\text{S}})]_n$ precipitates from solution and the methyl signals for the mesityl thiolate/selenolate ligand at 2.1 to 2.6 ppm decrease as new methyl signals appear around 2.3 ppm for the dichalcogenide oxidation product. The selenolate containing complex $(\text{PPh}_4)[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{Se}})]$ is oxidised more rapidly than the all sulfur complex: it is completely converted within 24 hours, compared with 96 h for $(\text{PPh}_4)[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{S}})]$. This observation supports the hypothesis that the fast reactivity of Se with O_2 prevents the nickel centre from being oxidised in $[\text{NiFeSe}]$ hydrogenases.^{9f-h}

Reactivity of the Ni complexes with HBF_4

Crystallographic evidence suggests that one of the terminal cysteine residues bound to the Ni centre in $[\text{NiFe}]$ hydrogenases (highlighted in Fig. 1a) acts as a proton relay during the catalytic cycle. This residue has increased crystallographic

disorder compared with the rest of the active site, suggesting various protonation states.^{9a-c} In the *D. baculatum* $[\text{NiFeSe}]$ hydrogenase the selenocysteine residue in the same position exhibits this crystallographic disorder suggesting that it is the proton relay in this case.⁴

Exposure of $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{Se}})]$ to one equivalent of HBF_4 in dichloromethane results in the protonation of the mesityl selenolate group (Scheme 2), in analogy to protonation in the enzyme active site. Mesityl selenol is released from the metal centre and has been isolated from the solution and characterised using ^1H NMR spectroscopy (Fig. S23†). The remaining ‘ $\text{Ni}(\text{L}^{\text{S}})$ ’ centre precipitates as the black solid $[\text{Ni}(\text{L}^{\text{S}})]_n$ (separated and characterised by ATR-IR and elemental analysis). Diethyl ether was added to the filtrate to precipitate $n\text{-Bu}_4\text{NBF}_4$, which was separated by filtration and characterised using ^1H NMR spectroscopy and mass spectrometry. The solvent was removed from the remaining filtrate to give mesityl selenol. A comparable reactivity pattern is observed with $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{S}})]$: the ‘ Mes^{S} ’ ligand is protonated and is released from the metal centre as mesityl thiol (Scheme 2; Fig. S24†). The products of protonation were separated and characterised in the same way as with $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{Se}})]$. The protonation of these complexes parallels the proposed protonation of the selenocysteine and cysteine residue in the hydrogenase.

Electrochemistry and electrodeposition of Ni complexes

The cyclic voltammograms (CVs) of all four complexes show an irreversible reduction process at approximately -1.4 V vs. NHE at a scan rate of 100 mV s^{-1} in acetonitrile, dichloromethane and dimethylformamide containing $n\text{-Bu}_4\text{NBF}_4$ as a supporting electrolyte (Fig. 4a black trace, Fig. S25–26†). This wave is assigned to the reduction of $\text{Ni}(\text{II})$ to $\text{Ni}(\text{I})$ based on previous electrochemical studies with Ni–thiolate complexes.^{12g,16} A catalytic wave grows at approximately -1.30 V vs. NHE with increasing amounts of triethylammonium chloride (Et_3NHCl) in acetonitrile indicating electrocatalytic H_2 production (Fig. 4a, Fig. S27†). The overpotential required for the reduction of protons is approximately 0.65 V under these conditions, calculated from the half wave potential for catalytic proton reduction in the presence of 10 equivalents of Et_3NHCl (-1.17 V vs. NHE, Fig. S28†) and the standard potential for reduction of protons from Et_3NHCl in acetonitrile (-0.51 V vs. NHE).¹⁷

However, the proton reduction activity does not stem from a molecular Ni species, but from a solid deposit formed on the electrode surface at the applied negative potential. Here, the Ni complexes act as molecular precursors to catalyst particles. The same electrocatalytic response was observed after removing the working electrode following electrocatalytic proton reduction in $n\text{-Bu}_4\text{NBF}_4$ –acetonitrile (0.1 M) with Et_3NHCl (10 mM) and a Ni complex, rinsing it with acetonitrile and immersing it in a fresh, Ni complex free electrolyte solution (Fig. S29†). Electronic absorption spectroscopy of the nickel complexes confirmed their stability in the presence of 10 equivalents of Et_3NHCl in acetonitrile in the absence of an



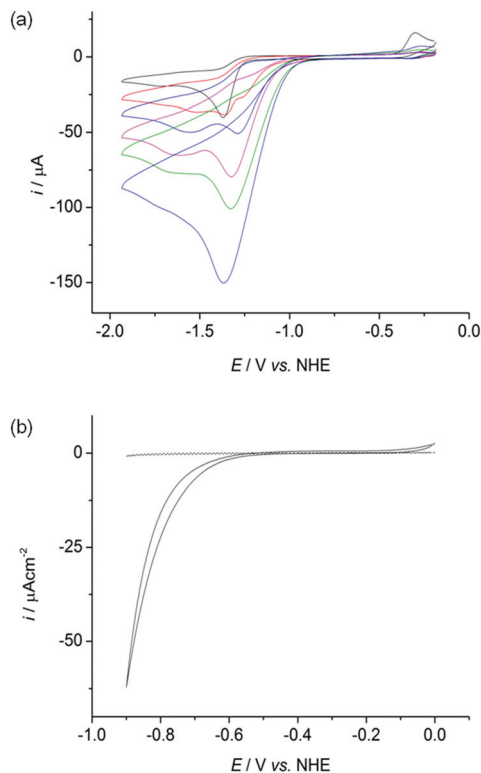


Fig. 4 (a) CV of $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{Se}})(\text{Mes}^{\text{S}})]$ (1 mM) in acetonitrile containing $n\text{-Bu}_4\text{NBF}_4$ (0.1 M) at a glassy carbon electrode with a scan rate of 100 mV s^{-1} with increasing concentrations of Et_3NHCl : no acid (black), 1 (red), 2 (blue), 3 (pink), 5 (green) and 10 (purple) mM. (b) CV of a Ni particle activated (solid trace) and unmodified (dashed trace) glassy carbon electrode in a phosphate buffered aqueous solution (0.1 M) at pH 7 at a scan rate of 100 mV s^{-1} at room temperature.

applied potential, demonstrating that the complexes were deposited electrochemically on the electrode at negative potentials (Fig. S30†).

The oxidation wave at $E = -0.25 \text{ V}$ in Fig. 4a and S25–S27† is attributed to the oxidation of a soluble product formed during the irreversible reduction of the complexes. The redox process cannot be assigned to the oxidation of deposited material as it is not visible in the CV of a nickel modified glassy carbon electrode in a fresh electrolyte solution.

Although the particle formation has no obvious relevance to the biomimetic features of the complexes, there is currently much interest in assembling heterogeneous electrocatalysts for proton reduction from molecular precursors.¹⁸ We therefore decided to investigate the Ni particle formation in more detail. Scanning electron microscopy (SEM) and energy-dispersive X-ray spectroscopy (EDX) analysis of a glassy carbon slide (total surface area 1.6 cm^2) modified with $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{Se}})(\text{Mes}^{\text{S}})]$ (1 mM) at -1.33 V vs. NHE for 0.5 h showed the deposition of nickel containing particles on the glassy carbon surface (Fig. 5a).

EDX of the particles revealed that they contain Ni, S and Se (Fig. 5b), but the exact composition and oxidation state of the Ni–S–Se containing deposit remains unknown. Using

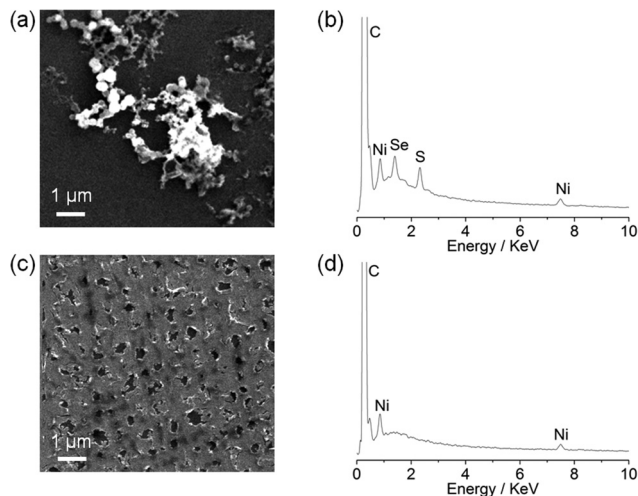


Fig. 5 SEM images and EDX spectra of a (a and b) Ni particle modified glassy carbon electrode prepared from the electro-deposition of $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{Se}})(\text{Mes}^{\text{S}})]$ (1 mM) and (c and d) Ni film on a glassy carbon electrode prepared from the electrodeposition of $\text{Ni}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ (1 mM). The deposits were formed with the compounds at an applied potential of -1.33 V vs. NHE for 0.5 h in acetonitrile (0.1 M $n\text{-Bu}_4\text{NBF}_4$) with Et_3NHCl (10 mM).

$\text{Ni}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ as a precursor resulted in a Ni film (no sulfur or selenium residues were detected) covering the electrode surface (Fig. 5c and 5d) rather than Ni particle formation as observed with the Ni complexes. Our observations demonstrate that the molecular Ni precursors allow for the incorporation of sulfur and selenium in the solid precipitate and the formation of a distinct morphology compared to using $\text{Ni}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$.

Electrocatalytic activity of Ni-activated electrodes in water

We subsequently investigated the activity of the Ni particle modified electrode, which was formed by holding the potential at -1.33 V vs. NHE for 0.5 h in a solution of the Ni complexes (1 mM) and Et_3NHCl (10 mM) in $n\text{-Bu}_4\text{NBF}_4$ -acetonitrile (0.1 M) and washing of the electrode. The Ni particle electrode displayed a respectable activity in a pH neutral Ni free aqueous phosphate solution (0.1 M; Fig. 4b). The onset of a catalytic wave was observed at approximately -0.60 V vs. NHE , which corresponds to a small overpotential of 180 mV ($E^0, \text{H}^+/\text{H}_2 = -0.42 \text{ V at pH 7}$).

The production of H_2 was confirmed using controlled potential electrolysis at -0.9 V vs. NHE with a glassy carbon rod or fluoride doped tin oxide (FTO) coated glass electrode (both approximately 1.6 cm^2) modified with the Ni precursor by the same procedure. SEM/EDX analysis also confirmed the deposition of Ni on the FTO substrate (Fig. S31†). A Faradaic yield of approximately 60 to 71% was observed for all four deposits on glassy carbon and FTO glass modified with the Ni. The total amount of H_2 generated varied widely on glassy carbon (10 to 2000 μmol after 20 h bulk electrolysis), but was more reproducible on FTO coated glass (9 to 55 μmol after 10 h electrolysis, Fig. S32 and Table S1†).



Modifying the FTO electrode with $\text{Ni}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ by the same procedure resulted in the formation of comparable amounts of H_2 , but with a reduced Faradaic yield of only 32%, suggesting that the molecular precursor does not only have an effect on the composition and morphology of Ni deposit, but also on the activity of the Ni-modified electrodes. We also compared the Ni particle electrodes with the benchmark proton reduction catalyst Pt.¹⁹ A metallic Pt foil with the same geometrical surface area (approximately 1.6 cm^2) was therefore tested as the working electrode under the same experimental conditions. The activity of the Pt foil is approximately an order of magnitude higher than the Ni particle modified FTO electrodes in an aqueous phosphate solution at pH 7 (Fig. S32†).

Conclusions

We have demonstrated the synthesis and characterisation of the first series of complexes, which resemble the primary coordination sphere of the Ni site in [NiFeSe] hydrogenases. Single crystal X-ray structures are reported for all complexes. Two complexes, $[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{Se}})]^-$ and $[\text{Ni}(\text{L}^{\text{Se}})(\text{Mes}^{\text{S}})]^-$, display the key structural features of the Se-containing enzyme such as a distorted four-coordinate 'NiS₃Se' environment.

Aerobic oxidation of the complex $[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{Se}})]^-$ results in the oxidation of the monodentate selenium ligand to form a dichalcogenide, whereas Ni remained in the +2 oxidation state. Comparable chemistry was observed in the active site of *D. vulgaris* [NiFeSe] hydrogenase after exposure to O_2 .^{9f,g} Reactivity of $[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{Se}})]^-$ with HBF_4 leads to protonation of the selenolate ligand, indicating that the selenium atom is indeed a plausible protonation site during H_2 cycling in [NiFeSe] hydrogenases. Although unrelated to the biomimetic composition of the Ni molecules, the complexes also act as precursors to Ni-containing particles on an electrode surface, which show high electroactivity in pH neutral aqueous solution. Work is currently in progress to introduce iron in our Ni site precursors and assemble a full structural [NiFeSe] hydrogenase model complex.

Experimental section

Materials and methods

Unless otherwise stated all compounds were prepared using an anhydrous and anaerobic MBraun glovebox or Schlenk techniques. All starting materials were purchased from commercial suppliers in the highest available purity for all analytical measurements and used without further purification. Organic solvents were dried and deoxygenated prior to use. Mesityl selenol²⁰ and $[\text{Ni}(\text{L}^{\text{S}})]_n$ ¹²ⁱ (Scheme 1) were prepared according to literature procedures.

Physical measurements

¹H and ¹³C NMR spectra were recorded on a Bruker DPX-400 MHz spectrometer and the spectra referenced against

the solvent peak. The ⁷⁷Se NMR spectrum was recorded on a Bruker Avance 500 MHz BroadBand NMR spectrometer and referenced against dimethyl selenide in *d*-benzene as an external reference at 0 ppm. The mass spectrum of $(\text{L}^{\text{Se}}\text{-H})_2$ was carried out on a Waters ZQ HPLC-MS. The mass spectra of $\text{L}^{\text{Se}}\text{-H}_2$ and dimesityl diselenide were recorded by the University of Cambridge Mass Spectrometry Service using a Bruker Bio Apex 4.0 FTICR EI MS. The mass spectra of the metal complexes and inorganic salts were recorded on a Waters Quattro LC electrospray ionisation mass spectrometer. Expected and experimental isotope distributions of $[\text{Ni}(\text{L}^{\text{E}})(\text{Mes}^{\text{E}})]^-$ were compared. Elemental analysis was carried out by the micro-analysis service of the Department of Chemistry, University of Cambridge. IR spectra were recorded on a Perkin Elmer SpectrumOne FTIR spectrometer with an ATR sampling accessory. Electronic absorption spectra were recorded on an Agilent Cary UV-Vis 50 Bio spectrometer. The SEM images were obtained using a Philips XL30 132-10 electron microscope. Energy-dispersive X-ray spectroscopy (edax PV7760/68 ME) was used at a 15 kV acceleration voltage, spot size 4.0 and an acquisition time of at least 100 s. The elements were assigned and atomic ratios were identified using the built in software (EDAX).

X-ray crystallographic studies

All data were recorded with Mo K_α radiation ($\lambda = 0.71073 \text{ \AA}$) on a Nonius Kappa CCD diffractometer fitted with an Oxford Cryosystems Cryostream cooling apparatus. The single crystals were mounted in Paratone N oil on the tip of a glass fibre and kept under a stream of N₂. Structure solution was carried out using direct methods and refined by least squares (*SHELXL-97*)²¹ using Chebyshev weights on F_o^2 . The weighted *R*-factor, *wR* and goodness of fit (GOF) are based on F^2 . The hydrogen atoms were assigned to idealised positions and given thermal parameters of 1.5 (methyl hydrogens) or 1.2 (non-methyl hydrogens) times the thermal parameter of the carbon atom to which they were attached. The tridentate L^{Se} in the complexes $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{Se}})(\text{Mes}^{\text{S}})]$ and $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{Se}})(\text{Mes}^{\text{Se}})]$ exhibits the same sort of disorder, in each case modelled as two poorly resolved S/Se sites. Crystal data, data collection parameters, and structure refinement details for the complexes are given in Table 2. Selected bond lengths and angles are shown in Table 1. The mean bond lengths and angles for the discussion in the paper were calculated as follows: for a sample of *n* observations x_i , a weighted mean value (x_u) with its standard deviation (σ) was calculated using the following equations: $x_u = \sum_i x_i/n$, $\sigma = \{\sum_i (x_i - x_u)^2/[n(n-1)]\}^{1/2}$.

Electrochemical measurements

CVs were recorded at room temperature under Ar using an IviumStat or CompactStat potentiostat. A standard three electrode cell was used for all measurements with a glassy carbon disc working (3 mm diameter), a platinum foil counter and a Ag/Ag⁺ (organic solutions) or a Ag/AgCl/KCl_(sat) (aqueous solutions) reference electrode. For CVs recorded in acetonitrile containing *n*-Bu₄NBF₄ (0.1 M, electrochemistry grade, Sigma



Table 2 Crystal data and structure refinement for complexes shown in Fig. 3 (for selected bond lengths and angles see Table 1)

Complex	(<i>n</i> -Bu ₄ N)[Ni(L ^{Se})(Mes ^S)]	(<i>n</i> -Bu ₄ N)[Ni(L ^{Se})(Mes ^{Se})]	(<i>n</i> -Bu ₄ N)[Ni(L ^S)(Mes ^{Se})]	(<i>n</i> -Bu ₄ N)[Ni(L ^S)(Mes ^S)]
Empirical formula	C ₃₇ H ₅₅ NNiS ₃ Se	C ₃₇ H ₅₅ NNiS ₂ Se ₂	C ₃₇ H ₅₅ NNiS ₃ Se	C ₃₇ H ₅₅ NNiS ₄
Formula weight	747.67	794.57	747.67	700.77
Temperature (K)	180(2)	220(2)	180(2)	180(2)
Space group	<i>P</i> 2 ₁ / <i>c</i>	<i>P</i> 2 ₁ / <i>n</i>	<i>P</i> 2 ₁ / <i>c</i>	<i>P</i> 2 ₁ / <i>c</i>
<i>a</i> (Å)	18.6006(4)	10.6327(2)	18.7224(2)	18.5275(2)
<i>b</i> (Å)	23.1712(5)	22.5152(5)	23.0398(2)	23.1544(2)
<i>c</i> (Å)	18.7401(5)	16.1237(5)	18.7818(2)	18.7038(2)
β (°)	109.268(1)	94.708(1)	109.810(1)	109.270(1)
<i>V</i> (Å ³)	7624.5(3)	3846.95(16)	7622.27(13)	7574.26(13)
<i>Z</i>	8	4	8	8
ρ_{calc} (g cm ⁻³)	1.303	1.372	1.303	1.229
μ (Mo K α , mm ⁻¹)	1.653	2.530	1.654	0.758
Crystal size (mm)	0.46 × 0.10 × 0.02	0.23 × 0.21 × 0.02	0.28 × 0.23 × 0.10	0.28 × 0.12 × 0.02
θ range (°)	3.51 to 25.34	3.55 to 25.66	3.51 to 22.49	3.52 to 27.50
Total number of data	43 959	35 314	52 166	73 103
Number of unique data	13 739	7112	9870	17 283
Number of parameters	393	304	602	602
Completeness	0.984	0.974	0.992	0.993
<i>R</i> ₁ ^a	0.0711	0.0452	0.0634	0.0757
<i>wR</i> ₂ ^b	0.1765	0.1123	0.1672	0.1819

$$^a R_1 = \sum ||F_o| - |F_c|| / \sum |F_o|. \quad ^b wR_2 = \{ \sum [w(F_o^2 - F_c^2)^2] / \sum [w(F_o^2)^2] \}^{1/2}.$$

Aldrich), the Fc/Fc⁺ couple was used as an external reference and potentials were converted to the normal hydrogen electrode (NHE) by adding 630 mV in acetonitrile.²² For CVs in a pH 7 aqueous phosphate solution (0.1 M), potentials were converted by adding 0.2 V to the potential against Ag/AgCl/KCl_(sat).²³

Controlled potential electrolysis (CPE)

CPE in phosphate solution (0.1 M, pH 7) was carried out using a fluoride doped tin oxide (FTO) coated glass electrode or glassy carbon rod electrode (geometrical surface area in contact with electrolyte solution approximately 1.6 cm²), a platinum mesh counter electrode and a Ag/AgCl reference electrode. CPE was carried out in an airtight electrochemical cell containing N₂ with 2% methane as the internal standard for gas chromatography, GC, analysis. The headspace gas was analysed using an Agilent 7890A GC equipped with a 5 Å molecular sieve column, using N₂ carrier gas with a flow rate of approximately 3 mL min⁻¹. The GC columns were kept at 40 °C and a thermal conductivity detector (TCD) was used. Measurements were taken every 2 h and the cell was purged to remove all hydrogen following each GC measurement. All measurements were repeated at least three times. Faradaic efficiency (%) = 100[H₂ (mol) × 2*F*/*Q* (C)].

Synthesis of 'L^{Se}-H₂

The first step of this reaction must be carried out under argon. Lithium (25% dispersion in oil, Sigma Aldrich) was washed with hexane (3 × 2 mL). To the de-oiled Li metal (0.080 g, 11.53 mmol) was added 4,4'-di-*tert*-butylbiphenyl (DTBB, 0.115 g, 432 μmol) in tetrahydrofuran (3 mL) and the reaction mixture was cooled to -90 °C (acetone-liquid N₂ bath), resulting in the formation of a bright blue solution of a radical anion of DTBB.¹⁴ Thianthrene (1.25 g, 5.76 mmol) in

tetrahydrofuran (15 mL) was added at -90 °C and the resulting beige solution was stirred for eight hours during which time the temperature of the reaction mixture was allowed to slowly reach -50 °C. Selenium powder (0.455 g, 5.76 mmol) was added in one batch and the orange solution was allowed to reach room temperature slowly overnight. Degassed water (50 mL) was slowly added and the aqueous layer was extracted with diethyl ether (4 × 50 mL) to remove DTBB and any remaining unreacted starting material. The aqueous layer was then acidified with aqueous HCl (2 M, 50 mL) and extracted with dichloromethane (3 × 25 mL). The combined organic layers were washed with aqueous HCl (2 M, 25 mL), and the solvent was removed under vacuum to give a yellow gum of crude ('L^{Se}-H')₂, which was used for the next step without further purification.²⁴ The crude ('L^{Se}-H')₂ was dissolved in tetrahydrofuran (15 mL) and added dropwise to a solution of LiAlH₄ (68 mg, 1.79 mmol) in tetrahydrofuran (3 mL) with stirring at room temperature. The resulting colourless solution was stirred overnight. Aqueous HCl (2 M, 12 mL) was then added dropwise and the product was extracted with diethyl ether (3 × 20 mL). The combined extracts were washed with aqueous HCl (2 M, 12 mL) and the solvent was removed under high vacuum at room temperature to give the product as a white solid, which was purified by recrystallisation from a saturated solution in tetrahydrofuran at -35 °C. Yield: 494 mg, 29%. ¹H NMR (400 MHz, CDCl₃) δ /ppm = 7.47 (1H, dd), 7.33 (1H, dd), 7.00-7.19 (6H, m), 4.05 (1H, s, SH), 1.91 (1H, s, SeH); EI-MS (CHCl₃) +ive: 297.94 (30%, L^{Se}); elemental analysis calculated (%) for C₁₂H₁₀S₂Se C 48.48, H 3.39; found C 48.67, H 3.39.

Synthesis of [NiL^{Se}]_{*n*}

A solution of 'L^{Se}-H₂ (200 mg, 671 μmol) in tetrahydrofuran (3 mL) was added dropwise to a solution of Ni(OCOCH₃)₂·4H₂O (167 mg, 671 μmol) in methanol (0.5 mL). The dark



brown suspension was heated to reflux for four hours, whereupon an insoluble black precipitate was separated by filtration, washed with tetrahydrofuran (3 × 2 mL) and dried under vacuum at room temperature. Yield 208 mg, 87%. Elemental analysis calculated (%) for $[C_{12}H_8NiS_2Se]_n$ C 40.72, H 2.28; found C 40.78, H 2.31. ATR-IR $\tilde{\nu}/cm^{-1}$ = 3040, 1568, 1441, 1424, 1239, 1154, 1085, 1038, 757, 746. $[NiL^{Se}]_n$ is insoluble in any common organic solvent.

General procedure for synthesis of $(n-Bu_4N)[Ni(L^E)(Mes^E)]$

To a solution of NaOMe in methanol was added mesityl thiol or selenol. The solution was stirred for 20 min and then added to a stirred suspension of $[NiL^E]_n$ in tetrahydrofuran at room temperature. The reaction mixture was stirred for one hour until $[NiL^E]_n$ fully dissolved giving a brown solution. A solution of $n-Bu_4NOH \cdot 30H_2O$ in methanol was added and the solution was stirred for an additional 20 min. All solvents were removed under high vacuum at room temperature and the brown residue was taken up in tetrahydrofuran and filtered through a Millex FG PTFE microfilter (pore size 20 μm). Hexane was layered on top of the tetrahydrofuran solution and the undisturbed mixture gave brown crystalline needles of the product after several days. The crystals were separated by filtration and washed with hexane. X-ray diffraction quality single crystals were selected directly from the reaction vessel.

$(n-Bu_4N)[Ni(L^{Se})(Mes^S)]$

NaOMe (28 mg, 525 μmol) in methanol (1 mL), mesityl thiol (79 μL , 525 μmol), $[Ni(L^{Se})]_n$ (186 mg, 525 μmol) in tetrahydrofuran (4 mL), $n-Bu_4NOH \cdot 30H_2O$ (420 mg, 525 μmol) in methanol (0.75 mL). Yield 280 mg, 71%. 1H NMR (400 MHz, THF- d_8) δ/ppm = 7.52 (1H, d, L^{Se}), 7.43 (1H, d, L^{Se}), 7.38 (1H, d, L^{Se}), 7.30 (1H, d, L^{Se}), 6.79–6.97 (4H, m, L^{Se}), 6.78 (2H, s, Mes^S), 3.34 (8H, m, $n-Bu_4N$), 2.79 (6H, s, Mes^S), 2.46 (3H, s, Mes^S), 1.59 (8H, m, $n-Bu_4N$), 1.56 (8H, m, $n-Bu_4N$), 1.37 (12H, t, $n-Bu_4N$); ESI-MS (CH_2Cl_2) –ive: 505 (100%, $[Ni(L^{Se})(Mes^S)]^-$), +ive: 242 (100%, $n-Bu_4N^+$); elemental analysis calculated (%) for $C_{37}H_{55}NiNS_3Se$ C 59.44, H 7.41, N 1.87; found C 59.17, H 7.36, N 1.90.

$(n-Bu_4N)[Ni(L^{Se})(Mes^{Se})]$

NaOMe (7.6 mg, 141 μmol) in methanol (2 mL), mesityl selenol (28.1 mg, 141 μmol), $[Ni(L^{Se})]_n$ (50 mg, 141 μmol) in tetrahydrofuran (3 mL), $n-Bu_4NOH \cdot 30H_2O$ (113 mg, 141 μmol) in methanol (1 mL). Yield 77 mg, 68%. 1H NMR (400 MHz, THF- d_8) δ/ppm = 7.59 (1H, d, L^{Se}), 7.52 (1H, d, L^{Se}), 7.26 (1H, d, L^{Se}), 7.17 (1H, d, L^{Se}), 6.78–6.88 (4H, m, L^{Se}), 6.66 (2H, s, Mes^{Se}), 3.41 (8H, m, $n-Bu_4N$), 2.71 (6H, s, Mes^{Se}), 2.11 (3H, s, Mes^{Se}), 1.66 (8H, m, $n-Bu_4N$), 1.35 (8H, m, $n-Bu_4N$), 0.90 (12H, t, $n-Bu_4N$); ESI-MS (CH_2Cl_2) –ive: 553 (100%, $[Ni(L^{Se})(Mes^{Se})]^-$), +ive: 242 (100%, $n-Bu_4N^+$); elemental analysis calculated (%) for $C_{37}H_{55}NiNS_2Se_2$ C 55.93, H 6.98, N 1.76; found C 55.85, H 6.94, N 1.81.

$(n-Bu_4N)[Ni(L^S)(Mes^{Se})]$

NaOMe (17.6 mg, 326 μmol) in methanol (2 mL), mesityl selenol (64.9 mg, 326 μmol), $[Ni(L^S)]_n$ (100 mg, 326 μmol) in

tetrahydrofuran (3 mL), $n-Bu_4NOH \cdot 30H_2O$ (261 mg, 326 μmol) in methanol (1 mL). Yield 175 mg, 72%. 1H NMR (400 MHz, THF- d_8) δ/ppm = 7.60 (2H, d, L^S), 7.09 (2H, d, L^S), 6.83 (2H, t, L^S), 6.72 (2H, t, L^S), 6.66 (2H, s, Mes^{Se}), 3.41 (8H, m, $n-Bu_4N$), 2.73 (6H, s, Mes^{Se}), 2.11 (3H, s, Mes^{Se}), 1.67 (8H, m, $n-Bu_4N$), 1.36 (8H, m, $n-Bu_4N$), 0.90 (12H, t, $n-Bu_4N$); ESI-MS (CH_2Cl_2) –ive: 505 (100%, $[Ni(L^S)(Mes^{Se})]^-$), +ive: 242 (100%, $n-Bu_4N^+$); elemental analysis calculated (%) for $C_{37}H_{55}NiNS_3Se$ C 59.44, H 7.41, N 1.87; found C 59.59, H 7.35, N 1.93.

$(n-Bu_4N)[Ni(L^S)(Mes^S)]$

NaOMe (65 mg, 1.21 mmol) in methanol (3 mL), mesityl thiol (182 μL , 1.21 mmol), $[Ni(L^S)]_n$ (400 mg, 1.30 mmol) in tetrahydrofuran (9 mL), $n-Bu_4NOH \cdot 30H_2O$ (966 mg, 1.21 mmol) in methanol (3 mL). Yield 672 mg, 79%. 1H NMR (400 MHz, THF- d_8) δ/ppm = 7.62 (2H, d, L^S), 7.12 (2H, d, L^S), 6.92 (2H, t, L^S), 6.79 (2H, t, L^S), 6.71 (2H, s, Mes^S), 3.49 (8H, m, $n-Bu_4N$), 2.77 (6H, s, Mes^S), 2.17 (3H, s, Mes^S), 1.75 (8H, m, $n-Bu_4N$), 1.43 (8H, m, $n-Bu_4N$), 0.98 (12H, m, $n-Bu_4N$); ESI-MS (CH_2Cl_2) –ive: 457 (100%, $[Ni(L^S)(Mes^S)]^-$), +ive: 242 (100%, $n-Bu_4N^+$); elemental analysis calculated (%) for $C_{37}H_{55}NiNS_4$ C 63.41, H 7.91, N 2.00; found C 63.46, H 7.92, N 2.08.

Synthesis of $(PPh_4)[Ni(L^S)(Mes^{Se})]$

Elemental Se (24 mg, 300 μmol) was added to mesityl magnesium bromide (300 μL of a 1 M solution in diethyl ether). The solution was stirred overnight until the Se powder dissolved and the colour changed from orange to yellow. Tetrahydrofuran (2 mL) was added followed by $[Ni(L^S)]_n$ (96 mg, 300 μmol) and this was stirred for one hour until $[Ni(L^S)]_n$ dissolved giving a brown solution. A solution of PPh_4Br (126 mg, 300 μmol) in methanol (0.5 mL) was added and the solution was stirred for 20 min. The solution was concentrated to dryness under reduced pressure. The brown residue was dissolved in tetrahydrofuran (3 mL) and filtered through a Millex FG PTFE microfilter (pore size 20 μm). Diethyl ether was added to precipitate the brown product which was separated by filtration, washed with diethyl ether and dried under vacuum. Yield 169 mg, 67%. 1H NMR (400 MHz, $CDCl_3$) δ/ppm = 7.66 (4 H, m, PPh_4), 7.52 (8H, m, PPh_4), 7.49 (8H, m, PPh_4), 7.42 (2H, d, L^S), 7.04 (2H, d, L^S), 6.79 (2H, t, L^S), 6.70 (4H, m, L^S and Mes^{Se}), 2.52 (6H, s, Mes^{Se}), 2.08 (3H, s, Mes^{Se}); ^{13}C NMR (500 MHz, $CDCl_3$) δ/ppm = 156.6, 143.8, 134.4 (PPh_4), 133.0, 130.8 (PPh_4), 129.8, 128.3, 127.0, 126.9, 120.2, 117.8 (PPh_4), 117.1 (PPh_4), 27.2 (Mes^{Se}), 21.1 (Mes^{Se}); ^{77}Se NMR (500 MHz, CD_2Cl_2) δ/ppm = 605.4; ESI-MS (CD_2Cl_2): –ive: 505 (100%, $[Ni(L^S)(Mes^{Se})]^-$), +ive: 339 (100%, PPh_4^+); elemental analysis calculated (%) for $C_{45}H_{39}NiP_3Se$ C 63.99, H 4.65, P 3.67; found C 63.60, H 4.50, P 3.60.

Synthesis of $(PPh_4)[Ni(L^S)(Mes^S)]$

To a solution of NaOMe (17 mg, 309 μmol) in methanol (1 mL) was added mesityl thiol (50 μL , 309 μmol). The solution was stirred for 20 min and then added to a stirred suspension of $[NiL^S]_n$ (100 mg, 309 μmol) in tetrahydrofuran (3 mL) at room temperature and the solution was stirred for one hour until



$[\text{NiL}^{\text{S}}]_n$ fully dissolved giving a brown solution. A solution of PPh_4Br in methanol was added and this was stirred for 20 min. The solution was concentrated to dryness under reduced pressure. The brown residue was dissolved in tetrahydrofuran (3 mL) and filtered through a Millex FG PTFE microfilter (pore size 20 μm). Diethyl ether was added to precipitate the brown product which was separated by filtration, washed with diethyl ether and dried under vacuum. Yield 158 mg, 64%. ^1H NMR (400 MHz, CD_2Cl_2) δ/ppm = 7.92 (4H, m, PPh_4), 7.78 (8H, m, PPh_4), 7.62 (8H, m, PPh_4), 7.49 (2H, d, L^{S}), 7.14 (2H, d, L^{S}), 6.89 (2H, t, L^{S}), 6.80 (2H, m, L^{S} and Mes^{S}), 2.59 (6H, s, Mes^{S}), 2.17 (3H, s, Mes^{S}); ^{13}C NMR (500 MHz, CD_2Cl_2) δ/ppm = 155.4, 142.9, 140.3, 135.7, 134.4 (PPh_4), 134.1, 130.7, 130.6 (PPh_4), 129.2, 127.3, 127.3, 120.4, 117.8 (PPh_4), 117.1 (PPh_4), 24.2 (Mes^{S}), 20.8 (Mes^{S}); ESI-MS (CH_2Cl_2); -ive: 457 (100%, $[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{S}})]^-$), +ive: 339 (100%, PPh_4^+); elemental analysis calculated (%) for $\text{C}_{45}\text{H}_{39}\text{NiPS}_4$ C 67.75, H 4.93, P 3.88; found C 67.40, H 4.95, P 3.91.

Reaction of $(\text{PPh}_4)[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{Se}})]$ with oxygen

$(\text{PPh}_4)[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{Se}})]$ (116 mg, 137 μmol) in dichloromethane (5 mL) was exposed to atmospheric O_2 for one day with stirring to give a brown solution containing dimesityl diselenide with a black precipitate characterised as $[\text{Ni}(\text{L}^{\text{S}})]_n$. *Isolation and characterisation of precipitate containing $[\text{Ni}(\text{L}^{\text{S}})]_n$* : The solid was separated by filtration and washed with dichloromethane, methanol and diethyl ether and dried under vacuum. ATR-IR spectroscopy, elemental analysis and single crystal X-ray analysis identified the solid product as $[\text{Ni}(\text{L}^{\text{S}})]_n^{121}$ (22 mg, 53%). ATR-IR $\tilde{\nu}/\text{cm}^{-1}$ = 3036, 1442, 1425, 1251, 1092, 759, 727. *Isolation and characterisation of filtrate containing $(\text{Mes}^{\text{Se}})_2$* : Diethyl ether (15 mL) was added to the filtrate solution to precipitate a tetraphenyl phosphonium salt as a pink solid which was separated by filtration and washed with diethyl ether (42 mg). ^1H NMR (400 MHz, CD_2Cl_2) δ/ppm = 7.66 (4 H, m, PPh_4), 7.52 (8H, m, PPh_4), 7.49 (8H, m, PPh_4), ESI-MS (CD_2Cl_2): +ive: 339 (100%); $\lambda_{\text{max}}/\text{nm}$ (acetonitrile) 226 nm. The solvent was removed from the resulting filtrate to give crude dimesityl diselenide as a yellow solid, which was purified by filtration through silica in diethyl ether (10 mg, 36%). ^1H NMR (400 MHz, CDCl_3) δ/ppm = 6.77 (2H, s), 2.18 (3H, s), 2.14, (6H, s), EI-MS 398.0; $\lambda_{\text{max}}/\text{nm}$ (acetonitrile) 289.

Reaction of $(\text{PPh}_4)[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{S}})]$ with oxygen

$(\text{PPh}_4)[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{S}})]$ (30.7 mg, 38.4 μmol) in dichloromethane (5 mL) was exposed to atmospheric O_2 for five days under stirring to give a brown solution containing dimesityl disulfide with a black precipitate, characterised as $[\text{Ni}(\text{L}^{\text{S}})]_n$. *Isolation and characterisation of precipitate containing $[\text{Ni}(\text{L}^{\text{S}})]_n$* : The solid was separated by filtration and washed with dichloromethane, methanol and diethyl ether and dried under vacuum. ATR-IR spectroscopy and elemental analysis identified the solid product as $[\text{Ni}(\text{L}^{\text{S}})]_n^{121}$ (9.6 mg, 86%). *Isolation and characterisation of filtrate containing $(\text{Mes}^{\text{S}})_2$* : Diethyl ether (15 mL) was added to the filtrate to precipitate a tetraphenyl phosphonium salt as a pink solid which was separated by filtration, washed

with diethyl ether and dried under vacuum (11 mg). The solvent was removed from the resulting filtrate to give crude dimesityl disulfide as a pale yellow solid, which was purified by filtration through silica in diethyl ether (5.8 mg, 80%). ^1H NMR (400 MHz, CDCl_3) δ/ppm = 6.82 (2H, s), 2.18 (3H, s), 2.14 (6H, s).

Reaction of $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{E}})(\text{Mes}^{\text{E}})]$ with oxygen

Reaction of $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{E}})(\text{Mes}^{\text{E}})]$ with a $n\text{-Bu}_4\text{N}$ cation gives the dichalcogenides $(\text{Mes}^{\text{E}})_2$ along with a mixture of insoluble products which could not be unequivocally isolated and characterised.

Reaction of $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{Se}})]$ with HBF_4

To a solution of $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{Se}})]$ (20 mg, 26.7 μmol) in dichloromethane (3 mL) was added $\text{HBF}_4\cdot\text{Et}_2\text{O}$ (3.7 μL , 26.7 μmol) in dichloromethane (2 mL) and the reaction mixture was stirred for 30 min, whereupon $[\text{Ni}(\text{L}^{\text{S}})]_n$ (5.5 mg, 64%) formed as a black precipitate. The solid was separated by filtration and washed with dichloromethane, methanol and diethyl ether, dried under vacuum and characterised by ATR-IR and elemental analysis. Diethyl ether was added to the remaining filtrate to precipitate crude $n\text{-Bu}_4\text{NBF}_4$, which was separated by filtration, washed with diethyl ether and dried under vacuum. ^1H NMR (400 MHz, CDCl_3) δ/ppm = 3.20 (8H, m), 1.62 (8H, m), 1.44 (8H, m), 1.00 (12 H, t), ESI MS (CHCl_3); -ive: 87 (100%, BF_4^-) +ive: 242 (100%, $n\text{-Bu}_4\text{N}^+$). Diethyl ether was removed from the remaining filtrate under vacuum to yield mesityl selenol (4.3 mg, 79%). ^1H NMR (400 MHz, CDCl_3) δ/ppm = 6.88 (2H, s), 2.34 (6H, s), 2.23 (3H, s), 1.24 (1H, s).

Reaction of $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{S}})]$ with HBF_4

To a solution of $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{S}})]$ (50 mg, 71.3 μmol) in dichloromethane (4 mL) was added $\text{HBF}_4\cdot\text{Et}_2\text{O}$ (9.7 μL , 71.3 μmol) in dichloromethane (2 mL) and this was stirred for 30 min. $[\text{Ni}(\text{L}^{\text{S}})]_n$ (20.9 mg, 83%) and mesityl thiol (8 mg, 74%) formed in analogy to the reaction of $(n\text{-Bu}_4\text{N})[\text{Ni}(\text{L}^{\text{S}})(\text{Mes}^{\text{Se}})]$ with HBF_4 and were characterised by the same methods. ^1H NMR of mesityl thiol (400 MHz, CDCl_3) δ/ppm = 6.81 (2H, s), 3.04 (1H, s), 2.27 (6H, s), 2.17 (3H, s).

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