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Electrocatalytic proton reduction by a model  
for [NiFeSe] hydrogenases†Gamze Gezer,<sup>a</sup> Dinesh Durán Jiménez,<sup>a</sup> Maxime A. Siegler<sup>b</sup> and  
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Two new heterodinuclear nickel–iron complexes [Ni(pbSmSe)FeCpCO]PF<sub>6</sub> and [Ni(xbSmSe)FeCpCO]PF<sub>6</sub> were synthesized as mimics of the [NiFeSe] hydrogenase active site (HCp = cyclopentadiene; H<sub>2</sub>pbSmSe = 1,9-diselenol-3,7-dithia-2,2,8,8-tetramethylnonane; H<sub>2</sub>xbSmSe = 1,2-bis(2-thiabutyl-3,3-dimethyl-4-selenol)benzene). The compounds were characterized by single crystal X-ray diffraction and cyclic voltammetry. X-ray structure determinations showed that in both NiFe complexes the nickel(II) center is in a square-planar S<sub>2</sub>Se<sub>2</sub> environment; the two selenolate donors are bridging to the iron(II) center that is further coordinated to an η<sup>5</sup>-cyclopentadienyl group and a carbon monoxide ligand. Electrochemical studies showed that the complex [Ni(pbSmSe)FeCpCO]PF<sub>6</sub> is an electrocatalyst for the production of H<sub>2</sub> in DMF in the presence of acetic acid at −2.1 V vs. Fc<sup>+</sup>/Fc; a foot-of-the-wave (FOWA) analysis of the catalytic currents yielded an estimation of *k*<sub>obs</sub> of 24 s<sup>−1</sup>.

## Introduction

Hydrogenase enzymes, catalyzing the reversible oxidation of dihydrogen, play an important role in the metabolism of bacteria.<sup>1</sup> In the past decades, hydrogenases have attracted the attention of synthetic chemists, since dihydrogen gas may be used as a sustainable source of energy. In order to produce dihydrogen gas for the application in fuel cells, new efficient electrocatalysts for the hydrogen-evolving reaction (HER) may be developed by using biomimetic, functional models of hydrogenases.<sup>2</sup>

Three types of hydrogenases are known, which are classified based on the metal center in the active site. The [FeFe] hydrogenases contain a dinuclear iron center linked to an Fe<sub>4</sub>S<sub>4</sub> cluster, comprising CO, CN<sup>−</sup> and a dithiolate ligand. These [FeFe] hydrogenases catalyze both H<sub>2</sub> evolution and uptake, but their predominant activity is in H<sub>2</sub> evolution. However, the [FeFe] hydrogenases generally are highly air sensitive. The [Fe] hydrogenases have a mononuclear iron catalytic center and do not contain Fe–S clusters. These enzymes catalyze the transfer of hydride groups and H<sub>2</sub> activation.<sup>3</sup> The third class of hydrogenases comprises the [NiFe] hydrogenases containing a heterodimetallic Ni–Fe active site with a nickel

center bound to four cysteine thiolates with two of the cysteines bridging between the nickel and an iron center (Fig. 1a).<sup>4</sup> Although this enzyme is mostly involved in the uptake of H<sub>2</sub>, it is also able to catalyze the production of H<sub>2</sub>.<sup>3</sup> The [NiFeSe] hydrogenases form a subclass of the [NiFe] hydrogenases, in which one of the non-bridging cysteines (Cys) in the active site of the enzyme is replaced by selenocysteine (Sec) (Fig. 1b).<sup>5</sup> [NiFeSe] hydrogenases show interesting properties for H<sub>2</sub> production such as their high catalytic rates and their activity at low overpotentials; as they are less air-sensitive they produce H<sub>2</sub> even in the presence of low concentrations of O<sub>2</sub>.<sup>6</sup> Compared to their cysteine homologues the [NiFeSe] hydrogenases have higher catalytic activity in the hydrogen evolution reaction.<sup>3,7</sup> This difference in activity may be explained by the differences in the physical properties of selenium compared to those of sulfur, such as its higher acidity and higher nucleophilicity, in addition to the lower redox potential of the selenocysteine redox couple. The p*K*<sub>a</sub> of Sec is 5.3 whereas the p*K*<sub>a</sub> of Cys is 8.3, which may help in the rapid exchange of protons.

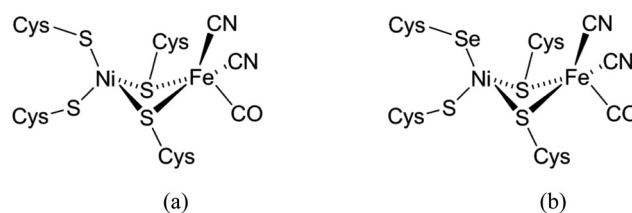


Fig. 1 Schematic representation of the active site in (a) [NiFe] and (b) [NiFeSe] hydrogenases.<sup>4,5</sup>

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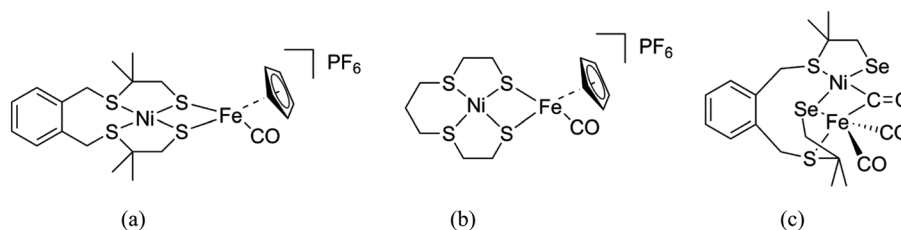


Fig. 2 Schematic drawings of the [NiFe] hydrogenase models reported by Artero (a)<sup>11</sup> and Schröder (b),<sup>20</sup> and the first structural model of [NiFeSe] hydrogenases reported by Reisner (c).<sup>7</sup>

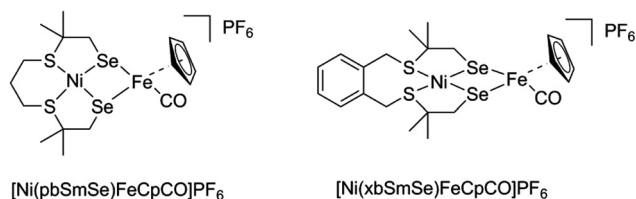


Fig. 3 Schematic drawing of the heterodimetallic compounds described in this paper.

Selenium is also a softer donor atom than sulfur; the polarizable volume of selenium is 3.8 Å in comparison to 2.9 Å for sulfur. Although these different properties can be the potential causes for the higher catalytic activity of the [NiFeSe] hydrogenases, the exact role of selenocysteine in the [NiFeSe] hydrogenases is still not completely clear.<sup>8,9</sup>

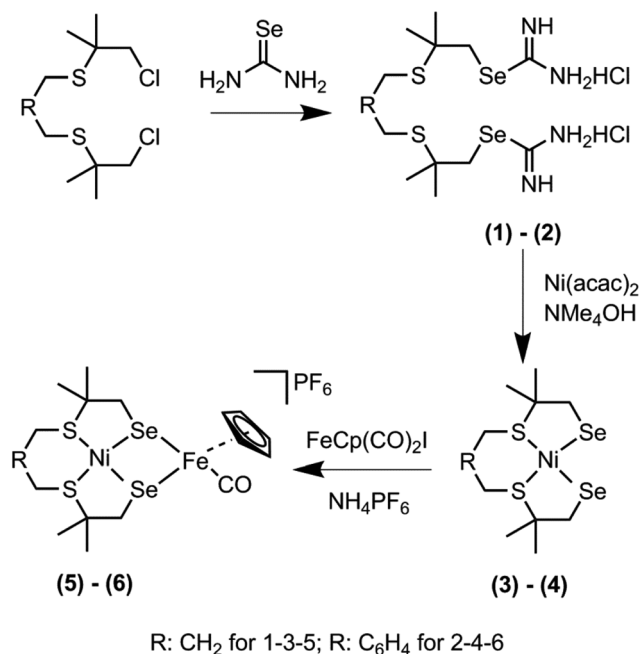
In the past decades a large number of structural and functional models for the active site in [NiFe] hydrogenases have been reported with overpotentials as low as 50 mV. From these studies, it was found that the addition of a cyclopentadienyl (Cp<sup>-</sup>) ligand resulted in increased catalytic rates and stability of the catalyst.<sup>10–14</sup> Apart from these models, a number of mononuclear Ni/Co/Fe complexes and several heterodimetallic [NiRu] complexes have been reported as functional models of the [NiFe] hydrogenases active site.<sup>15–19</sup> Two heterodinuclear compounds related to our work described in this paper have been reported by Artero and Schröder, comprising Ni<sub>2</sub>S<sub>2</sub> centers bound to a {FeCpCO} group (Fig. 2a and b).<sup>11,20</sup> However, so far only one heterodimetallic nickel–iron complex was reported comprising a selenolate ligand coordinated to the nickel center, as a potential model of the active site in [NiFeSe] hydrogenases (Fig. 2c).<sup>7</sup> In this paper, we describe the synthesis and characterization of two new heterodimetallic nickel–iron complexes (Fig. 3) as mimics of the [NiFeSe] hydrogenase active site. The electrochemical properties and electrocatalytic activity for H<sub>2</sub> production of these NiFe complexes are reported.

## Results and discussion

### Synthesis and characterization

The novel heterodinuclear compounds [Ni(pbSmSe)FeCpCO]PF<sub>6</sub> and [Ni(xbSmSe)FeCpCO]PF<sub>6</sub> were synthesized following

the procedure shown in Scheme 1. The precursors to two different selenouronium ligands were synthesized based on reported procedures. The compounds 1,9-dichloro-3,7-dithia-2,2,8,8-tetramethylnonane and bis(3-chloro-2,2-methyl-1-thiapropryl)-*o*-xylene were treated with two equivalents of selenourea in ethanol to give the selenouronium ligand precursors (1) and (2) as white powders in good purities in 86% and 90% yield, respectively. The ligand precursor (2) has been reported earlier.<sup>7</sup> The compounds [Ni(pbSmSe)] (3) and [Ni(xbSmSe)] (4) were synthesized by heating the selenouronium ligand precursors in refluxing ethanol with [Ni(acac)<sub>2</sub>] in the presence of NMe<sub>4</sub>OH. The compounds [Ni(pbSmSe)] (3) and [Ni(xbSmSe)] (4) were obtained as green solids in 64% and 83% yield, respectively. The compounds were characterized with <sup>1</sup>H and <sup>13</sup>C NMR spectroscopy, mass spectrometry, elemental analysis and single crystal X-ray diffraction for compound (3); the X-ray structure of compound (4) has been reported.<sup>7</sup> The nickel compounds (3) and (4) give rise to sharp resonances in the <sup>1</sup>H



Scheme 1 Synthesis scheme of the selenouronium salts (1) and (2), the mononuclear Ni complexes (3) and (4) and the heterodinuclear NiFe complexes (5) and (6).



NMR spectra indicating that the nickel(II) centers in these compounds are in low-spin, square-planar geometries, which are retained in solution. The clear NMR spectra are in contrast with those of the corresponding sulfur-based compounds  $[\text{Ni}(\text{pbSmS})]^{21}$  and  $[\text{Ni}(\text{xbSmS})]^{22}$  which generally show broad signals. Such broadening of the NMR signals is ascribed to fluxional behavior of the ligand surrounding the nickel ion; as a result part of the time the nickel centers are in more or less tetrahedral geometries giving rise to the paramagnetic high-spin state. Reaction of the mononuclear nickel complexes (3) and (4) with one equivalent of commercially available  $[\text{FeCp}(\text{CO})_2]$  in dichloromethane provided the corresponding  $[\text{Ni}(\text{L})\text{FeCpCO}]$  complexes; subsequently the counter anion was exchanged by the addition of  $\text{NH}_4\text{PF}_6$  in acetonitrile resulting in the compounds  $[\text{Ni}(\text{pbSmSe})\text{FeCpCO}](\text{PF}_6)$  (5) and  $[\text{Ni}(\text{xbSmSe})\text{FeCpCO}](\text{PF}_6)$  (6). These heterodinuclear complexes were characterized with NMR and FTIR spectroscopy, mass spectrometry, elemental analysis and single crystal X-ray diffraction. The  $^1\text{H}$  NMR spectra of the NiFe complexes recorded in dichloromethane are weak and poorly resolved, but show the expected signals of the cyclopentadienyl, aromatic and methyl protons; however, the methylene groups which are in close proximity to nickel center are not clearly discernible. Several attempts have been undertaken to obtain better quality  $^1\text{H}$  NMR spectra of the NiFe complexes; spectra were recorded in another solvent (DMSO) and were recorded at different temperatures, but unfortunately to no avail (ESI, Fig. S12–16†). Again, the broadening of these signals might be explained by tetrahedral distortions of the square-planar geometry, resulting in a fraction of the nickel centers to be in the high-spin state. In contrast to the mononuclear compounds this distortion is not fluxional, but immobilized by the bridging of the two selenolate donor atoms between the  $\text{Ni}(\text{II})$  and  $\text{Fe}(\text{II})$  centers (see below). In addition, the results of the mass analysis and IR spectra (see below) indicate that a dynamic equilibrium may exist of compounds that are the result of a disproportionation reaction of (5) or (6)  $[\text{Ni}(\text{L})\text{FeCpCO}]^+$  to form  $[\text{Ni}(\text{L})\text{FeCp}]^+$  and  $[\text{Ni}(\text{L})\text{FeCp}(\text{CO})_2]^+$ , which would also give rise to broad signals.

### Description of the structures

Single crystals of (3) suitable for X-ray structure determination were obtained by vapor diffusion of pentane into a dichloromethane solution of the complex; crystallographic and refinement data are provided in Table S1.† A projection of the molecular structure is given Fig. 4 and selected bond distances and angles are listed in Table 1. Complex (3) crystallizes in the space group  $P2_1$ ; the asymmetric unit contains one molecule of the mononuclear compound  $[\text{Ni}(\text{pbSmSe})]$ . The  $\text{Ni}(\text{II})$  center is in a square-planar geometry by the coordination of two selenolate and two thioether donor atoms. The Ni–Se bond distances are 2.2898(6) and 2.2910(6) Å; as expected for the larger ionic radius of Se these distances are longer than the Ni–S bond distances of 2.180(8) Å in the thiolate analog  $[\text{Ni}(\text{pbSmS})]^{21}$ . As a result, the Ni–S thioether bond distances at 2.1707(10) and 2.1608(11) Å are slightly shorter than those of

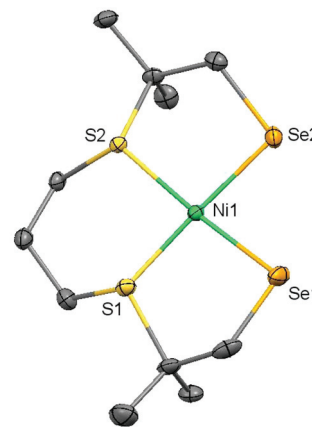


Fig. 4 Displacement ellipsoid plot (50% probability level) of (3) at 110(2) K. Hydrogen atoms are omitted for clarity.

Table 1 Selected bond lengths (Å) and angles (°) for the complexes (3), (5) and (6)

	(3)	(5)	(6)
<b>Distances (Å)</b>			
Ni1–Se1	2.2898(6)	2.2768(8)	2.2837(5)
Ni1–Se2	2.2910(6)	2.2978(7)	2.2933(5)
Ni1–S1	2.1608(11)	2.1996(11)	2.1835(7)
Ni1–S2	2.1707(10)	2.1817(12)	2.1820(8)
Fe1–Se1		2.3859(7)	2.4043(5)
Fe1–Se2		2.4018(8)	2.3923(5)
Fe1–Cp(centroid)		1.71(4)	1.70(3)
Fe1–C12/C22		1.741(4)	1.773(3)
<b>Angles (°)</b>			
S1–Ni1–Se1	89.79(3)	92.75(4)	91.31(2)
S2–Ni1–Se2	90.79(3)	93.00(4)	91.74(2)
Se1–Ni1–Se2	87.62(2)	82.52(3)	80.550(17)
S1–Ni1–S2	91.80(4)	91.27(4)	95.02(3)
S2–Ni1–Se1	170.94(4)	173.92(4)	167.66(3)
S1–Ni1–Se2	177.40(4)	172.33(4)	169.19(3)
Se1–Fe1–Se2		78.11(2)	76.174(15)
Ni1–Se1–Fe1		85.40(3)	81.074(17)
Ni1–Se2–Fe1		84.58(3)	81.139(17)

2.1711(3) and 2.1668(3) Å in  $[\text{Ni}(\text{pbSmS})]^{21}$ . The square-planar geometry reveals a slight tetrahedral distortion with a dihedral angle of 8.91°, defined by the planes S1–Ni1–S2 and Se1–Ni1–Se2, which is slightly larger than in the thiolate analog having a dihedral angle of 5.17°. This larger tetrahedral distortion in the solid state of the selenolate compound is rather surprising, as the NMR spectra of the thiolate compound are broadened due to the fluxionality of the ligand, which results in the nickel ion in the low-spin square-planar structure to be in equilibrium with a nickel center in a more tetrahedral high-spin state. In contrast, the NMR spectra show the selenolate compound to be clearly low-spin and diamagnetic, which may indicate that the tetrahedral distortion in the solid state is merely due to packing effects.

Single crystals of the complexes (5) and (6) were obtained by vapor diffusion of diethyl ether into dichloromethane solutions of the complexes; crystallographic and refinement data



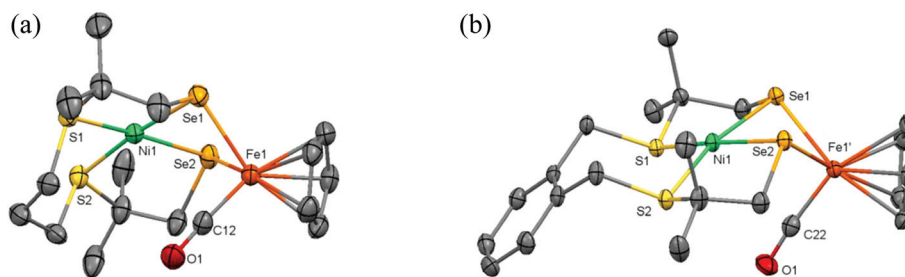


Fig. 5 Displacement ellipsoids plots (50% probability level) of the cationic complex in (a) (5) and (b) (6) at 110(2) K. Hydrogen atoms,  $\text{PF}_6^-$  anions, lattice solvent molecules, and disorder (in (6)) are omitted for clarity.

are provided in Table S1.† Projections of the molecular structures of the heterodinuclear complexes are shown in Fig. 5; selected bond distances and angles are listed in Table 1. Complex (5) crystallizes in the space group  $R\bar{3}$  and the crystal lattice contains some amount of significantly disordered solvents molecules. Complex (6) crystallizes in the space group  $P2_1/c$ ; the coordination spheres around Ni1 and the CO coordinated to Fe1 are found to be slightly disordered over two orientations. The nickel(II) ions in the complexes (5) and (6) are in square-planar geometries bound to two thioether and two selenolate donor atoms. In both compounds the two Se donor atoms are bridging to the Fe(II) ion, resulting in a Ni–Se–Fe ‘butterfly’ core with a ‘hinge’ angle (defined by the angle between the planes through  $\text{NiSe}_2$  and  $\text{FeSe}_2$ ) of  $120.87^\circ$  for (5) and  $108.36^\circ$  for (6). The Fe(II) ion in both complexes is further coordinated to a symmetrically bound  $\eta^5$ -cyclopentadienyl ligand and a CO ligand. The Ni–Se distances of 2.2837 (5) and 2.2933(5) Å in  $[\text{Ni}(\text{xbSmSe})\text{FeCpCO}]\text{PF}_6$  are longer than the corresponding Ni–S thiolate distances of 2.1670(9) and 2.1717(8) Å found in  $[\text{Ni}(\text{xbSmS})\text{FeCpCO}]\text{BF}_4$ ,<sup>11</sup> as a result of the larger ionic radius of the selenolate donor atom. For both NiFe complexes the square-planar geometry of the nickel centers is slightly distorted, with dihedral angles of  $7.39^\circ$  and  $12.63^\circ$  for complexes (5) and (6) respectively. This distortion seems to be caused by the bridging of both selenolate atoms between the Ni(II) and Fe(II) centers, resulting in larger S–Ni–Se and significantly smaller Se–Ni–Se angles. In both complexes the molecule of CO is directed towards the Ni center with a Ni–C(O) distance of 3.1 Å for complex (5) and 2.9 Å for complex (6). The major difference between the two heterodinuclear compounds is the relative orientation of the  $\{\text{FeCpCO}\}$  group. Whereas in complex (5) the  $\{\text{FeCpCO}\}$  group and the bridge between the sulfur atoms are on the same side of the Ni square plane, in complex (6) they are on opposite sides.

### IR spectroscopy of the NiFe complexes

The carbonyl stretching bands in the solid state IR spectra of compound (5) appear at  $1918\text{ cm}^{-1}$  (m),  $1987\text{ (s)}$  and  $2035\text{ (s)}$ , whereas the IR spectrum of compound (6) reveals one strong band at  $1923\text{ cm}^{-1}$  in addition two weaker bands at  $1989$  and  $2037\text{ cm}^{-1}$  (ESI, Fig. S7 and 8†). Based on its structure complex (5) should show only a single CO band. Indeed, the compound  $[\text{Ni}(\text{xbSmS})\text{FeCpCO}]\text{BF}_4$  has been reported to show a single car-

bonyl stretching band in the IR spectrum at  $1939\text{ cm}^{-1}$ .<sup>11</sup> However, the compound  $[\text{Ni}(\text{xbSmS})\text{FeCp}(\text{CO})_2]\text{BF}_4$  was reported to be an intermediate in the formation of  $[\text{Ni}(\text{xbSmS})\text{FeCpCO}]\text{BF}_4$ , showing two CO stretching bands at  $2008$  and  $2054\text{ cm}^{-1}$ . Interestingly, the mass spectra recorded of crystals of compound (5) show three major fragments: two of them are assigned to the expected monocationic compound  $[\text{M} - \text{PF}_6]^+$  and the compound after loss of the CO ligand  $[\text{M} - (\text{PF}_6) - (\text{CO})]^+$  (ESI Fig. S19†). Unexpectedly, also a fragment that can be assigned to the monocationic compound with two CO ligands ( $[\text{M} - (\text{PF}_6) + (\text{CO})]^+$ ) is observed. This fragment must be the result of disproportionation of  $[\text{Ni}(\text{pbSmSe})\text{FeCpCO}]^+$  to form  $[\text{Ni}(\text{pbSmSe})\text{FeCp}]^+$  and  $[\text{Ni}(\text{pbSmSe})\text{FeCp}(\text{CO})_2]^+$ , the former compound would contain a 16-electron  $\text{Fe}^{\text{II}}$  center, whereas the second species would be asymmetrically bridged by only one of the selenolate atoms. In contrast, whereas the IR of compound (6) indicates that a product containing two molecules of CO bound to iron may be present, the mass spectrum of (6) does not show a peak that can be assigned to a fragment  $[\text{M} - (\text{PF}_6) + (\text{CO})]^+$ . We therefore have to conclude that whereas for compound (6) the product with one CO ligand bound to iron is the most stable one, for compound (5) a mixture of  $[\text{Ni}(\text{pbSmSe})\text{FeCpCO}](\text{PF}_6)$  and  $[\text{Ni}(\text{pbSmSe})\text{FeCp}(\text{CO})_2](\text{PF}_6)$  is obtained, from which a single crystal of the monocarbonyl complex was picked. Because of the small mass difference of one molecule of CO the elemental analysis is not conclusive.

Similar values of IR stretching bands have been reported for the carbonyl ligands in the active sites of the  $[\text{NiFe}]$  and  $[\text{NiFeSe}]$  hydrogenases as well as for another structural model of  $[\text{NiFeSe}]$  hydrogenases.<sup>7</sup> The relatively lower energy of the CO stretching frequencies in the selenolate compounds has been attributed to an increase of electron density at the Fe center, as the selenolate donor atoms are more electron-donating than thiolate donor atoms.<sup>7</sup>

### Electrochemical analyses

The electrochemical properties of the nickel and nickel-iron complexes were investigated using cyclic voltammetry; the relevant data are presented in Table 2. For the mononuclear complex (3) one reversible reduction wave is observed with an  $E_{\text{pc}}$  at  $-2.1\text{ V vs. Fc}^+/\text{Fc}$  at a scan rate of  $200\text{ mV s}^{-1}$  in DMF (Fig. 6a), which is tentatively ascribed to the  $\text{Ni}^{\text{II}}/\text{Ni}^{\text{I}}$  redox



**Table 2** Electrochemical data of the Ni and NiFe complexes (vs.  $\text{Fc}^+/\text{Fc}$ )<sup>a</sup>

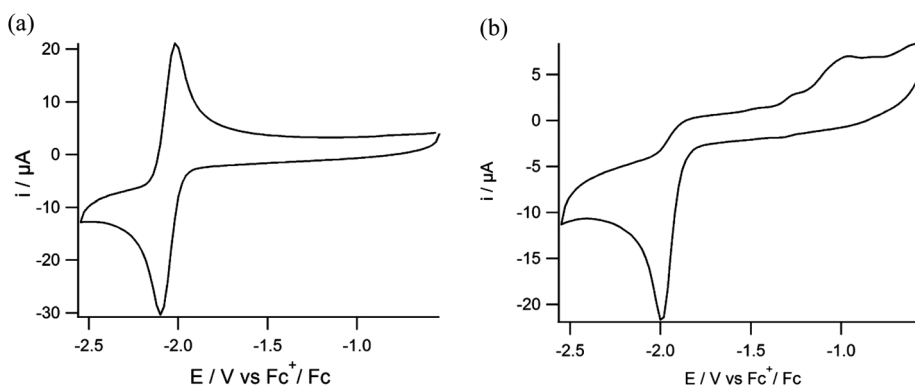
Compound	$E_{\text{pa}}$ (V)	$E_{\text{pc}}$ (V)
(3)	-2.02	-2.10
(4)		-1.99
(5)	-2.03	-2.10
(6)		-1.99

<sup>a</sup> Experimental conditions: 1 mM solutions of complexes in DMF containing 0.1 M TBAPF<sub>6</sub> as the supporting electrolyte. Glassy carbon working electrode, platinum counter electrode, Ag/AgCl reference electrode and scan rate 200 mV s<sup>-1</sup>. The values have been calculated using  $\text{Fc}/\text{Fc}^+$  as an internal reference, which was found to have  $E_{1/2}$  of 0.54 V vs. Ag/AgCl in our conditions.

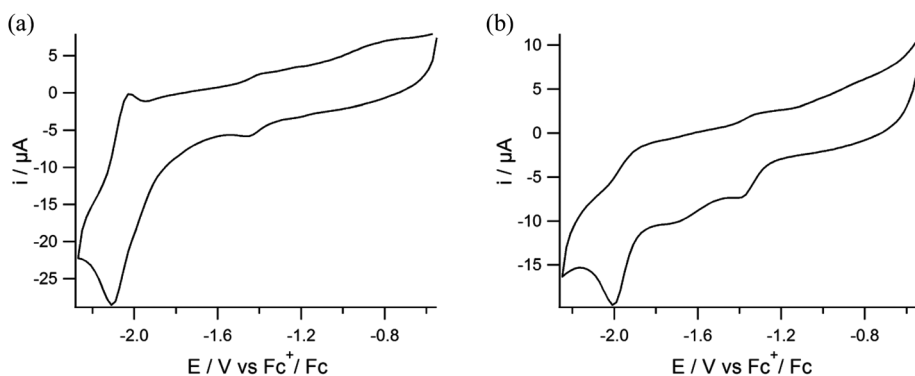
couple. In contrast, the corresponding thiolate-containing compound [Ni(pbSmS)] has been reported to show two irreversible waves at -1.05 V and -1.5 V vs.  $\text{Fc}^+/\text{Fc}$  at a scan rate of 100 mV s<sup>-1</sup> in dichloromethane solution.<sup>21</sup> We could not readily find an explanation for the large difference of nearly 1 V between the observed reduction potential of (3) and the values reported for [Ni(pbSmS)]. Therefore a CV of complex (3) was also recorded in dichloromethane solution (ESI, Fig. S3†). Although the reduction of (3) appeared to be irreversible in

dichloromethane, the reduction potential of (3) was found to be similar in both dichloromethane and in DMF solution. For the nickel complex (4) one irreversible wave is observed with an  $E_{\text{pc}}$  at -1.99 V vs.  $\text{Fc}^+/\text{Fc}$  at a scan rate of 200 mV s<sup>-1</sup> in DMF (Fig. 6b), similar to the irreversible electrochemical behavior reported for the analogous thiolate compound [Ni(xbSmS)] at -2.03 V vs.  $\text{Fc}^+/\text{Fc}$  in DMF solution.<sup>11</sup> The slightly less negative reduction potential for the nickel center in compound (4) relative to that in (3) may tentatively be ascribed to larger flexibility of the 7-membered chelate ring of the xylyl backbone, facilitating a change in redox state of the nickel center. We do not have an explanation for the differences in reversibility of the reduction wave of the nickel centers in (3) and (4), nor for the observation that the reduction wave for (3) is reversible in DMF, whereas it is irreversible in DCM. Furthermore, we cannot give a reason for the apparent 1 V difference in the reduction potentials of [Ni(pbSmSe)] (3) and [Ni(pbSmS)], especially as the difference in reduction potentials for the xylene-bridged compounds [Ni(xbSmS)] and [Ni(xbSmSe)] (4) is negligible.

The same electrochemical conditions with a scan rate of 200 mV s<sup>-1</sup> were also used to study the electrochemical behavior of the NiFe complexes (5) and (6) in DMF solutions. For complex (5) one quasi-reversible wave is observed with an  $E_{\text{pc}}$  at -2.1 V vs.  $\text{Fc}^+/\text{Fc}$  (Fig. 7a) with an  $i_{\text{pc}}$  that is nearly two times



**Fig. 6** Cyclic voltammograms of (a) compound (3) and (b) compound (4) in DMF solutions containing TBAPF<sub>6</sub> (0.1 M) as the supporting electrolyte on a glassy carbon electrode at 200 mV s<sup>-1</sup>.



**Fig. 7** Cyclic voltammograms of (a) compound (5) and (b) compound (6) in DMF solutions containing TBAPF<sub>6</sub> (0.1 M) as the supporting electrolyte on a glassy carbon electrode at 200 mV s<sup>-1</sup>.



higher than the  $i_{pa}$ . For complex (6) one quasi-reversible wave is observed with an  $E_{pc}$  at  $-1.99$  V vs.  $Fc^+/Fc$  (Fig. 7b). At first sight it thus seems that the presence of the cyclopentadienyl-iron moiety does not influence the reduction potential of the nickel(II) ion. In addition for both complexes one small wave is observed at around  $-1.4$  V vs.  $Fc^+/Fc$ .

In order to better understand the electrochemical properties of the nickel-iron complexes, the electrochemical behavior of  $[FeCp(CO)_2I]$  was also examined (Fig. S4†). For this compound one irreversible reduction is observed with an  $E_{pc}$  at  $-2.05$  V vs.  $Fc^+/Fc$  at a scan rate of  $200$  mV  $s^{-1}$  in DMF, which is tentatively ascribed to the  $Fe^{II}/Fe^I$  redox couple. Similar to the nickel-iron complexes, an additional small wave is observed at  $-1.3$  V vs.  $Fc^+/Fc$ . This feature thus indicates that the small wave around  $-1.4$  V vs.  $Fc^+/Fc$  in the voltammograms of the nickel-iron complexes is related to the presence of the  $[FeCp(CO)I]$  moiety. It seems that the quasi-reversible reduction of complex (5) with an  $E_{pc}$  at  $-2.1$  V vs.  $Fc^+/Fc$  is an overlap of the reduction of the nickel ion in (3) ( $E_{pc}$  at  $-2.1$  V vs.  $Fc^+/Fc$ ) with the reduction wave of iron center in  $[FeCp(CO)_2I]$  (at  $-2.05$  V vs.  $Fc^+/Fc$ ); upon closer inspection of the reduction wave for compound (5) it looks as if this wave indeed reveals a shoulder around  $-2.0$  V. This overlap of two redox events may also explain why the reversibility of this reductive peak changed from reversible in the mononuclear complex  $[Ni(pbSmSe)]$  to quasi-reversible in the heterodinuclear compound (5). In comparison, the complex  $[Ni(xbSmS)FeCpCO]BF_4$  has been reported to show one reversible redox couple at  $-1.43$  V vs.  $Fc^+/Fc$  and one irreversible wave at  $-2.01$  V vs.  $Fc^+/Fc$  in DMF.<sup>11</sup> In contrast to the CVs of the complexes (5) and (6), in this report the reduction wave at  $-1.43$  V vs.  $Fc^+/Fc$  has the same intensity as the one at  $-2.01$  V vs.  $Fc^+/Fc$ . The reduction wave of complex (6) appears to become more reversible compared to that of the corresponding mononuclear nickel complex  $[Ni(xbSmSe)]$ .

### Electrocatalytic hydrogen evolution in the presence of HOAc

The activity of the new compounds in electrocatalytic proton reduction was studied using cyclic voltammetry with addition of varying amounts of HOAc to DMF solutions of the Ni and NiFe complexes. The reversible reduction observed for complex

(3) with an  $E_{pc}$  at  $-2.1$  V vs.  $Fc^+/Fc$  becomes irreversible with the addition of HOAc (ESI, Fig. S1†). On the other hand, for complex (4) the irreversible reduction peak with an  $E_{pc}$  at  $-1.99$  V does not change upon addition of HOAc and no catalytic current is observed (ESI, Fig. S2†). The quasi-reversible reduction peak of complex (5) with an  $E_{pc}$  at  $-2.1$  V vs.  $Fc^+/Fc$  becomes irreversible with increasing concentrations of HOAc while the  $E_{pc}$  shifts to more negative potentials indicating electrocatalytic activity (Fig. 8). Again, for complex (6) no catalytic wave is observed upon the addition of different equivalents of acid.

To quantify the rate of the hydrogen evolution reaction the foot-of-the-wave (FOWA) analysis was applied.<sup>23,24</sup> The FOWA analysis was developed by Costentin and Savéant and can be used for the analysis of voltammograms that do not show an S-shaped curve with a fixed plateau current.<sup>23</sup> Using the FOWA analysis a  $k_{obs}$  of  $24$  s $^{-1}$  was calculated for complex (5) (ESI, Fig. S5†). In order to confirm that indeed dihydrogen gas is formed in the catalytic reaction, a controlled-potential coulometry (CPC) experiment was carried out on a  $0.5$  mM solution of complex (5) in DMF ( $5$  ml) in the presence of  $17.5$   $\mu$ l of HOAc ( $50$  equivalents of  $H^+$  per NiFe compound) at  $-2.1$  V vs.  $Fc^+/Fc$ . The produced dihydrogen gas was quantified volumetrically by GC analysis. The CPC experiment was run for  $50$  min, while the solution was stirred continuously. Using complex (5) as the electrocatalyst for proton reduction, a total of  $64$   $\mu$ l  $H_2$  was produced by  $0.5$  mM complex in  $50$  min with  $72\%$  faradaic yield. In a control experiment at this potential formation of  $H_2$  is not observed in the absence of the catalyst. In order to compare the activity of the mononuclear  $[Ni(pbSmSe)]$  and the dinuclear compound  $[Ni(pbSmSe)FeCpCO]PF_6$  a CPC experiment was also run for complex (3). After  $50$  min the amount of  $H_2$  produced by compound (3) appeared to be very low compared to the NiFe complex (5); the formed  $H_2$  was barely detectable with a concentration falling out of the lower range of the calibration line. A CPC experiment was also carried out using complex (6) in DMF solution in the presence of HOAc at  $-1.9$  V vs.  $Fc^+/Fc$ . In this case dihydrogen evolution was not observed and it can be concluded that this compound is not active as an electrocatalyst for proton reduction with a weak acid such as HOAc. In contrast, the

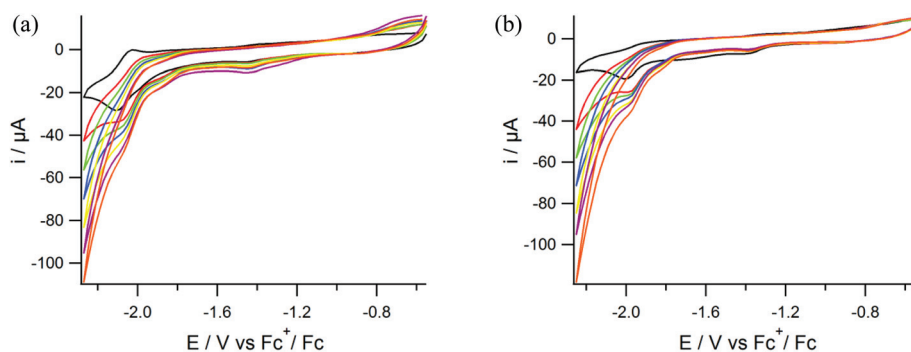


Fig. 8 Cyclic voltammograms of (a) compound (5) and (b) compound (6) in DMF solutions of TBAPF<sub>6</sub> ( $0.1$  M) on a glassy carbon electrode at  $200$  mV  $s^{-1}$  in the presence of  $0$  (black),  $10$  (red),  $20$  (green),  $30$  (blue),  $40$  (yellow),  $50$  (purple),  $60$  (orange) mM of acetic acid.



complex  $[\text{Ni}(\text{xbsmS})\text{FeCpCO}]\text{BF}_4$ , the thiolate analogue of complex (6), has been reported to be an electrocatalyst for  $\text{H}_2$  evolution, which achieved 20 turnovers in 4 h in the presence of the stronger acid trifluoroacetic acid.<sup>11</sup>

## Conclusion

Two novel NiFe complexes are reported as mimics of the  $[\text{NiFeSe}]$  hydrogenase active site. Both  $[\text{Ni}(\text{pbSmSe})\text{FeCpCO}]\text{PF}_6$  and  $[\text{Ni}(\text{xbsmSe})\text{FeCpCO}]\text{PF}_6$  complexes have structural similarities with active site of  $[\text{NiFeSe}]$  hydrogenase. The compounds contain nickel and iron centers coordinated with two selenolate and two thioether donors. The bond distances between the nickel centers and the selenolate donors are 2.29 Å for complexes (5) and (6), whereas the Ni–Se distance in the active site of  $[\text{NiFeSe}]$  hydrogenase has been reported to be 2.46 Å.<sup>5</sup> The major drawback of the selenium complexes compared to the reported sulfur analogues is their higher air sensitivity. The compound  $[\text{Ni}(\text{pbSmSe})\text{FeCpCO}]\text{PF}_6$  catalyzes the electrocatalytic reduction of protons in the presence of acetic acid, as shown by CV and CPC experiments, whereas  $[\text{Ni}(\text{xbsmSe})\text{FeCpCO}]\text{PF}_6$  does not. The reason for this difference in activity may lie in the disproportionation reaction that seemingly occurs for compound (5) but not for (6), generating an active 16-electron species for (5). To the best of our knowledge compound (5) is thus the first functional model of the  $[\text{NiFeSe}]$  hydrogenase active site. Although this NiFe compound is just a very modest catalyst, this work is an initial step for the development of more efficient mimics of the  $[\text{NiFeSe}]$  hydrogenase active site.

## Experimental

### Materials

All experiments were performed using standard Schlenk techniques or in a glovebox under an argon atmosphere unless otherwise noted. Chemicals were purchased from Acros or Aldrich and were used without further purification. Organic solvents were deoxygenated by the freeze–pump–thaw method and were dried over molecular sieves prior to use. The NMR solvent  $\text{CD}_2\text{Cl}_2$  for the metal complexes was deoxygenated by the freeze–pump–thaw method and was stored over molecular sieves in a glovebox. The compounds 1,9-dichloro-3,7-dithia-2,2,8,8-tetramethylnonane,<sup>21</sup> bis(3-chloro-2,2-methyl-1-thiopropyl)-*o*-xylene,<sup>22</sup> and  $[\text{Ni}(\text{xbsmSe})]_7$  were synthesized according to published methods.  $[\text{FeCp}(\text{CO})_2]_2$  was purchased from Aldrich. The synthesis of the  $[\text{NiFe}]$  complexes is based on a method described in literature.<sup>20</sup>

### Physical measurements

NMR spectra were recorded on a 300 MHz Bruker DPX 300 spectrometer and chemical shifts were referenced against the solvent peaks. Mass spectra were obtained with a Finnigan TSQ quantum instrument using ESI. HRMS was recorded

on a Thermo Scientific LTQ Orbitrap XL high resolution FT-MS system. Elemental analyses were performed by the Microanalytical Laboratory Kolbe in Germany. IR spectra were recorded on a PerkinElmer UATR Two FT-IR spectrometer. Electrochemical measurements were performed at room temperature under argon using an Autolab PGstat10 potentiostat controlled by GPES4 software. A three-electrode cell system was used with a glassy carbon working electrode, a platinum counter electrode and an Ag/AgCl reference electrode. All electrochemistry measurements were done in DMF solution with tetrabutylammonium hexafluoridophosphate as the supporting electrolyte. All potentials are referenced to the internal reference system  $\text{Fc}^+/\text{Fc}$ , which under these conditions was found at 0.54 V vs. Ag/AgCl in DMF. Electrocatalysis experiments were carried out by adding different concentrations of acetic acid to the DMF solution of complexes. Controlled-potential coulometry (CPC) experiments were done with the same three-electrode cell system and electrodes. CPC experiments were recorded with an Autolab PGstat10 potentiostat controlled by GPES4 software. Gas chromatographic analysis was performed on a Shimadzu gas chromatograph GC-2010 at 35 °C fitted with a Supelco Carboxen 1010 molecular sieve column. Helium was used as the carrier gas, and compounds were detected using a thermal conductivity detector operated at 80 mA. The total volume of  $\text{H}_2$  produced during the reaction was calculated using a calibration line, which was obtained using the external reference method by injection of known amounts of  $\text{H}_2$  into the GC using a Hamilton gas-tight syringe (see ESI Fig. S6†). A solution of complexes (5) or (6) in DMF (5 ml, 0.5 mM) was placed into a three-electrode cell and prior to each measurement the system was deaerated by bubbling with helium for 10 min. The system was closed, and the headspace was pumped through the solution for 1 min. Afterward, the headspace pumping was temporarily stopped to allow equilibration of the pressure, then the GC measurement was started with a 0.5 mL sample of the headspace injection. The GC valve and the pump (KNF NMS 010 L micro diaphragm pump) were enclosed in a helium-purged housing to prevent air from leaking into the system.

### Crystal structure determinations

All reflection intensities were measured at 110(2) K using a SuperNova diffractometer (equipped with Atlas detector) with Mo  $\text{K}\alpha$  radiation ( $\lambda = 0.71073$  Å) for (3) and Cu  $\text{K}\alpha$  radiation ( $\lambda = 1.54178$  Å) for (5) and (6) under the program CrysAlisPro (Version 1.171.36.32 Agilent Technologies, 2013). The same program was used to refine the cell dimensions and for data reduction. The structures were solved with the program SHELXS-2014/7 and were refined on  $F^2$  with SHELXL-2014/7.<sup>25</sup> Numerical absorption correction based on gaussian integration or Analytical numeric absorption correction over a multifaceted crystal model was applied using CrysAlisPro. The temperature of the data collection was controlled using the system Cryojet (manufactured by Oxford Instruments). The H atoms were placed at calculated positions using the instructions AFIX 23, AFIX 43 or AFIX 137 with isotropic displacement



parameters having values 1.2 or 1.5  $U_{\text{eq}}$  of the attached C atoms.

#### Additional notes on the structure determinations

(3) The structure was refined in the space group  $P2_1$ . The absolute configuration was established by anomalous-dispersion effects in diffraction measurements on the crystal. The Flack parameter refines to  $-0.002(5)$ . CCDC 1537790 contains the supplementary crystallographic data for  $[\text{Ni}(\text{pbSmSe})]$ .

(5) The crystal lattice contains some amount of significantly disordered solvent molecules found in 'channels' along the  $c$  direction. Their contribution has been taken out using the SQUEEZE (Spek, 2009) procedure in the final refinement.<sup>26</sup>

(6) The coordination sphere around the Ni center in the complex as well as the CO coordinated to the iron center are found to be slightly disordered over two orientations; the occupancy factor of the major component of the disorder refines to 0.9585(6). Disorder may occur as both orientations have very similar space-filling requirements. CCDC 1537791 and 1537792 contain the supplementary crystallographic data for  $[\text{Ni}(\text{pbSmSe})\text{Fe}(\text{CO})\text{Cp}](\text{PF}_6)$  and  $[\text{Ni}(\text{xbSmSe})\text{Fe}(\text{CO})\text{Cp}](\text{PF}_6)$ .

#### Synthesis of ligand precursor (1)

A solution of selenourea (1.2 g, 9.6 mmol) in 10 ml ethanol was added to a solution of 1,9-dichloro-3,7-dithia-2,2,8,8-tetramethylnonane (1.4 g, 4.83 mmol) in 5 ml ethanol; the reaction mixture was refluxed for 30 min. The solution was allowed to cool, and the solid product was isolated by filtration. The product was washed with cold ethanol and diethyl ether, and dried *in vacuo* yielding a white powder. Yield: 2.2 g (86%).  $^1\text{H}$  NMR (300 MHz,  $(\text{CD}_3)_2\text{SO}$ )  $\delta$  (ppm): 9.41 (d,  $J = 11.3$  Hz, 8H, NH), 3.61 (s, 4H,  $\text{CH}_2$ ), 2.66 (t,  $J = 7.2$  Hz, 4H,  $\text{CH}_2$ ), 1.73 (q,  $J = 7.1$  Hz, 2H,  $\text{CH}_2$ ), 1.66 (s, 12H,  $\text{CH}_3$ ).  $^{13}\text{C}$  NMR (75 MHz,  $(\text{CD}_3)_2\text{SO}$ ):  $\delta$  (ppm) = 167.02 (C(NH)NH<sub>2</sub>), 46.31 ( $\text{CH}_2$ -Se), 40.57 ( $\text{CH}_2$ -S), 29.67 ( $\text{CH}_2$ - $\text{CH}_2$ S), 28.07 ( $\text{CH}_3$ ). ESI-MS ( $\text{H}_2\text{O}$ ): 233.1, calcd: 233.01  $[\text{M} - 2\text{Cl}]^{2+}$ .

#### Synthesis of $[\text{Ni}(\text{pbSmSe})]$ (3)

A solution of  $\text{NMe}_4\text{OH}$  (558 mg, 3.08 mmol), ligand precursor (1) (824 mg, 1.54 mmol) and  $\text{Ni}(\text{acac})_2$  (396 mg, 1.54 mmol) in 170 ml ethanol was refluxed for 1 h. The solvent was evaporated until approximately 30 ml solvent remained, resulting in a green precipitate. The solid was collected by filtration and washed with ethanol. Yield: 429 mg (64%).  $^1\text{H}$  NMR (300 MHz,  $\text{CD}_2\text{Cl}_2$ )  $\delta$  (ppm): 2.64 (t,  $J = 7.1$  Hz, 4H,  $\text{CH}_2$ -S), 2.41 (s, 4H,  $\text{CH}_2$ -Se), 2.22 (m, 2H,  $\text{CH}_2$ - $\text{CH}_2$ S), 1.57 (s, 12H,  $\text{CH}_3$ ).  $^{13}\text{C}$  NMR (75 MHz,  $\text{CD}_2\text{Cl}_2$ ):  $\delta$  (ppm): 29.19 ( $\text{CH}_2$ ,  $\text{CH}_2$ - $\text{CH}_2$ S), 28.80 ( $\text{CH}_2$ ,  $\text{CH}_2$ -Se), 26.70 ( $\text{CH}_3$ ), 25.39 ( $\text{CH}_2$ - $\text{CH}_2$ S). HR-MS ( $\text{CH}_2\text{Cl}_2$ ): 436.89255, calcd: 436.89196  $[\text{M} + \text{H}]^+$ . Elemental analysis calcd (%) for  $\text{C}_{11}\text{H}_{22}\text{NiS}_2\text{Se}_2 \cdot 0.2 \text{N}(\text{CH}_3)_4\text{Cl}$ : C: 31.02 H: 5.38; found C: 30.73 H: 5.86.

#### Synthesis of $[\text{Ni}(\text{pbSmSe})\text{Fe}(\text{CO})\text{Cp}](\text{PF}_6)$ (5)

A solution of  $[\text{Ni}(\text{pbSmSe})]$  (400 mg, 0.91 mmol) and  $[\text{FeCp}(\text{CO})_2\text{I}]$  (279 mg, 0.91 mmol) in 25 ml dichloromethane was stirred at RT for 2 days. The mixture was filtered to remove

an insoluble precipitate and the solvent was evaporated using the Schlenk line. The resulting solid was washed with diethyl ether resulting in a brown precipitate which was dried *in vacuo*. A solution of  $\text{NH}_4\text{PF}_6$  (300 mg, 1.84 mmol) in 10 ml acetonitrile was added to the brown solid and the mixture was stirred for 4 h. The solvent was evaporated until dryness, the remaining solid was dissolved in dichloromethane (5 ml) and the solution was filtered to remove  $\text{NH}_4\text{I}$ . A large amount (~30 ml) of diethyl ether was added into the dichloromethane solution, and the mixture was cooled at  $-35$  °C overnight. The resulting brown precipitate was collected by filtration and dried *in vacuo*. Yield: 105 mg (20%).  $^1\text{H}$  NMR (300 MHz,  $\text{CD}_2\text{Cl}_2$ )  $\delta$  (ppm): 4.75 (s, Cp). ESI-MS ( $\text{CH}_3\text{CN}$ ): 584.9, calcd: 584.9  $[\text{M} - \text{PF}_6]^+$ , 612.9, calcd: 612.9  $[\text{M} - (\text{PF}_6) + (\text{CO})]^+$  and 556.9, calcd: 556.9  $[\text{M} - (\text{PF}_6) - (\text{CO})]^+$ . Elemental analysis calcd (%) for  $\text{C}_{17}\text{H}_{27}\text{F}_6\text{FeNiOPS}_2\text{Se}_2 \cdot 0.25(\text{C}_2\text{H}_5)_2\text{O}$ : C: 28.92 H: 3.98; found C: 29.00 H: 3.93. IR (neat):  $\tilde{\nu} = 2035$  (s), 1987 and 1918 (CO stretch)  $\text{cm}^{-1}$ , 830 ( $\text{PF}_6$  stretch)  $\text{cm}^{-1}$ .

#### Synthesis of $[\text{Ni}(\text{xbSmSe})\text{Fe}(\text{CO})\text{Cp}](\text{PF}_6)$ (6)

A solution of  $[\text{Ni}(\text{xbSmSe})]$  (200 mg, 0.40 mmol) and  $[\text{FeCp}(\text{CO})_2\text{I}]$  (122 mg, 0.40 mmol) in 15 ml dichloromethane was stirred at RT for 2 days. The mixture was filtered to remove an insoluble precipitate and the solvent was evaporated using the Schlenk line. The residue was washed with diethyl ether, resulting in a brown powder which was dried *in vacuo*. A solution of  $\text{NH}_4\text{PF}_6$  (132 mg, 0.81 mmol) in 5 ml acetonitrile was added to the brown solid and the resulting mixture was stirred for 4 h. The solvent was evaporated until dryness, the remaining solid was dissolved in dichloromethane (5 ml) and the solution was filtered to remove  $\text{NH}_4\text{I}$ . A large amount (~30 ml) of diethyl ether was added into the dichloromethane solution, and the mixture was cooled at  $-35$  °C overnight. The resulting brown precipitate was collected by filtration and dried *in vacuo*. Yield: 18 mg (6%).  $^1\text{H}$  NMR (300 MHz,  $\text{CD}_2\text{Cl}_2$ )  $\delta$  (ppm): 7.93 (m, Ar), 7.31 (m, Ar), 4.74 (s, Cp), 1.76 (s,  $-\text{CH}_3$ ), 1.66 (s,  $-\text{CH}_3$ ). ESI-MS ( $\text{CH}_3\text{CN}$ ): 619.0, calcd: 618.9  $[\text{M} - \text{CO} - \text{PF}_6]^+$ , 647.0, calcd: 646.9  $[\text{M} - \text{PF}_6]^+$ . Elemental analysis calcd (%) for  $\text{C}_{22}\text{H}_{29}\text{F}_6\text{FeNiOPS}_2\text{Se}_2 \cdot 0.3\text{CH}_2\text{Cl}_2$ : C: 32.56 H: 3.64; found C: 32.33 H: 3.77. IR (neat):  $\tilde{\nu} = 1923$  (CO stretch)  $\text{cm}^{-1}$ , 828 ( $\text{PF}_6$  stretch)  $\text{cm}^{-1}$ .

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