

Cite this: *Chem. Sci.*, 2019, 10, 5079

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# Regiodivergent hydrosilylation, hydrogenation, $[2\pi + 2\pi]$ -cycloaddition and C–H borylation using counterion activated earth-abundant metal catalysis†

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The widespread adoption of earth-abundant metal catalysis lags behind that of the second- and third-row transition metals due to the often challenging practical requirements needed to generate the active low oxidation-state catalysts. Here we report the development of a single endogenous activation protocol across five reaction classes using both iron- and cobalt pre-catalysts. This simple catalytic manifold uses commercially available, bench-stable iron- or cobalt tetrafluoroborate salts to perform regiodivergent alkene and alkyne hydrosilylation, 1,3-diene hydrosilylation, hydrogenation,  $[2\pi + 2\pi]$ -cycloaddition and C–H borylation. The activation protocol proceeds by fluoride dissociation from the counterion, *in situ* formation of a hydridic activator and generation of a low oxidation-state catalyst.

Received 3rd December 2018

Accepted 7th April 2019

DOI: 10.1039/c8sc05391j

rsc.li/chemical-science

## Introduction

The ubiquity of catalytic protocols using precious metals such as platinum, palladium and rhodium can be ascribed to the highly robust and reliable nature of these methods and widespread commercial availability of the catalyst precursors. Operational simplicity has made reaction screening and optimisation routine using these metals. However, beyond high value applications the low natural abundance, volatile cost and toxicity of these metals remains problematic. Earth-abundant transition metals offer an alternative, sustainable platform for catalysis, particularly for bulk and dispersive technologies, but are yet to achieve widespread adoption even though excellent catalytic activity has been achieved in exemplar industrial reactions.<sup>1</sup> Thus, the operational simplicity of earth-abundant-metal-catalysed reactions must be addressed to enable the widespread use and development of these powerful methodologies.

The global silicone industry is forecast to be worth \$18.3 billion in 2021 and finds applications in areas as diverse as soft materials, cosmetics and food additives.<sup>2</sup> Alkene and alkyne hydrosilylation reactions underpin this industry and the homogenous nature of these processes results in the loss of over 5 tonnes of platinum annually.<sup>3</sup> Thus a transition to earth-

abundant metal catalysis would be beneficial both environmentally and economically. Seminal studies using isolated, low oxidation-state iron- and cobalt pre-catalysts have shown the potential of these metals for alkene hydrosilylation,<sup>4</sup> and *in situ* activation of metal(II/III) pre-catalysts using organometallic reagents<sup>1b,5</sup> has decreased the operational barrier to use. Methods have also been developed using bench-stable reductants such as alkoxide reagents<sup>6</sup> or amines<sup>7</sup> (Scheme 1a). However, an additional, external reagent is still required for pre-catalyst activation. Iron- and cobalt carboxylate salts have been shown to act as pre-catalysts for alkene hydrosilylation that do not require an external activator (Scheme 1b).<sup>8</sup> Similarly, Huang showed that a tridentate PNN-cobalt(II) dichloride pre-catalyst could be activated thermally for alkene hydrosilylation.<sup>5b</sup> However, these are limited to a single reaction class and the carboxylate counterions are pre-catalyst specific.

The alkoxide activation of iron- and cobalt pre-catalysts (Scheme 1a) was proposed to proceed by reaction of the alkoxide and silane to form a hydridic silicon 'ate' complex which reduces the pre-catalyst by a hydride transfer.<sup>6a</sup> In contrast, the carboxylate and thermal activation methods were proposed to proceed by a  $\sigma$ -bond metathesis reaction between the metal carboxylate and silane (Scheme 1b).<sup>8</sup> We postulated that using a counterion which is known to dissociate a nucleophile would allow an activation method that combined the operational simplicity of carboxylate activation and the broad scope of alkoxide activation. The tetrafluoroborate counterion is known to dissociate to  $\text{BF}_3$  and fluoride.<sup>9</sup> The fluoride could react with the silane to give a hydridic silicon 'ate' complex,<sup>10</sup> and activate a pre-catalyst by hydride transfer (Scheme 1c).<sup>11</sup>

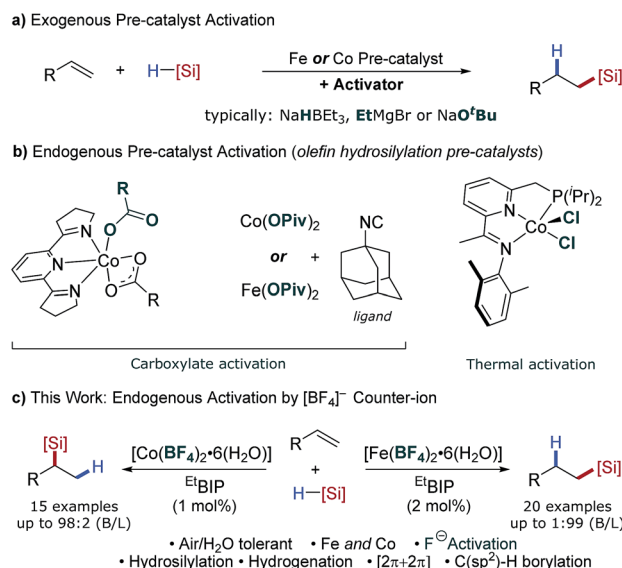
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† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8sc05391j

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**Scheme 1** Overview of prior alkene hydrosilylation reactions using iron- and cobalt catalysts. (a) Pre-catalyst activation using additives. (b) Catalyst activity facilitated by carboxylate ligands or thermal activation. (c) This work: catalyst activation through use of weakly coordinating tetrafluoroborate counterions.

## Results and discussion

In order to establish the utility of the tetrafluoroborate counterion for activation of iron(II) pre-catalysts we selected alkene hydrosilylation as a model reaction. Baseline reactivity was determined by the use of iron pre-catalysts bearing strongly coordinating chloride anions  $[^tBIPFeCl_2]$  (Table 1, entry 1) and weakly coordinating triflate anions  $[^tBIPFe(OTf)_2]$  (entry 2). Both reactions showed no catalytic activity. The tetrafluoroborate pre-catalyst, formed *in situ* by the reaction of the commercially available hydrate salt  $Fe(BF_4)_2 \cdot 6H_2O$  with bisiminopyridine ligand ( $^tBIP$ ), showed excellent catalytic activity without an external activator, giving the linear silane product **2a** in excellent yield and regioselectivity (Table 1, entry 3). Lower catalyst loadings showed good, but reduced, reactivity in the production of **2a** (Table 1, entries 4–5). Using iron(II) tetrafluoroborate, with less hindered bisiminopyridine ligands ( $^HBIP$ ,  $^MeBIP$ ,  $^MesBIP$ ) led to low reactivity, with only the *N*-mesityl ligand giving good yield, with similar linear:branched selectivity to that observed with  $^tBIP$ , while the more hindered *N*-isopropyl ligand gave only trace reactivity (Table 1, entries 6–9). Control reactions showed the need for both iron and ligand to achieve catalysis (see ESI, Table SI 1.1†).

Having established the tetrafluoroborate activation for iron-catalysed hydrosilylation, we next attempted to apply the same protocol to cobalt-catalysed alkene hydrosilylation. Using commercially available cobalt(II) tetrafluoroborate and the *N*-dimethylphenyl bisiminopyridine ligand gave the linear silane **3a** in excellent yield and good regioselectivity (Table 1, entry 11); the opposite to that observed under iron catalysis. Variation of the ligand *N*-aryl substituent showed that all cobalt catalysed systems selectively gave the branched silane, with the more hindered

**Table 1** Reaction optimisation for iron- and cobalt-catalysed hydro-silylation using tetrafluoroborate pre-catalyst activation<sup>a</sup>

$n-C_8H_{13} + PhSiH_3 \xrightarrow[THF, r.t., 4 h]{^tBIP (n \text{ mol\%}), [Fe^{II}] \text{ or } [Co^{II}] (n \text{ mol\%})} n-C_8H_{13}-CH_2-CH_2-PhSiH_2 + n-C_8H_{13}-CH_2-CH_2-PhSiH_3$

**2a**      **3a**

$^HBIP$  Ar = Ph  
 $^MeBIP$  Ar = 2,6-Me-Ph  
 $^MesBIP$  Ar = 2,4,6-Me-Ph  
 $^tBIP$  Ar = 2,6-Et-Ph  
 $^iPrBIP$  Ar = 2,6-*i*-Pr-Ph

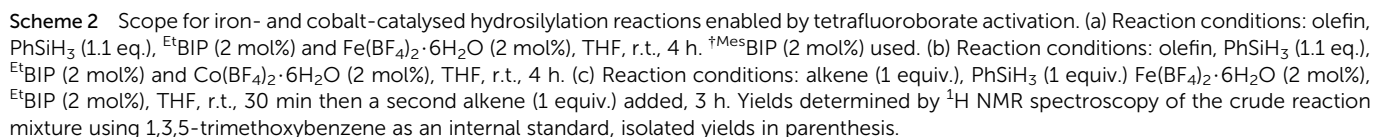
Entry	[M]	Loading (mol%)	Ligand	Yield (%) ( <b>2a</b> : <b>3a</b> )
1	$FeCl_2$	2	$^tBIP$	0
2	$Fe(OTf)_2$	2	$^tBIP$	0
3	$Fe(BF_4)_2 \cdot 6H_2O$	2	$^tBIP$	87 (93 : 7)
4	$Fe(BF_4)_2 \cdot 6H_2O$	0.5	$^tBIP$	67
5	$Fe(BF_4)_2 \cdot 6H_2O$	1	$^tBIP$	82
6	$Fe(BF_4)_2 \cdot 6H_2O$	2	$^HBIP$	0
7	$Fe(BF_4)_2 \cdot 6H_2O$	2	$^MeBIP$	Trace
8	$Fe(BF_4)_2 \cdot 6H_2O$	2	$^MesBIP$	78 (>95 : 5)
9	$Fe(BF_4)_2 \cdot 6H_2O$	2	$^iPrBIP$	Trace
10	$Co(BF_4)_2 \cdot 6H_2O$	2	$^tBIP$	84 (86 : 14)
11	$Co(BF_4)_2 \cdot 6H_2O$	2	$^HBIP$	72 (4 : 96)
12	$Co(BF_4)_2 \cdot 6H_2O$	2	$^MeBIP$	68 (3 : 97)
13	$Co(BF_4)_2 \cdot 6H_2O$	2	$^tBIP$	82 (5 : 95)
14	$Co(BF_4)_2 \cdot 6H_2O$	2	$^iPrBIP$	31 (16 : 84)
15	$Co(BF_4)_2 \cdot 6H_2O$	1	$^tBIP$	90 (8 : 92)

<sup>a</sup> Reaction conditions: 1-octene (1.00 equiv.), phenylsilane (1.10 equiv.) and metal tetrafluoroborate (*n* mol%), THF (1 M), r.t., 1 h. Yields determined by  $^1H$  NMR spectroscopy of the crude reaction mixture using 1,3,5-trimethoxybenzene as an internal standard.

catalysts giving reduced yields of silane **3a** (Table 1, entries 10–14). Notably, and to the best of our knowledge, this is the first example of a regioselective alkene hydrosilylation where high levels of regioselectivity are observed using an identical ligand and the same reaction conditions, but only varying the metal used.<sup>5b</sup> Use of a lower catalyst loading of 1 mol% gave the branched silane product in excellent yield and selectivity (Table 1, entry 15).

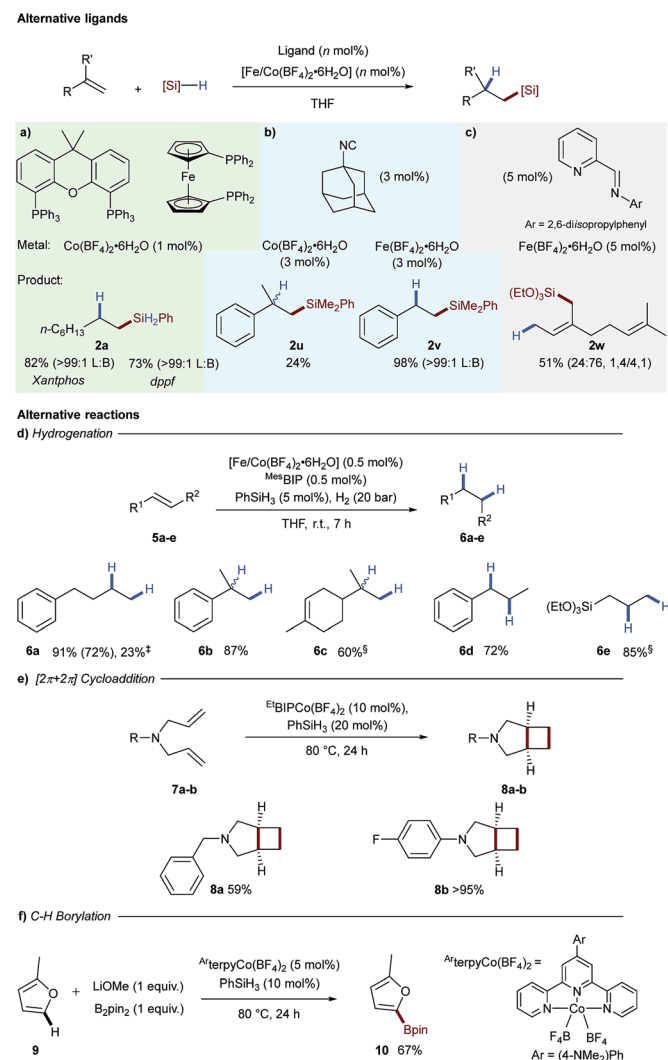
Having established optimal conditions for olefin hydrosilylation, we set out to investigate the scope and limitations of these reactions (Scheme 2). A variety of alkyl- and alkoxy-silane reagents were successfully used for alkene hydrosilylation with both iron- (anti-Markovnikov) and cobalt (Markovnikov) pre-catalysts providing the linear- **2a–p** and branched alkyl silanes **3a–l** in excellent yield and regioselectivity, respectively. Iron-catalysed hydrosilylation of 1-octene with a tertiary silane (triethoxysilane) gave the linear silane **2d** with a turnover number of 425 and a turnover frequency of  $5100 \text{ h}^{-1}$ . In the cobalt-catalysed system, using only 110 ppm cobalt with phenylsilane gave the branched silane **3a** with an overall turnover number of 5940, and turnover frequency of  $2970 \text{ h}^{-1}$ . As all reaction components are air- and moisture stable the reactions can be set up without the need for specialist equipment. Therefore the iron- and cobalt catalysed hydrosilylation of 1-octene **1a** was carried out in air with only a limited loss of catalyst activity and regioselectivity (Scheme 2, **2a** and **3a**). Alkyl- and aryl substituted alkenes underwent hydrosilylation in





The potential of the developed tetrafluoroborate activated hydrosilylations for polymer crosslinking and late-stage functionalisation was demonstrated by iterative homo- **2s** and

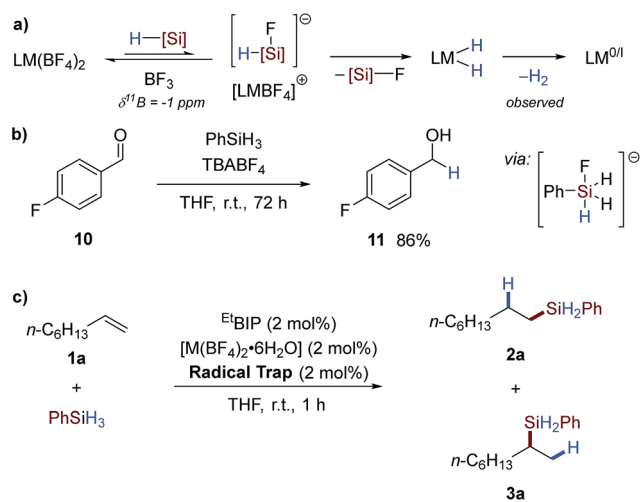
To assess the generality of tetrafluoroborate iron- and cobalt-salts for hydrosilylation we opted to perform alkene hydrosilylation reactions with ligands distinct from bisiminopyridine. We initially selected bis-phosphine ligands, xantphos and dppf, as these had been used previously by Ge for cobalt-catalysed alkene hydrosilylation.<sup>12</sup> Use of both of these ligands in combination with  $\text{Co}(\text{BF}_4)_2 \cdot 6\text{H}_2\text{O}$  effectively catalysed the hydrosilylation of 1-octene **1a** with phenylsilane to generate linear silane **2a** in excellent yield as a single exclusive regioisomer (Scheme 3a). Adamantyl isocyanide as a ligand additionally proved effective for both iron- and cobalt-catalysed alkene hydrosilylation to generate **2u** and **2v** respectively (Scheme 3b).<sup>13</sup> The bidentate iminopyridine ligand, used previously by Ritter for



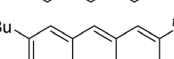


**Scheme 3** Application of tetrafluoroborate activation to other ligand classes for alkene hydrosilylation (top). Reaction conditions: (a) xantphos or dpfp (1 mol%),  $Co(BF_4)_2 \cdot 6H_2O$  (1 mol%), 1-octene (1 mmol), phenylsilane (1.1 mmol), THF (2 M), r.t., 4 h. (b) Adamantyl isocyanide (9 mol% [Co] or 6 mol% [Fe]), metal tetrafluoroborate (3 mol%),  $\alpha$ -methylstyrene ([Co], 1 mmol) or styrene ([Fe], 1 mmol), phenyldimethylsilane (1.3 mmol), THF (2 M), 80 °C, 3 h. (c)  $iPrIP$  (5 mol%),  $Fe(BF_4)_2 \cdot 6H_2O$  (5 mol%), myrcene (1 mmol), triethoxysilane (1.2 mmol), THF (2 M), r.t., 16 h. (d) Alkene (1 equiv.),  $Co(BF_4)_2 \cdot 6H_2O$  (0.5 mol%),  $Me^sBIP$  (1 mol%),  $PhSiH_3$  (5 mol%),  $H_2$  (20 bar), r.t., 7 h. <sup>†</sup>Alkene (1 equiv.),  $Fe(BF_4)_2 \cdot 6H_2O$  (2 mol%),  $Me^sBIP$  (2 mol%),  $PhSiH_3$  (5 mol%),  $H_2$  (20 bar), r.t., 7 h. <sup>§</sup> $Co(BF_4)_2 \cdot 6H_2O$  (1 mol%) and  $Me^sBIP$  (1 mol%). (e) 1,6-Diene (1 equiv.),  $EtBIPCo(BF_4)_2 \cdot 6H_2O$  (10 mol%),  $PhSiH_3$  (20 mol%), 80 °C, 24 h. (f) Arene (15 equiv.),  $(4-NMe_2-Ph-terpy)Co(BF_4)_2$  (5 mol%),  $PhSiH_3$  (20 mol%), LiOMe (1 equiv.),  $B_2pin_2$  (1 equiv.), 80 °C, 24 h.

1,4-hydrosilylation of 1,3-dienes,<sup>14</sup> was effective using  $Fe(BF_4)_2 \cdot 6H_2O$  to give allylic silane **2w** (Scheme 3c).

As we presumed the active hydrosilylation catalyst was a low oxidation-state metal species, we next explored the use of the tetrafluoroborate activation as a general platform to access these species. We postulated that mixing the tetrafluoroborate pre-catalysts with a substoichiometric amount of silane reagent would give a generic low oxidation-state catalyst that would be



Entry	M	Radical Trap	Yield (%) (3a:4a)
1	Fe		72 (92:8)
2	Co		68 (6:94)
3	Fe		86 (94:6)
4	Co		69 (3:97)
5	Fe		72 (90:10)
6	Co		35 (14:86)



boryl complexes have been proposed to be the key catalytic intermediate.<sup>18</sup> In order to apply tetrafluoroborate activation to a range of mechanistically distinct reactions, C–H borylation of 2-methylfuran **9** was carried out using a cobalt-terpyridine tetrafluoroborate pre-catalyst, to give the boronic ester **10** in 67% yield (Scheme 3f).

The facile activation observed using this protocol was thought to result from pre-catalyst reduction by an *in situ* formed hydridic silicon ‘ate’; formed by reaction of the silane reagent with fluoride dissociated from the counterion. This putative ‘ate’ complex then transfers hydride to the pre-catalyst, facilitating reductive elimination of dihydrogen (Scheme 4a).<sup>4b</sup> Silane reagents have been shown to be hydridic reagents in the presence of suitable nucleophiles, such as fluoride, and this has been applied in the reduction of carbonyls.<sup>10</sup> To examine whether similar reactivity could be obtained in this case, *n*-butylammonium tetrafluoroborate (TBABF<sub>4</sub>) was reacted with phenylsilane in the presence of 4-fluorobenzaldehyde **10** to give the primary alcohol reduction product **11** in excellent yield (Scheme 4b). This underlines the possibility of pre-catalyst reduction by a hydridic silicon ‘ate’ complex<sup>6a</sup> formed by reaction of fluoride and phenylsilane.<sup>11</sup> Although the exact nature of the active catalyst is not known, the reaction was tested in the presence of radical inhibitors and was unaffected (Scheme 4c and Table SI 10†), suggesting that reduction occurs by a two-electron mechanism.<sup>4b</sup>

## Conclusions

A procedure for the regiodivergent hydrosilylation of olefins, a highly valuable industrial reaction, has been developed using iron- and cobalt tetrafluoroborate catalysts without the need for an external activator or the use of isolated low oxidation-state complexes. This has been used as an activation platform to access the low oxidation-state catalysts in a range of iron- and cobalt-catalysed reactions including hydrosilylation, 1,3-diene hydrosilylation, alkene hydrogenation, [2π + 2π]-cycloaddition and C–H borylation. The developed tetrafluoroborate activation represents a versatile platform for activation, and serves as a generic strategy for accessing low oxidation-state reactivity with both iron and cobalt. It is hoped that this work will streamline the discovery of new reactivity, development of novel synthetic methodology and, ultimately, the replacement of precious metals with their earth-abundant counterparts.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

RA and SPT thank the Royal Society for funding a PhD studentship. AJC and SPT thank Syngenta for part funding a PhD studentship. SPT thanks the Royal Society for a University Research Fellowship. JD and SPT thank Prof. M. Shaver for useful discussions. All thank the University of Edinburgh, School of Chemistry and Technical Staff for generous support.

## Notes and references

- (a) I. Bauer and H.-J. Knölker, *Chem. Rev.*, 2015, **115**, 3170–3387; (b) X. Du and Z. Huang, *ACS Catal.*, 2017, **7**, 1227–1243; (c) R. I. Khusnutdinov, A. R. Bayguzina and U. M. Dzhemilev, *Russ. J. Org. Chem.*, 2012, **48**, 309–348; (d) J. R. Carney, B. R. Dillon and S. P. Thomas, *Eur. J. Org. Chem.*, 2016, **23**, 3912–3929; (e) B. D. Sherry and A. Fürstner, *Acc. Chem. Res.*, 2008, **41**, 1500–1511; (f) A. Fürstner, *ACS Cent. Sci.*, 2016, **2**, 778–789.
- Global Silicones Market by Market, Product and Country, 4th edn, Summary, 1.
- A. J. Holwell, *Platinum Met. Rev.*, 2008, **52**, 243–246.
- (a) A. M. Tondreau, C. C. H. Atienza, K. J. Weller, S. A. Nye, K. M. Lewis, J. G. P. Delis and P. J. Chirik, *Science*, 2012, **335**, 567–570; (b) S. C. Bart, E. Lobkovsky and P. J. Chirik, *J. Am. Chem. Soc.*, 2004, **126**, 13794–13807.
- (a) W. Y. Chu, R. Gilbert-Wilson, T. B. Rauchfuss, M. Van Gastel and F. Neese, *Organometallics*, 2016, **35**, 2900–2914; (b) X. Du, Y. Zhang, D. Peng and Z. Huang, *Angew. Chem., Int. Ed.*, 2016, **55**, 6671–6675; (c) M. D. Greenhalgh, D. J. Frank and S. P. Thomas, *Adv. Synth. Catal.*, 2014, **356**, 584–590; (d) X. Jia and Z. Huang, *Nat. Chem.*, 2016, **8**, 157–161; (e) J. Guo and Z. Lu, *Angew. Chem., Int. Ed.*, 2016, **55**, 10835–10838; (f) C. Chen, M. B. Hecht, A. Kavara, W. W. Brennessel, B. Q. Mercado, D. J. Weix and P. L. Holland, *J. Am. Chem. Soc.*, 2015, **137**, 13244–13247; (g) M. D. Greenhalgh, A. S. Jones and S. P. Thomas, *ChemCatChem*, 2015, **7**, 190–222; (h) J. Sun and L. Deng, *ACS Catal.*, 2016, **6**, 290–300; (i) J. Guo and Z. Lu, *Angew. Chem., Int. Ed.*, 2016, **55**, 10835–10838.
- (a) J. H. Docherty, J. Peng, A. P. Dominey and S. P. Thomas, *Nat. Chem.*, 2017, **9**, 595–600; (b) I. Buslov, S. C. Keller and X. Hu, *Org. Lett.*, 2016, **18**, 1928–1931; (c) J. Peng, J. H. Docherty, A. P. Dominey and S. P. Thomas, *Chem. Commun.*, 2017, **53**, 4726–4729; (d) J. L. Boyer and A. K. Roy, *US pat.* 0343311 A1, 2014.
- A. J. Challinor, M. Calin, G. S. Nichol, N. B. Carter and S. P. Thomas, *Adv. Synth. Catal.*, 2016, **358**, 2404–2409.
- (a) D. Noda, A. Tahara, Y. Sunada and H. Nagashima, *J. Am. Chem. Soc.*, 2016, **138**, 2480–2483; (b) A. Sanagawa and H. Nagashima, *Organometallics*, 2018, **17**, 2859–2871; (c) C. H. Schuster, T. Diao, I. Pappas and P. J. Chirik, *ACS Catal.*, 2016, **6**, 2632–2636; (d) G. Wu, U. Chakraborty and A. Jacobi von Wangelin, *Chem. Commun.*, 2018, **54**, 12322–12325; (e) H. L. Sang, S. Yu and S. Ge, *Chem. Sci.*, 2018, **9**, 973–978.
- (a) A. J. Cresswell, S. G. Davies, P. M. Roberts and J. E. Thomson, *Chem. Rev.*, 2015, **115**, 566–611; (b) C. A. Wamser, *J. Am. Chem. Soc.*, 1948, **70**, 1209–1215; (c) M. G. Freire, C. M. S. S. Neves, I. M. Manucho, J. A. P. Coutinho and A. M. Fernandes, *J. Phys. Chem. A*, 2010, **114**, 3744–3749; (d) R. Agahi, A. J. Challinor, N. B. Carter and S. P. Thomas, *Org. Lett.*, 2019, **21**, 993–997.



- 10 (a) J. Boyer, R. Corriu, R. Perz, M. Poirier and C. Reye, *Synthesis*, 1981, 7, 558–559; (b) M. Das and D. F. O'Shea, *Tetrahedron*, 2013, 69, 6448–6460.
- 11 J. M. Larsson and K. J. Szabó, *J. Am. Chem. Soc.*, 2013, 135, 443–455.
- 12 C. Wang, W. J. Teo and S. Ge, *ACS Catal.*, 2017, 7, 855–863.
- 13 D. Noda, A. Tahara, Y. Sunada and H. Nagashima, *J. Am. Chem. Soc.*, 2016, 138, 2480–2483.
- 14 J. Y. Wu, B. N. Stanzl and T. Ritter, *J. Am. Chem. Soc.*, 2010, 132, 13214–13216.
- 15 (a) J. G. de Vries and C. J. Elsevier, *The Handbook of Homogeneous Hydrogenation*, Wiley-VCH Verlag GmbH, Weinheim, Germany, 2006; (b) P. Etayo and A. Vidal-Ferran, *Chem. Soc. Rev.*, 2013, 42, 728–754; (c) C. Pettinari, F. Marchetti and D. Martini, in *Comprehensive Coordination Chemistry II*, Elsevier, 2003, pp. 75–139.
- 16 For examples of olefin hydrogenation using iron and cobalt see: (a) P. J. Chirik, *Acc. Chem. Res.*, 2015, 48, 1687–1695; (b) D. Gärtner, A. Welther, B. R. Rad, R. Wolf and A. Jacobi von Wangelin, *Angew. Chem., Int. Ed.*, 2014, 53, 3722–3726; (c) L. J. Murphy, M. J. Ferguson, R. McDonald, M. D. Lumsden and L. Turculet, *Organometallics*, 2018, 37, 4814–4826; (d) A. J. MacNair, M.-M. Tran, J. E. Nelson, G. U. Sloan, A. Ironmonger and S. P. Thomas, *Org. Biomol. Chem.*, 2014, 12, 5082–5088; (e) D. J. Frank, L. Guet, A. Käslin, E. Murphy and S. P. Thomas, *RSC Adv.*, 2013, 3, 25698; (f) T. N. Gieshoff, U. Chakraborty, M. Villa and A. Jacobi von Wangelin, *Angew. Chem., Int. Ed.*, 2017, 56, 3585–3589.
- 17 (a) J. M. Hoyt, V. A. Schmidt, A. M. Tondreau and P. J. Chirik, *Science*, 2013, 775, 772–775; (b) V. A. Schmidt, J. M. Hoyt, G. W. Margulieux and P. J. Chirik, *J. Am. Chem. Soc.*, 2015, 137, 7903–7914.
- 18 (a) J. V. Obligation, M. J. Bezdek and P. J. Chirik, *J. Am. Chem. Soc.*, 2017, 139, 2825–2832; (b) J. V. Obligation and P. J. Chirik, *ACS Catal.*, 2017, 7, 4366–4371; (c) N. G. Léonard, M. J. Bezdek and P. J. Chirik, *Organometallics*, 2017, 36, 142–150; (d) J. V. Obligation, H. Zhong and P. J. Chirik, *Isr. J. Chem.*, 2017, 57, 1032–1036; (e) W. N. Palmer, J. V. Obligation, I. Pappas and P. J. Chirik, *J. Am. Chem. Soc.*, 2016, 138, 766–769.

