PERSPECTIVE

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Tuning steric and electronic effects in transition-metal β-diketiminate complexes†

Chi Chen,[a] Sarina M. Bellows[b] and Patrick L. Holland*[a]

β-Diketiminates are widely used supporting ligands for building a range of metal complexes with different oxidation states, structures, and reactivities. This Perspective summarizes the steric and electronic influences of ligand substituents on these complexes, with an eye toward informing the design of new complexes with optimized properties. The backbone and *N*-aryl substituents can give significant steric effects on structure, reactivity and selectivity of reactions. The electron density on the metal can be tuned by installation of electron withdrawing or donating groups on the β-diketiminate ligand as well. Examples are shown from throughout the transition metal series to demonstrate different types of effects attributable to systematic variation of β-diketiminate ligands.

1. Introduction

The properties and reactions of metal complexes are highly dependent on the choice of supporting ligand, and this choice is one of the keys to successful coordination chemistry. Since its introduction in 1968,[1–3] the β-diketiminate (often called "NacNac" because of its addition of two Nitrogen atoms to the common acac ligand) has gained great popularity as a support-

ing ligand. Unlike acetylacetonate (acac), the β-diketiminate ligand scaffold offers steric protection at the metal center through the choice of *N*-substituents; this makes β-diketiminates less labile and more suitable as spectator ligands. β-Diketiminate ligands are typically synthesized from condensation of a β-diketone and an amine, and chemists have only scratched the surface of the thousands of potential combinations.[4]

N-Aryl β-diketiminate ligands have been most widely used, and they support a variety of metals in many oxidation states. Complexes of *N*-aryl β-diketiminates have shown great reactivity and selectivity for a variety of methodologies,[4,5] including polymerization and functionalization of alkenes and cross-coupling reactions. In addition, late transition metal

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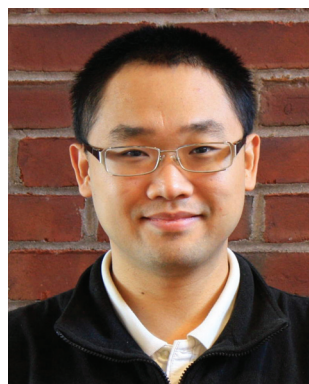
Chi Chen received his Bachelor of Science degree at Peking University in 2009 and did additional research at the University of Texas – Arlington before starting graduate research at the University of Rochester in 2011. In a joint project with Daniel Weix and Patrick Holland, he is developing and studying new β-diketiminate supported cobalt catalysts for alkene transformations such as isomerization and hydrosilylation. In 2013, he moved to Yale University where he is completing his PhD research.

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Sarina Bellows received her Bachelor of Science degree from Syracuse University in 2008, and pursued PhD research at the University of Rochester with Patrick Holland. In her research, she synthesized iron complexes of new β-diketiminate ligands, and also performed computations to explain the mechanisms of their reactions. Since receiving her PhD in 2014, she has been a postdoctoral fellow at Rochester with Thomas Cundari and William Jones through the Center for Enabling New Technologies through Catalysis.
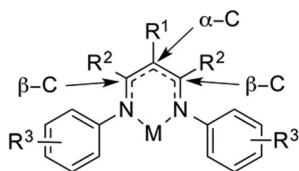
Sarina M. Bellows

Fig. 1 Substituent patterns in β-diketiminate ligands.

β-diketiminate complexes have been used to build low coordinate metal centers, mimicking the active sites of metalloproteins.[6–14] A vast number of ligand variations and different coordination modes have been reported, and some examples are shown in Fig. 1. In this Perspective, the focus will be solely on complexes of the type shown in Fig. 1 with d block transition metals in a η^2 binding mode. We summarize trends from systematic variations in these complexes with examples, though we make no claim that our coverage is complete. This Perspective is intended to serve as a guide to chemists who are interested in tuning the properties of β-diketiminate complexes to achieve their specific goals. We also refer the interested reader to another Perspective by Budzelaar which gives more depth on *N*-aryl β-diketiminate complexes of Ru, Os, Rh, Ir, Pd, and Pt.[15]

2.   Nomenclature

In this Perspective, the ligand abbreviation $^{R1}L^{R2,R3}$ is used to specify the substituents on a β-diketiminate ligand. R^1 refers to the substituent on the central backbone carbon (α-C), R^2 refers to the substituents on the nitrogen-bearing carbon atoms (β-C), and R^3 refers to the substituents on the *N*-aryl group. For the R^3 aryl substituents *meta*- and *para*-substitutions of *N*-aryl are specified as *m*- and *p*-, respectively, while the common *ortho*-substituents are given without the

Patrick Holland completed an AB at Princeton University, and a PhD at UC Berkeley with Richard Andersen and Robert Bergman. In postdoctoral work at Minnesota with William Tolman, he learned to love β-diketiminates through the synthesis of copper complexes. In his independent career, he has explored the use of β-diketiminate complexes of iron, cobalt and nickel, as applied to N₂ reduction, C–H oxidation, redox-active ligands, new bonding environments, and novel reactivity. He was on the faculty at the University of Rochester from 2000–2013, and is now a Professor of Chemistry at Yale University.

Patrick L. Holland

Dipp	2,6-diisopropylphenyl
Tipp	2,4,6-triisopropylphenyl
Dep	2,6-diethylphenyl
Mes	2,4,6-trimethylphenyl
An	1-anthracenyl
ArF	3,5-bis(trifluoromethyl)phenyl
Tbt	2,4,6-tris[bis(trimethylsilyl)methyl]phenyl

Chart 1 Abbreviations used in this Perspective.

o-abbreviation for convenience. Some other abbreviations can be found in Chart 1.

3.   Steric effects on β-diketiminates

The steric demands of β-diketiminate ligands can be tuned by substitution of functional groups on the backbone (β-C) or the *N*-aryl substituents. Typical backbone (β-C) substituents are *tert*-butyl, phenyl, trifluoromethyl and methyl; unsubstituted (β-dialdiminate) ligands are also known. Two approaches can be used to tune the sterics of the *N*-aryl groups: first, to change the size of *ortho*-substituents on the *N*-aryl; or second, to relocate the substituents from *ortho*- position to the *meta*- or *para*-position.

The modification of β-diketiminate steric hindrance can bring changes in the structure and reactivity. The structural differences include changes on the coordination number, bond angles and bond lengths, geometry and conformation of metal complexes. We highlight three types of reactivity differences: different structures of β-diketiminate complexes, different outcomes of stoichiometric reactions of β-diketiminate complexes, and different activity in catalytic reactions.

3.1.   Steric effects on structural properties

Generally, using smaller substituents on the β-C and *N*-aryl, or relocation of the *N*-aryl substituents farther from the metal center, reduces the overall steric coverage of the metal coordination sphere. As a result, dimeric/polymeric metal complexes are more often formed with less sterically hindered β-diketiminate ligands. For example, comparisons with more hindered monomeric analogues were reported for $[LScCl_2]_n$ ($L^{tBu,iPr}$,[16] $n = 1$; $L^{Me,iPr}$,[17] $n = 2$), $[LSc(CH_3)_2]_n$ ($L^{tBu,iPr}$,[16] $n = 1$; $L^{Me,iPr}$,[17] $n = 2$), $[LFeCl]_n$ ($L^{tBu,iPr}$,[18] $n = 1$; $L^{Me,iPr}$,[19] $^{Me}L^{Me,Me}$,[20] $n = 2$), $[LFeF]_n$ ($L^{tBu,iPr}$, $n = 1$; $L^{Me,iPr}$, $n = 2$),[21] $[LCoCl]_n$ ($L^{tBu,iPr}$,[22] $n = 1$; $L^{Me,iPr}$,[23] $n = 2$), $[LNiCl]_n$ ($L^{tBu,iPr}$,[22] $n = 1$; $L^{Me,iPr}$,[24] $^{Me}L^{Me,Me}$,[25] $n = 2$), $[LNi(CO)]_n$, ($L^{tBu,iPr}$,[26] $L^{Me,iPr}$,[27] $n = 1$; $^{Me}L^{Me,Me}$,[28] $n = 2$), $[L^{R,iPr}CuCl]_n$ ($L^{Me,iPr}$,[29] $^{Cl}L^{Me,iPr}$,[29] $n = 1$; $^{Ph}L^{H,iPr}$,[30] $L^{Me,Cl}$[31] $n = 2$), and $[LPd(μ-OAc)]_n$ ($L^{Me,iPr}$,[32] $n = 1$; $L^{Me,H}$,[32] $^{Cl}L^{Me,H}$[33] $n = 2$). The angle between the two β-diketiminate ligand planes

Table 1 Selected examples of steric effects on ligand plane orientation of bimetallic complexes complexes

Complex	Ligand	Dihedral angle between two ligand planes	Ref.
[LV]$_2$	L^{Me,An}	66°	34
	L^{Me,Et}	0°	34
	L^{Me,Me}	0°	34
[LCr(μ-Cl)]$_2$	L^{tBu,iPr}	32°	35
	L^{Me,iPr}	0°	36
	L^{Me,Me}	0°	37
LCr(η^5-Cp)(μ-O)Cr(η^5-Cp)L	L^{Me,Me}	9°	38
	L$^{Me,m\text{-}TIPP}$	17°	38
[LFe(μ-H)]$_2$	L^{tBu,iPr_3}	67°	39
	L^{tBu,iPr}	69°	40
	L^{Me,iPr}	71°	21
	^{Me}L^{Me,Me}	82°	20
LFe(tBuPy)(NN)Fe(tBuPy)L	L^{tBu,iPr}	82°	41
	L^{Me,iPr}	50°	41
LFeNNFeL	L^{tBu,iPr}	87°	6
	L^{Me,iPr}	0°	41
[LFeNNFeL]K$_2$	L^{tBu,iPr}	36°	6
	L^{Me,iPr}	34°	41
LNi(P$_4$)NiL	L^{Me,iPr}	40°	42
	L^{Me,Et}	51°	42
[LCu(μ-Cl)]$_2$	L^{Me,Et}	0°	43
	L^{Me,Cl}	81°	31
	^{Cl}L^{Me,Me}	75°	43
[LCu(μ-OH)]$_2$	L^{CF_3,Me}	60°	44
	L^{Me,Me}	0°	45
	^{CN}L^{H,Et}	0°	46
	^{CN}L^{H,Me_3}	11°	43
	^{NO_2}L^{H,Me_3}	41°	30

Table 2 Selected examples of steric effect on distance of metal to ligand plane

Complex	Ligand	Distance from M to ligand plane (Å)	Ref.
LScCl$_2$(THF)$_n$	L^{tBu,iPr}	1.295	16
	L^{Me,iPr}	0.694	47
LSc(alkyl)$_2$	L^{tBu,iPr}	1.154	16
	L^{Me,iPr}	1.116	16
	L$^{Me,m\text{-}tBu}$	0.489	69
	L$^{Me,m\text{-}Tipp}$	0.204	69
LZrCl$_3$	L^{tBu,iPr}	1.650	70
	L^{Me,iPr}	0.820	71
LVCl$_2$(THF)$_n$	L^{Me,Me}	0.528	52
	L^{Me,H}	0.227	55
LCr(Cp)(Me)	L^{Me,iPr}	0.702	62
	L^{Me,Et}	0.699	72
	L^{Me,Me}	0.650	72
LCr(Cp)(Cl)	L^{Me,iPr}	0.719	62
	L^{Me,Et}	0.751	72
	L^{Me,Me}	0.680	63
	L^{Me,H}	0.087	73
LCr(Cp)(μ-O)Cr(Cp)L	L^{Me,Me}	0.858	38
		0.848	
	L$^{Me,m\text{-}TIPP}$	0.771	38
		0.726	
[LCr(μ-Cl)(THF)]$_2$	L^{Me,iPr}	0.668	36
	L^{Me,Me}	0.554	37
[LFe(μ-H)]$_2$	L^{tBu,iPr}	0.565	40
	L^{Me,iPr}	0.540	21
	^{Me}L^{Me,Me}	0.260	20
LFe(μ-H)$_2$B(Et)$_2$	L^{tBu,iPr}	0.093	74
	L^{Me,iPr}	0.000	74
LFe(μ-Cl)$_2$Li(THF)$_2$	L^{Me,iPr}	0.381	18
	L^{Me,Me_3}	0.000	20
LFe(F)(tBuPy)	L^{tBu,iPr}	0.339	21
	L^{Me,iPr}	0.294	21
LFe(tBuPy)(NN)Fe(tBuPy)L	L^{tBu,iPr}	0.394	41
		0.553	
	L^{Me,iPr}	0.250	41
		0.250	
LFe-(η^3-N$_3$Ad)	L^{tBu,iPr}	0.762	75
	L^{Me,iPr}	0.753	75
[LFeNNFeL]K$_2$	L^{tBu,iPr}	0.290	6
		0.111	
	L^{Me,iPr}	0.072	41
		0.004	
LFe–alkyl	L^{tBu,iPr}	0.065	76
	L^{Me,iPr}	0.019	77
LFe–alkyne	L^{tBu,iPr}	0.097	78
	L^{Me,iPr}	0.008	79
LCo(μ-Cl)$_2$Li(THF)$_2$	L^{tBu,iPr}	0.362	22
	L^{Me,iPr}	0.314	80
LNi(P$_4$)NiL	L^{Me,iPr}	0.184	42
		0.184	
	L^{Me,Et}	0.215	42
		0.030	
LCu(CNAr)	L^{Me,iPr}	0.342	29
	L^{Me,Me}	0.144	81
[LCu(μ-S)]$_2$	^{Ph}L^{H,iPr}	0.349	67
	^{Ph}L^{H,Et}	0.302	67
	^{ArF}L^{H,iPr}	0.271	67
	^{ArF}L^{H,Me}	0.002	67
LCu(NCCH$_3$)	L^{tBu,iPr}	0.046	8
	L^{CF_3,iPr}	0.028	82
	L$^{CF_3/Me,iPr}$	0.022	82
LRu(Cl)(η^6-Benzene)	L^{CF_3,Me}	0.624	83
	L^{Me,Me}	0.635	84
	L$^{CF_3,m\text{-}CF_3}$	0.246	83
	L$^{Me,m\text{-}Me}$	0.207	85
	L^{Me,H}	0.048	83
LRu(Cl)(η^5-Cp*)	L^{Me,Me}	0.628	86
	L$^{Me,m\text{-}Me}$	0.343	86

in dimeric metal complexes is often influenced by the different substituents on the ligand (Table 1). However, there is no clear correlation between the substituent size and the angle, indicating that this angle is dependent on the bonding at the metal as well as steric interactions between the ligands on the two sides.

One trend that emerges is that higher coordination numbers can be achieved with smaller β-diketiminate supporting ligands. For example, more solvent molecules (THF, arene, *etc.*) and neutral ligands (CO, PPh$_3$, *etc.*) can be coordinated to a metal center with less sterically hindered β-diketiminate in LScCl$_2$(THF)$_n$ (L^{tBu,iPr},[16] $n = 0$; L^{Me,iPr},[47] $n = 1$), LSc(CH$_3$)$_2$(THF)$_n$ (L^{tBu,iPr},[16] $n = 0$; L^{Me,iPr},[16] $n = 1$), LSc(Cl)(NHAr)(THF)$_n$ (L^{tBu,iPr},[48] $n = 0$; L^{Me,iPr},[49] $n = 1$), [LSc(CH$_3$)(arene)$_n$]$^+$ (L^{tBu,iPr},[50] $n = 0$; L^{Me,iPr},[50] $n = 1$), LTiCl$_2$(THF)$_n$ (L^{tBu,iPr},[51] L^{tBu,Me_3},[52] L$^{Me,Tbt/Me_3}$,[53] $n = 0$; L^{Me,iPr},[54] $n = 1$; L^{Me,H},[55] $n = 2$), LVCl$_2$(THF)$_n$ (L^{Me,iPr},[52,56] L^{Me,Et},[34] L^{Me,Me_3},[34] L^{Ph,iPr},[34] $n = 0$; L$^{Me, H}$,[55] $n = 2$), [LCr(μ-Cl)(Solvent)$_n$]$_2$ (L^{tBu,iPr},[35] $n = 0$; L^{Me,iPr},[36] L^{Me,Me},[37] $n = 1$; Solvent = THF, benzene), LFe(NHdipp)(THF)$_n$ (L^{tBu,iPr},[19] $n = 0$; L^{Me,iPr},[28] $n = 1$), and LCu(PPh$_3$)$_n$ (^{Ph}L^{H,iPr},[57] L^{Me,Me},[58] L^{Me,iPr},[59] L^{Me,Me_3},[60] $n = 1$; ^{Ph}L^{H,Me},[57] L$^{CF_3,m\text{-}CF_3}$,[61] $n = 2$). Steric conflict between *N*-aryl substituents and metal can also push the metal center out of the β-diketiminate ligand plane in some metal complexes, especially for early transition metals (Table 2). However, exceptions can be found in L^{R,Mes}TiCl$_2$,[52] L^{Me,R}Cr(η^5-Cp),[62,63] L^{R,iPr}FeNNFeL,[6,41] [L^{Me,R}Ni(μ-Cl)]$_2$,[24,25]

Table 3 Steric effects of backbone (β-C) substituents on structural properties

Complex	Ligand	N–M–N bite angle	C(aryl)–N–C(β) bond angle	M–N distance (Å)	Ref.
LScCl₂(THF)ₙ	$L^{tBu,iPr}$	95.9°	125.3° / 126.9°	2.046 / 2.099	16
	$L^{Me,iPr}$	86.8°	116.9° / 117.8°	2.107 / 2.175	47
LSc(alkyl)₂	$L^{tBu,iPr}$	93.5°	125.5° / 126.2°	2.091 / 2.144	16
	$L^{Me,iPr}$	90.7°	120.1° / 120.8°	2.113 / 2.133	16
LFe(μ-H)₂BEt₂	$L^{tBu,iPr}$	97.35°	127.80° / 129.28°	1.971 / 1.969	74
	$L^{Me,iPr}$	95.91°	120.58°	1.971	74
LFeX	$L^{tBu,iPr}$	96.35°	128.39°	1.946	18
	$L^{Me,iPr}$	94.50°	116.61° / 116.72°	2.002 / 2.006	19
LFe(F)(tBuPy)	$L^{tBu,iPr}$	97.80°	124.80° / 126.43°	2.015 / 2.007	21
	$L^{Me,iPr}$	95.00°	118.38° / 119.53°	2.012 / 2.009	21
LFe(tBuPy)-(NN)Fe-(tBuPy)L	$L^{tBu,iPr}$	99.23°	123.02° / 124.13°	2.005	41
		97.33°	124.22° / 124.76°	2.000	
	$L^{Me,iPr}$	95.86°	118.59° / 119.99°	2.005 / 1.993	41
LFe(N₃Ad)	$L^{tBu,iPr}$	98.84°	123.88° / 123.39°	2.043 / 2.018	75
	$L^{Me,iPr}$	97.95°	118.34° / 117.40°	2.021 / 2.016	75
LFeNNFeL	$L^{tBu,iPr}$	96.01°	129.11° / 127.00°	1.965 / 1.970	6
	$L^{Me,iPr}$	94.78°	121.57° / 118.66°	1.945 / 1.984	41
LFeiPr	$L^{tBu,iPr}$	94.25°	126.33° / 128.11°	1.990 / 1.989	76
	$L^{Me,iPr}$	92.78°	119.84° / 120.60°	1.983 / 1.983	77
LFe-(η²-PhC≡CH)	$L^{tBu,iPr}$	96.16°	123.65° / 124.62°	1.975 / 2.005	78
	$L^{Me,iPr}$	93.67°	119.31° / 118.57°	1.973 / 1.990	79
LCo(μ-Cl)₂Li-(THF)₂	$L^{tBu,iPr}$	99.42°	124.78° / 125.81°	1.968 / 1.961	22
	$L^{Me,iPr}$	98.19°	120.23° / 120.38°	1.957 / 1.962	80
LCo(alkyl)	$L^{tBu,iPr}$	97.68°	127.59° / 125.04°	1.960 / 1.950	88
	$L^{Me,iPr}$	95.60°	119.70° / 118.82°	1.948 / 1.946	89
LNi(CO)	$L^{tBu,iPr}$	98.85°	126.33° / 129.40°	1.924 / 1.856	26
	$L^{Me,iPr}$	96.41°	119.89° / 122.58°	1.917 / 1.868	27
LCu(η²-OAc)	$^{CN}L^{Me,iPr}$	96.63°	119.68° / 120.45°	1.905 / 1.914	90
	$^{CN}L^{H,iPr}$	94.79°	116.9° / 116.9°	1.944 / 1.944	46
[LCu(μ-OH)]₂	$L^{CF_3,Me}$	95.28°	122.69° / 122.87°	1.940 / 1.943	44
	$L^{Me,Me}$	94.83°	117.36° / 117.61°	1.937 / 1.945	45
LCu(NCCH₃)	$L^{tBu,iPr}$	102.33°	128.75° / 127.68°	1.936 / 1.931	8
	$L^{CF_3,iPr}$	98.98°	124.74° / 125.00°	1.940 / 1.935	68
	$L^{Me,iPr}$	98.98°	118.94° / 119.21°	1.940 / 1.942	8
	$^{Ph}L^{H,iPr}$	97.25°	118.46° / 116.59°	1.964 / 1.950	8

Table 3 *(Contd.)*

Complex	Ligand	N–M–N bite angle	C(aryl)–N–C(β) bond angle	M–N distance (Å)	Ref.
LRu(Cl)-(η⁵-Cp*)	$L^{CF_3,m\text{-}Me}$	90.18°	118.55° / 118.42°	2.069 / 2.055	86
	$L^{Me,m\text{-}Me}$	87.83°	116.43° / 115.98°	2.050 / 2.051	86
	$L^{CF_3,m\text{-}CF_3}$	89.67°	117.47° / 118.21°	2.070 / 2.071	86
	$L^{Me,m\text{-}CF_3}$	87.99°	114.91° / 115.46°	2.071 / 2.071	86
LRu(η⁵-Cp*)	$L^{CF_3,m\text{-}Me}$	90.08°	116.95° / 117.42°	2.050 / 2.050	86
	$L^{Me,m\text{-}Me}$	87.92°	115.62° / 115.29°	2.060 / 2.063	86
	$L^{CF_3,m\text{-}CF_3}$	89.55°	116.09° / 116.53°	2.055 / 2.056	86
	$L^{Me,m\text{-}CF_3}$	87.37°	114.08° / 114.07°	2.045 / 2.040	86

$L^{Me,R}Cu(OAc)$,[64,65] $[LCu(μ-OH)]_2$,[44–46] $[LCu(μ-S)]_2$,[66,67] and $L^{R,iPr}Cu(CO)$.[68]

When the backbone (β-C) substituent size increases (H < Me < CF₃ < tBu, Ph), the steric conflict between backbone (β-C) substituents and *N*-aryl groups escalates, pushing the *N*-aryl rings closer to the metal and forcing them into a more rigid configuration. As a consequence of this "buttressing effect", the metal center often moves deeper into the β-diketiminate binding pocket. This brings three changes to the structure: it typically increases the N–M–N bite angle, increases the C(aryl)–N–C(β) bond angle, and shortens the N–M bond length (see Table 3). Bulky substituents on the *N*-aryl may also affect the bonding to other ligands (see Table 4). Exceptions to this trend, however, are seen with $LTiCl_2$,[52] $LZrCl_3$,[70,87] $[LCr(μ-Cl)]_2$, and $K_2[LFeNNFeL]$,[6,41] due to cation coordination or conformational changes at the metal center. The distances from the metal to the non-diketiminate co-ligand can also be affected by the backbone substituents (see ESI† for details).

The choice of *N*-aryl substituent has a smaller influence on the bite angle, C(aryl)–N–C(β) bond angle and N–M bond length in most cases. However, changing *N*-aryl substituents can build up steric bulk above and below the N–M–N plane, which can significantly influence the distance from the metal to the other ligands. In general, more hindered *N*-aryl substituents lead to a longer M–L bond (Table 4).

Other modifications of β-diketiminate ligands, including installation of functional groups on the backbone α-C, or on the *para*-position of the *N*-aryl substituents, have little influence on the core structural parameters of β-diketiminate metal complexes.

The geometry and conformation of metal complexes can also be changed with modification of the supporting β-diketiminate ligand. The zirconium center in $L^{Me,R}Zr(CH_2Ph)_3$ (R = iPr, *p*-Me)[94] adopts a square pyramidal geometry with a crystallographic mirror plane passing through it. However, the relative orientation of the ligand planes shows differences (Fig. 2). Without *ortho*-substitution on *N*-aryl, the β-diketiminate ligand plane in $L^{Me,pMe}Zr(CH_2Ph)_3$ forms an angle of 67.7(3)° with the

Table 4 Steric effects of *N*-aryl substituents on structural properties

Complex	Ligand	N–M–N bite angle	M–N distance (Å)	C(aryl)–N–C(β) bond angle	Selected bond length (Å)	Ref.
LSc(CH$_2$TMS)$_2$	L^{Me,iPr}	90.7°	2.113	120.1°	Sc–C: 2.244	16
			2.133	120.8°	2.194	
	L$^{Me,m\text{-}^tBu}$	83.1°	2.128	121.6°	Sc–C: 2.210	69
			2.128	122.1°	2.215	
	L$^{Me,m\text{-}Tipp}$	84.9°	2.127	120.4°	Sc–C: 2.203	69
			2.123	119.2°	2.202	
[LV]$_2$	L^{Me,Et}	88.69°	2.066	115.84°	V–arene: 1.422	34
			2.041	114.05°		
	L^{Me,Me}	88.73°	2.057	115.98°	V–arene: 1.411	34
			2.034	113.22°		
	L^{Me,An}	88.83°	2.025	117.05°	V–arene: 1.744	34
			2.020	117.01°		
LCr(Cl)(η^5-Cp)	L^{Me,iPr}	89.9°	2.036	117.3°	Cr–Cp: 1.929	62
			2.036	117.3°		
	L^{Me,Et}	90.3°	2.022	118.0°	Cr–Cp: 1.901	72
			2.016	117.9°		
	L^{Me,Me}	90.5°	2.019	117.7°	Cr–Cp: 1.897	63
			2.018	119.0°		
LCr(Cp)(alkyl)	L^{Me,iPr}	90.7°	2.039	118.3°	Cr–Cp: 1.972	62
			2.039	118.8°		
	L^{Me,Et}	90.2°	2.029	118.7°	Cr–Cp: 1.963	72
			2.017	118.3°		
	L^{Me,Me}	90.7°	2.024	116.9°	Cr–Cp: 1.966	72
			2.026	117.6°		
LFe(μ-Cl)$_2$Li(THF)$_2$	L^{Me,iPr}	93.22°	2.021	120.27°	Fe–Cl: 2.338	18
			2.006	118.59°	2.324	
	^{Me}L^{Me,Me}	93.19°	1.983	119.19°	Fe–Cl: 2.325	91
			1.983	119.19°	2.325	
[LNi(μ-Cl)]$_2$	L^{Me,iPr}	93.66°	1.946	117.11°	Ni–Cl: 2.350	24
			1.938	116.42°	2.325	
	L^{Me,Me}	94.7°	1.915	117.88°	Ni–Cl: 2.313	25
			1.913	117.30°	2.300	
LNi(μ-P$_4$)NiL	L^{Me,iPr}	94.98°	1.947	117.74°	Ni–P: 2.339	42
			1.968	116.94°	2.217, 2.195	
	L^{Me,Et}	96.44°	1.931	119.86°	Ni–P: 2.203	42
			1.928	115.87°	2.329, 2.167	
[LCu(μ-S)]$_2$	L^{Me,Et}	99.30°	1.907	118.43°	Cu–S: 2.197	66
			1.910	118.18°	2.193	
	L^{Me,Me}	99.43°	1.899	119.65°	Cu–S: 2.184	67
			1.896	119.17°	2.187	
	^{Ph}L^{H,iPr}	96.95°	1.913	116.70°	Cu–S: 2.205	67
			1.905	115.97°	2.198	
	^{Ph}L^{H,Et}	96.92°	1.911	116.96°	Cu–S: 2.195	67
			1.909	117.21°	2.194	
	^{ArF}L^{H,iPr}	97.07°	1.921	115.47°	Cu–S: 2.194	67
			1.905	116.00°	2.206	
	^{ArF}L^{H,Me}	98.07°	1.906	115.21°	Cu–S: 2.198	67
			1.912	117.26°	2.198	
[LCu(μ-OH)]$_2$	^{CN}L^{H,Et}	93.63°	1.955	115.90°	Cu–O: 1.926	46
			1.943	115.44°	1.926, 1.909	
	^{CN}L^{H,Me_3}	93.35°	1.962	117.62°	Cu–O: 1.922	46
			1.958	117.29°	1.920, 1.904	
			1.946			
LRu(Cl)(η^6-Benzene)	L^{Me,Me}	86.56°	2.099	116.80°	Ru–Cl: 2.521	84
			2.099	116.80°	Ru–benzene: 1.688	
	L$^{Me,m\text{-}Me}$	88.21°	2.098	117.53°	Ru–Cl: 2.453	85
			2.091	117.38°	Ru–benzene: 1.683	
LRu(Cl)(η^5-Cp*)	L^{Me,Me}	87.51°	2.089	114.98°	Ru–Cl: 2.461	86
			2.075	115.14°	Ru–Cp*: 1.889	
	L$^{Me,m\text{-}Me}$	87.83°	2.050	116.43°	Ru–Cl: 2.451	86
			2.051	115.98°	Ru–Cp*: 1.869	
LRu(η^5-Cp*)	L^{Me,Me}	87.23°	2.070	114.36°	Ru–Cp*: 1.819	86
			2.060	113.70°		
	L$^{Me,m\text{-}Me}$	87.92°	2.060	115.62°	Ru–Cp*: 1.809	86
			2.063	115.29°		
	L^{Me,H}	87.68°	2.053	113.89°	Ru–Cp*: 1.800	92
			2.046	113.74°		

Table 4 (Contd.)

Complex	Ligand	N–M–N bite angle	M–N distance (Å)	C(aryl)–N–C(β) bond angle	Selected bond length (Å)	Ref.
[LPd(μ-Cl)]₂	$L^{Me,iPr}$	91.78°	2.023	118.65°	Pd–Cl: 2.366	93
			2.013	117.87°	2.354	
	$L^{Me,m\text{-}CF_3}$	90.93°	2.006	118.57°	Pd–Cl: 2.350	93
			1.989	118.97°	2.352	
	$L^{Me,H}$	91.30°	2.000	118.20°	Pd–Cl: 2.342	33
			2.001	120.61°	2.356	
LPd(Cl)(Py)	$L^{Me,iPr}$	91.70°	2.031	118.19°	Pd–Cl: 2.315	93
			2.014	116.65°	Pd–Py: 2.078	
	$L^{Me,m\text{-}CF_3}$	90.08°	2.026	119.46°	Pd–Cl: 2.302	93
			2.013	120.11°	Pd–Py: 2.039	
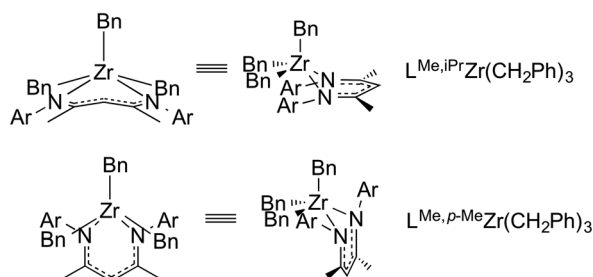
Fig. 2 Structural influence of sterically different aryl groups on the conformation of Zr complexes.
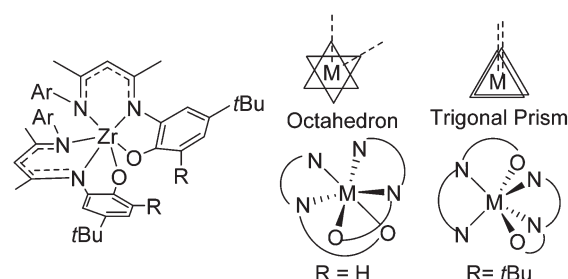
Fig. 3 Structural differences between bis(ligand) complexes on zirconium, with different ortho substituents.

least squares plane defined by C(Bn)–C(Bn)–N–N. In contrast, the angle between the ligand planes in $L^{Me,Me}Zr(CH_2Ph)_3$ is only 7.0(3)°. Presumably, this difference is due to steric conflict between the benzyl and N-aryl substituents. N-Aryloxy-β-diketiminate zirconium complexes also showed a different orientation depending on steric bulk (Scheme 1).[95] Bridged aryloxides were observed with one meta-tBu on the N-aryl, but the presence of a second meta-tBu group gave steric conflict

that resulted in the isolation of a dimer with bridging chlorides instead. In the same system, the L₂Zr complexes also showed conformational differences where the bulkier ligand adopted a trigonal prismatic geometry (Fig. 3).

The solution structure of the metal complex can be affected by different steric bulk as well. For example, two sets of peaks were observed in ¹H NMR and ¹²⁵Te NMR spectra of $L^{tBu,iPr}Sc$-$(TeCH_2TMS)_2$,[96] suggesting exo and endo tellurolates that are static on the NMR time scale. In contrast, the two tellurolate
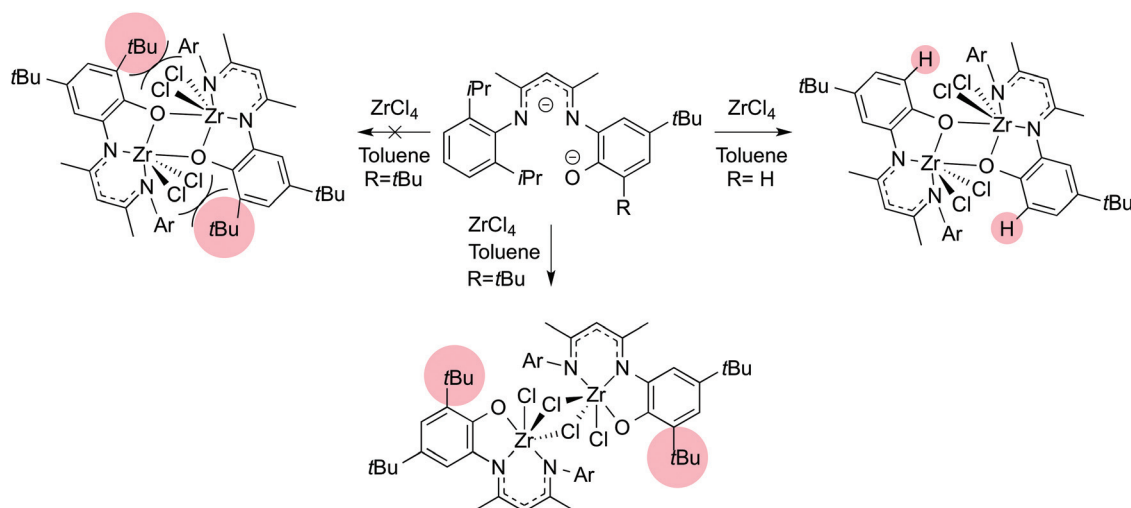
Scheme 1

groups are equivalent for $L^{Me,iPr}Sc(TeCH_2TMS)_2$,[96] indicating rapid *endo/exo* flipping. Thus, larger groups create more difficulty for $Sc(TeR)_2$ to flip through the channel restricted by the *N*-aryl groups. In another example, ^1H NMR peaks of a molybdenum imido alkylidene supported by $L^{Me,m-Me}$ was broadened compared with that of its $L^{Me,Me}$ analogue, suggesting the relatively free rotation of *N*-aryl in the less sterically hindered *meta*-substituted ligand.

3.2. Steric effects on reactivity and product formation

Here, we highlight other cases where different choices of steric bulk of the supporting β-diketiminate ligand give structurally different products under the same reaction conditions. In general, bulkier groups restrict the available conformations. For example, treatment of $L^{tBu,iPr}ScCl_2$ or $[L^{Me,iPr}ScCl(\mu-Cl)]_2$ with LiNHtBu in hexanes generated different products (Scheme 2).[48,49] The authors proposed that the less sterically hindered $L^{Me,iPr}$ allows the formation of a dimeric transition state that is necessary for ligand exchange and disproportionation.

Extrusion of $Te(CH_2TMS)_2$ from $L^{R,iPr}Sc(TeCH_2TMS)_2$ (R = tBu, Me) under photolysis formed different products depending on R (Scheme 3).[96] Crossover between $(LSc(TeCH_2SiMe_3)_2$ and $LSc(TeCH_2CMe_3)_2)$ showed that the product came from a

bimolecular process. It is likely that the tellurolate–telluride $(LSc(TeCH_2TMS))_2(\mu-Te)$ is an intermediate on the way to the bridging telluride complex. However, the greater steric bulk of $L^{tBu,iPr}$ stabilized the tellurolate–telluride species, preventing the loss of a second molecule of $Te(CH_2TMS)_2$.

Reduction of $L^{Me,R}VCl_2$ (R = Me, Et, anthracenyl) with 2 equivalents of KC_8 in THF gave dimeric vanadium(I) complexes, while reaction of $L^{Ph,iPr}VCl_2$ gave extrusion of the imido fragment from diketiminate under the same conditions (Scheme 4).[34] This was not only from having an available arene for binding, because reduction of $L^{Me,iPr}VCl_2$ in toluene gave an inverted sandwich complex. Rather, the authors surmised that the steric conflict between *N*-aryl and backbone phenyl group twisted the *N*-aryl group, destabilizing the LV intermediate and bringing about the reductive C–N bond cleavage of the ligand.

In another example, oxidation of a chromium(II) complex gave a highly reactive chromium oxo complex. However, the attempt to generate a chromium oxo complex gave different products depending on the steric bulk of different β-diketiminate ligands (Scheme 5).[38] Reaction of $L^{Me,Me}CrCp$ or $L^{Me,m-TIPP}CrCp$ with pyridine *N*-oxide gave a μ-oxo dimer, while the bulkier $L^{Me,Et}Cr–Cp$ generated a product from hydrogen atom transfer. The sterically more hindered *ortho*-ethyl substituents may prevent the μ-oxo dimer from forming, and rather the highly reactive terminal oxo ($L^{Me,Et}(Cp)Cr=O$) can abstract a hydrogen atom from its own ligand, ultimately generating a new C–C bond.

Upon addition of O_2, copper(I) complexes supported by different β-diketiminate ligands form different products (Scheme 6). More sterically hindered $L^{tBu,iPr}Cu(NCCH_3)$ and $L^{Me,iPr}Cu(NCCH_3)$ formed a copper(II) peroxo $LCu(O_2)$ while less bulky $^{R'}L^{H,R}Cu$ (R = iPr, Me, Et; R' = H, Ph) complexes gave a bis(μ-oxo)dicopper(III) complex.[8,30] These reactivity differences between the two systems were attributed to the steric effect of the backbone (β-C) substituents, which rigidify the *N*-aryl substituents and prevent the dimer from forming.

The dinitrogen ligand in $L^{R,iPr}FeNNFeL^{R,iPr}$ (R = tBu, Me) can be replaced by other neutral ligands like carbon monoxide or isocyanide.[41] When exposing with excess CO, $L^{Me,iPr}FeNN$-FeL converted to square pyramidal $L^{Me,iPr}Fe(CO)_3$, while the
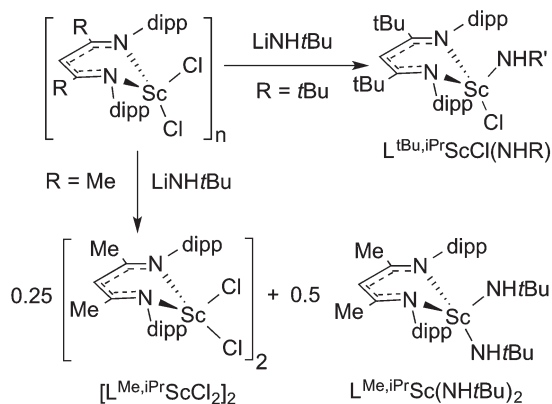
Scheme 2
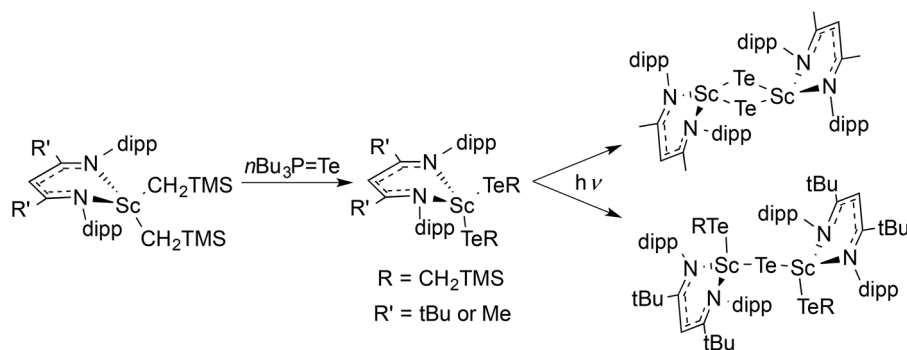
Scheme 3
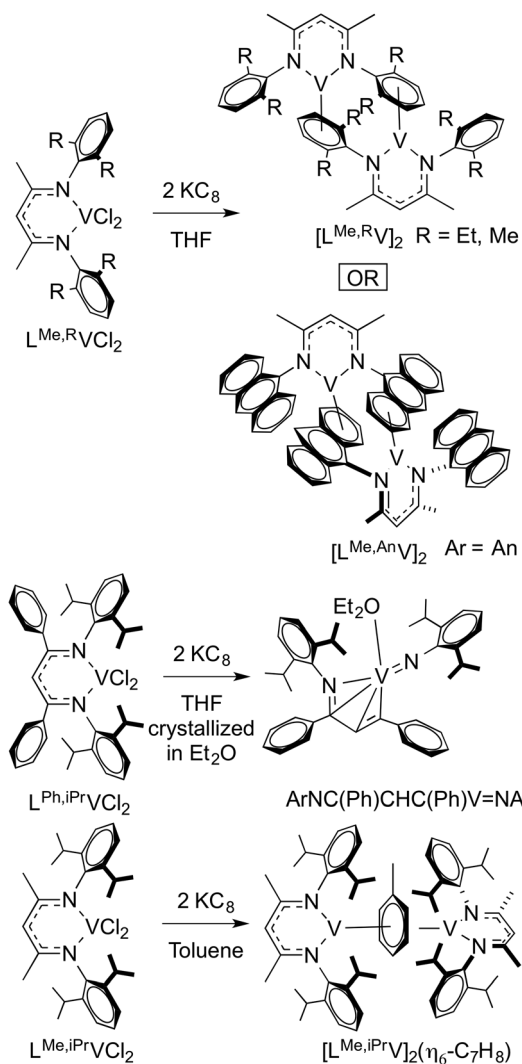
Scheme 4

Scheme 5

Scheme 6

L^{tBu,iPr} analogue gave a mixture of L^{tBu,iPr}Fe(CO)$_3$ and L^{tBu,iPr}Fe-(CO)$_2$. Since the two N-dipp substituents are closer in L^{tBu,iPr}, binding the third axial CO may bring steric tension between iPr and CO, which explains the formation of square planar L^{tBu,iPr}Fe(CO)$_2$. Similarly, N$_2$ exchange in L^{R,iPr}FeNNFeL^{R,iPr} is much more rapid with R = Me than R = ^{t}Bu, implying that transient species with axial N$_2$ are also accessible but only with the smaller R = Me.[41] In a more deep-seated difference in reactivity, attempts to make analogous ^{Me}L^{Me,Me}FeNNFe^{Me}L^{Me,Me} complexes gave N$_2$ cleavage to a tetra-iron bis(nitride) complex, with complete cleavage of the N–N bond (Scheme 7).[20] The authors proposed that the smaller supporting ligand allows access to an intermediate in which three LFe units can interact simultaneously with the same molecule of N$_2$.

3.3.   Steric effect on activity of metal complexes

Varying the steric bulk of the β-diketiminate ligand has a significant effect on activity of metal complexes in both stoichio-metric and catalytic reactions. In most cases, a more sterically hindered β-diketiminate ligand builds up steric tension in transition states or intermediates, which raises the activation barrier and slows the reaction rates. However, the added steric bulk has advantages because it can enable the isolation of transient intermediates.
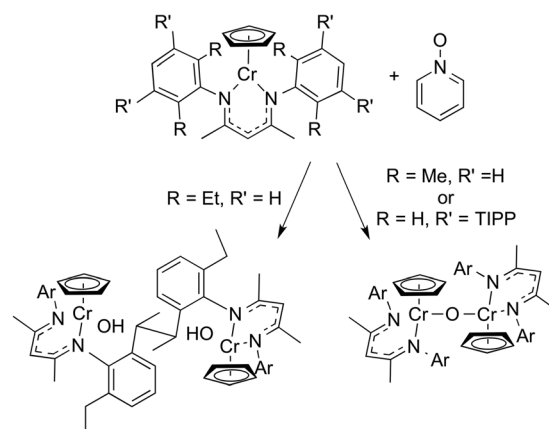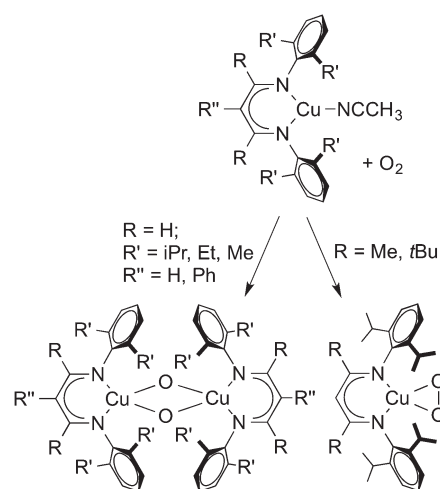
The single-electron oxidative addition of organic halides to chromium(II) complexes (Scheme 8) illustrated the steric effect of *ortho*-substituents on the N-aryl group.[62,72,97] The less hindered asymmetric L$^{Me,iPr/p-Y}$Cr(Cp) gave a rate constant of 0.5–1.0 M^{-1} s^{-1} (depending on the electronic properties of Y; see section 4.2 below),[97] whereas L^{Me,iPr}Cr(Cp) and its L^{Me,Me}, L^{Me,Mes}, and L^{Me,Et} analogues gave rate constants that were more than an order of magnitude smaller, ranging from 0.02–0.03 M^{-1} s^{-1}.[72] Thus, removing the *ortho*-alkyl groups from one of the N-aryl groups greatly enhanced the reactivity of chromium(II) by increasing the accessibility of methyl iodide.

Catalytic 1-hexene isomerization and dimerization was reported with [L^{Me,R}NiBr]$_2$ (R = iPr, Me), where the less steri-

Scheme 7

Scheme 8
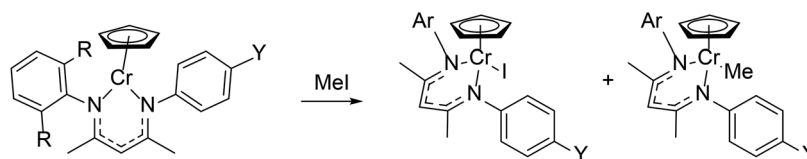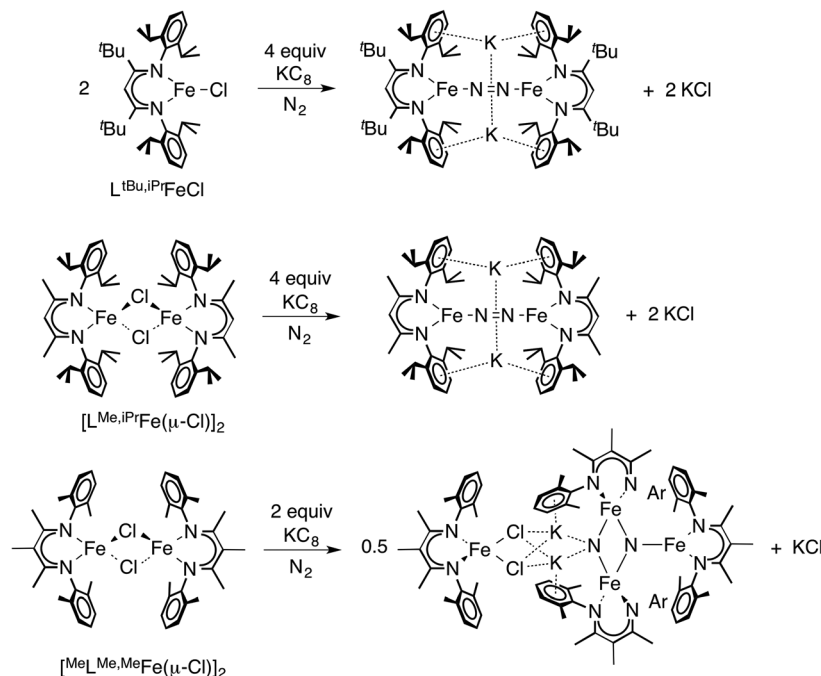
cally hindered $[L^{Me,Me}NiBr]_2$ gave higher conversions under the same conditions.[98] The authors proposed that a β-diketiminate nickel hydride complex was the active catalyst, which would proceed through insertion, β-hydride elimination and chain walking to generate internal alkenes. This makes sense if β-hydride elimination is the rate-limiting step, because larger β-diketiminate substituents would prevent the increase in coordination number. In a demonstration of this idea in a stoichiometric reaction, $L^{tBu,iPr}Fe$-tBu isomerized to $L^{tBu,iPr}Fe$-CH_2iBu only at elevated temperatures, while $L^{Me,iPr}Fe$-tBu isomerized at room temperature to $L^{Me,iPr}Fe$-CH_2iBu (Scheme 9).[76]

The mechanism of alkyne insertion was also studied in detail with isolated β-diketiminate iron hydride complexes. The rate of alkyne insertion was first order in [FeH] and zero order in [alkyne], with $k_{obs} = 1.7(2) \times 10^{-3}$ s^{-1} for $[L^{Me,iPr}FeH]_2$[99] and $5.0(5) \times 10^{-4}$ s^{-1} for $[L^{tBu,iPr}FeH]_2$;[40] again the less hindered complex had higher reactivity. In a related B–C bond cleavage reaction, two mechanisms were proposed: the less hindered iron complex undergoes single iron-hydride opening followed by insertion, while the more hindered $L^{tBu,iPr}$ system can completely dissociate to a reactive monomer.[74]

β-Diketiminate iron imido complexes are prone to hydrogen atom transfer (HAT) from the *ortho* isopropyl substituents of the supporting ligand. To solve the problem, $L^{Me,Ph_3}Fe = NR$ was prepared.[100] The second-order rate constants for hydrogen atom transfer to LFe = NAd from 1,4-cyclohexadiene in C_6D_6 were $2.0(2) \times 10^{-2}$ M^{-1} s^{-1} for $L^{Me,Ph_3}Fe = NAd$, $1.4(2) \times 10^{-4}$ M^{-1} s^{-1} for $L^{Me,iPr}Fe = NAd$ and ~0 for $L^{tBu,iPr}Fe = NAd$ (Scheme 10). Clearly the most bulky $L^{tBu,iPr}Fe = NAd$ gave the slowest HAT reactivity. However, the relative sizes of $L^{Me,iPr}$ and L^{Me,Ph_3} were not obvious. The authors measured the size using the G parameter, which estimates the fraction of the metal overshadowed by the ligand.[101] The results indicated very similar G parameter for $L^{Me,iPr}Fe = NAd$ ($G = 63.8\%$) over $L^{Me,Ph_3}Fe = NAd$ ($G = 62.2\%$), but different shapes (Fig. 4). The different orientation of *N*-aryl with respect to the ligand backbone shows more opening above the imido nitrogen, which results in a larger binding pocket for hydrocarbon substrates (Fig. 5).

Increasing the steric bulk of the β-diketiminate can also prevent formation of certain metal complexes due to steric blocking. In an example, β-diketiminate zirconium tribenzyl complex ($L^{Me,p-Me}Zr(CH_2Ph)_3$) can be synthesized through

Scheme 9

Scheme 10

Fig. 4 Differences in ligand coverage in $L^{Me,iPr}$ vs. L^{Me,Ph_3} in iron(III) imido complexes. The G parameter quantifies the ligand coverage, as described in ref. 100. Thus, even though the overall coverage is similar between the two ligands, the shape of the coverage is different.
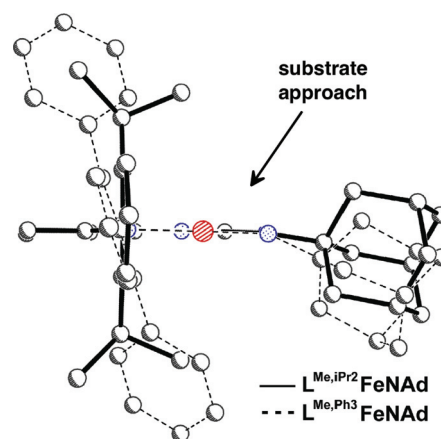

Fig. 5 Side view of the complexes in Fig. 4, showing the greater access to the Fe=N bond when using L^{Me,Ph_3}.

alkane elimination between tetra-alkyl zirconium(IV) and β-diketimines. For its bulkier analogue $L^{Me,iPr}Zr(CH_2Ph)_3$, sterically hindered iPr groups prevent $Zr(CH_2Ph)_4$ from accessing the β-diketiminate binding pocket. Therefore, it was necessary to develop a different synthetic method for $L^{Me,iPr}Zr(CH_2Ph)_3$ involving salt metathesis of LLi and $ZrCl_4$ followed by alkylation (Scheme 11).[94] In another example, $L^{Me,iPr}FeNNFeL^{Me,iPr}$ releases the labile dinitrogen ligand immediately in aromatic solvents forming $L^{Me,iPr}Fe(\eta^6\text{-}C_6H_6)$. However, the more sterically hindered $L^{tBu,iPr}FeNNFeL^{tBu,iPr}$ retains its structure in C_6H_6 up to 100 °C, without coordination of benzene.[41]

However, more sterically hindered metal complexes are favored in some cases because a sterically crowded environment can facilitate intramolecular reactions or increase the concentration of key unsaturated species. An example comes

Scheme 11

R' = tBu, Me; R" = H, Me, Ph, TMS, tBu
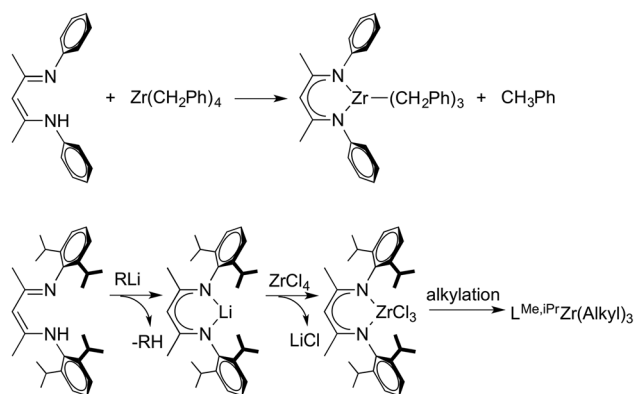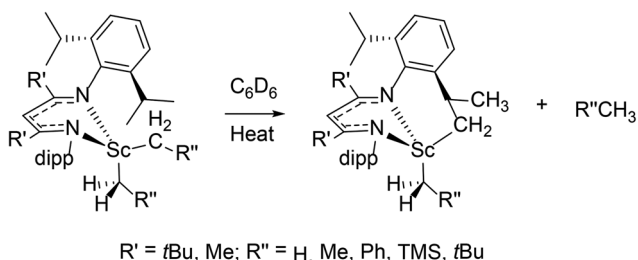
Scheme 12

Scheme 13

in reactions where metalation of ligand C–H bonds involves intramolecular C–H insertion. Upon heating in aromatic solvent, the four-coordinate dialkyl complexes $L^{R,iPr}ScR'_2$ (R = tBu, Me; R' = alkyl) (Scheme 12) underwent C–H metalation and eliminated alkane. The half-life of $L^{Me,iPr}ScR_2$ in metalation was significantly longer than its $L^{tBu,iPr}$ analogue, suggesting lower reactivity with the less sterically hindered metal complex.[16]

$L^{R,R'}NiBr$, $L^{R,R'}NiPh(PPh_3)$ and $L^{Me,R'}Ni(alkyl)$ (R = CF_3, Me; R' = iPr, Me) were reported to be active catalysts for ethylene,[102,103] styrene,[104] norbornene[105,106] polymerization and their copolymerization.[107,108] The polymer yield was significantly higher with more hindered ligand systems. Presumably, alkyl insertion into coordinated alkene is greatly facilitated by the more sterically hindered coordination environment.[105]

Reductive elimination is another process facilitated by a crowded coordination environment. With a β-diketiminate-supported Pd(II) methyl phosphine complex, catalytic Castro–Stephens coupling,[109] Stille coupling[110] and Hiyama coupling[111] were more rapid with a more sterically hindered β-diketiminate ligand ($L^{Me,Me}$ vs. $L^{Me,H}$) which gave faster reductive elimination.

In addition, metal-alkyl homolysis is influenced by ligand size. Since chromium(III) alkyl mediated radical polymerization often involves homolysis of the Cr–C bond to gain chain growth, more sterically hindered β-diketiminate ligand increases the Cr–C bond distances (see Table 4), giving a lower BDE, and increasing the rate of homolysis and thus rate of polymerization.[112,113]
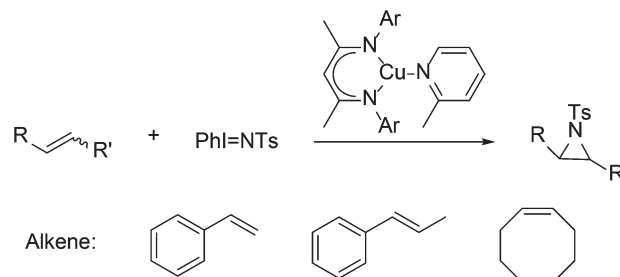
Catalytic carbodiimide formation from isocyanide and organic azide with a diketiminate-iron(I) catalyst gave significantly higher yields with a more sterically bulky catalyst ($L^{tBu,iPr} > L^{Me,Ph_3} > L^{Me,iPr}$). The proposed mechanism involves loss of one molecule of coordinated isocyanide before turning over the catalytic cycle. Not surprisingly, more hindered complexes favor a lower coordination number, which facilitates the loss of isocyanide, production of an active site, and turnover of the catalytic reaction.[114]

LCrCp catalyzed oxygen atom transfer[38] (eqn (1)) and LCu(2-methylpyridine)-catalyzed alkene aziridination[115] (Scheme 13) are also more rapid with more hindered complexes because the smaller catalysts have more rapid rates for corresponding side reactions. Upon formation of the catalytically active [LCr=O] intermediate, $L^{Me,Me}Cr–Cp$ generates $L^{Me,Me}Cr$-$(Cp)(\mu$-O)Cr(Cp)$L^{Me,Me}$ which is inactive towards catalytic oxygen atom transfer from O_2 to PPh_3. In contrast, more hindered $L^{Me,Et}Cr(Cp)$=O is less reactive towards formation of the μ-oxo complex and more catalytically active. Under catalytic aziridination conditions, smaller $L^{Me,Me}Cu(2$-methylpyridine) underwent a side reaction generating $TsNH_2$, which lowered the reactivity and yield of aziridination compared with $L^{Me,Me/iPr}Cu(2$-methylpyridine).

$$PPh_3 + \text{excess } O_2 \xrightarrow{LCr(\eta^5-Cp)} OPPh_3 \qquad (1)$$
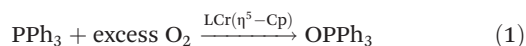
Ethylene polymerization with L_2TiCl_2 complexes supported by different ligands have been studied. $L^{Me,iPr_2}TiCl_2$ and $L^{CF_3,iPr_2}TiCl_2$ showed significantly higher activity than their corresponding $L^{Me,Me}$, $L^{Me,H}$ and $L^{CF_3,Me}$ analogues. In this case, it is possible that bulky N-aryl substituents prohibit β-hydride elimination and thus maintain chain growth.[116] In contrast, $LTiMe_2$ showed a different steric effect, where the less hindered $L^{Me,Me_3}TiMe_2$ was an order of magnitude more reactive than its more hindered $L^{tBu,Me_3}TiMe_2$ and $L^{Me,iPr}Me_2$ analogues.[52]

The steric effect for C–P cross-coupling catalyzed by LCrCp complexes is another interesting example, because the influence is different depending on the relative rate of oxidative addition and Cr–C homolysis.[117] For more reactive alkyl bromide substrates, more hindered $L^{Me,Me}CrCp$ or $L^{Me,Me}Cr(Cp)Br$ gave higher yields than less hindered asymmetric $L^{Me,iPr/p-Me}CrCp$ and $L^{Me,iPr/p-Me}Cr(Cp)Br$. Because these substrates undergo rapid single electron oxidative addition, the rate determining step is homolysis of the Cr–C bond. As previously mentioned,

the Cr–C BDE is lower with more hindered ligands, so these ligands speed the catalytic rate. On the other hand, for less active substrates like Cy–Cl, oxidative addition is rate limiting, and the rate is faster with the less hindering $L^{Me,iPr/p\text{-}Me}$ where one of the N-aryl groups has no ortho-substituents.

3.4. Steric effects on selectivity of metal complexes

Changing steric bulk can also influence the selectivity of reactions of β-diketiminate complexes. This is due to the conformational differences in the energy of the intermediate/transition state with different steric hindrance. In one example, a vanadium(I) β-diketiminate complex catalyzed cyclotrimerization of terminal alkynes at room temperature to give trisubstituted benzenes, with a mixture of isomers.[34] Catalysis with $[L^{Me,Me}V]_2$ gave a 65 : 35 ratio of 1,3,5-trisubstituted benzene over 1,2,4-trisubstituted benzene, whereas the more sterically hindered $[L^{Me,iPr}V]_2$ gave a slightly lower yield with 80 : 20 regioselectivity. The steric restrictions in the transition states or intermediates apparently can prevent formation of products with adjacent substituents.

As mentioned in section 3.3.3, changing the steric bulk can affect the reactivity of alkene polymerization and isomerization catalyzed by $[LNiBr]_2$. Less bulky supporting ligands lead to more rapid β-hydride elimination, giving polyethylene with more branching. In alkene isomerization, the steric hindrance of the ligand can have important influences on the selectivity between cis and trans alkene products. More sterically hindered $[L^{Me,iPr}NiBr]_2$ gave more cis product (44%) compared with $[L^{Me,Me}NiBr]_2$ (28%).[98] It is believed that the crowded coordination environment restricted the rotation of C–C bond in Ni–alkyl complex, hindering the formation of trans-transition states. A bulkier $L^{tBu,iPr}Co$–alkyl complex isomerized alkenes with much higher cis selectivity, often greater than 6 : 1 cis/trans, but the $L^{Me,iPr}Co$ analogue gave poor selectivity. In this cobalt(II) system, the preference of the $L^{tBu,iPr}$ complex for isomerization of terminal alkenes to only the 2 position was also attributed to the bulk of the ligand above and below the N_2Co plane.[88]

4. Electronic effects on β-diketiminate complexes

To tune the electronic properties of β-diketiminate ligands, various groups have been installed on the backbone (α-C and β-C) or on the N-aryl substituents. These modify the electron density at the metal center, which can affect the redox potential, IR frequency of other ligands, UV-Vis absorption maxima, and NMR chemical shifts. In addition, these electronic changes can also affect the reactivity through perturbation of the energy of transition states or intermediates. It should be borne in mind that many of the substituents used to change the electronic effects can also influence sterics as well, particularly on the backbone (β-C) and ortho positions of N-aryl groups.

4.1. Electronic effects on electron density and core structure of the metal center

Changes in electron density on the metal center can be monitored by various methods. Often, electron-withdrawing groups lead to more positive redox potentials, lower field chemical shifts in NMR spectra, and less backbonding into coordinated ligands, consistent with less electron density at the metal ion.

Copper and nickel complexes supported by β-diketiminate ligands bearing different electronic properties have been studied with cyclic voltammetry (Table 5). Judging from the redox potentials in Table 5, NO_2 and CF_3 have the strongest electronic effect, followed by CN and 3,5-bis(trifluoromethyl)-phenyl substituents. In addition, greater electronic effects result from substitutions on α-C and β-C, and less with N-aryl substituents. This is reasonable because the aryl ring is roughly perpendicular to the MN_2C_3 plane, and thus there is little conjugation of the π-systems. In contrast, backbone substituents are in the plane of the ligand backbone, and thus can have a greater impact on the electron density of the metal center. The exception is the relatively small electronic effect from 3,5-bis(trifluoromethyl)phenyl substituents on the backbone (α-C), which is presumably again from lack of conjugation between the perpendicular π-systems. However, the electronic influence of N-aryl substituents is not negligible. For example, alkyl substituents on the N-aryl behaved as electron-donating groups when $^{Ph}L^{H,iPr}$-supported copper complexes had a more negative copper(II/I) potential than $^{Ph}L^{H,Me}$ and $^{Ph}L^{H,Et}$ (Table 5).[30]

Another consequence of the changing redox potentials is the relative stability of certain oxidation levels. In L_2Cu complexes, irreversible reductions were observed with $^{Me}L^{H,H}$ and $^{H}L^{H,H}$ while reversible redox couples were observed in $^{CN}L^{H,H}$

Table 5 Dependence of reduction potential on substituents

Complex	Ligand	Reduction potential[a] (V)	Ref.
$LCu(NCCH_3)^b$	$^{Ph}L^{H,iPr}$	0.384	30
	$^{Ar\text{-}CF_3}L^{H,iPr}$	0.449	30
	$^{Ph}L^{H,Et}$	0.420	30
	$^{Ar\text{-}CF_3}L^{H,Et}$	0.428	30
	$^{Ph}L^{H,Me}$	0.388	30
	$^{CF_3}L^{H,Me}$	0.400	30
	$^{NO_2}L^{H,Mes}$	0.520	30
$LCu(NCCH_3)^c$	$L^{Me,iPr}$	−0.096	68
	$L^{Me/CF_3, iPr}$	0.11	68
	$L^{CF_3, iPr}$	0.411	68
$LCu(OAc)^b$	$L^{Me,iPr}$	−1.29	118
	$L^{Me,iPr/iPr\text{-}CN}$	−1.26	118
	$L^{Me,iPr/Et\text{-}CN}$	−1.24	118
L_2Cu^c	$^{Me}L^{H,H}$	−1.62	46
	$^{H}L^{H,H}$	−1.46	46
	$^{CN}L^{H,H}$	−0.97	46
	$^{NO_2}L^{H,H}$	−0.68	46
L_2Ni^c	$^{Me}L^{H,H}$	−2.42	119
	$^{H}L^{H,H}$	−2.16	119
	$^{Br}L^{H,H}$	−1.89	119
	$^{CN}L^{H,H}$	−1.64	119
	$^{NO_2}L^{H,H}$	−1.28	119

[a] Bu_4NPF_6 was used as electrolyte. [b] All values reported with Fc/Fc^+ in CH_3CN. [c] All values reported with Fc/Fc^+ in THF.

and $^{NO_2}L^{H,H}$, suggesting that the reduced Cu(I) state of the bis (β-diketiminate) complex is unstable in the complexes with more electron rich ligands. Conversely, with LCu(NCCH$_3$) complexes, the Cu(II) state was less stable with a more electron withdrawing group.[46] Ruthenium(II) complexes of $L^{CF_3,m\text{-}CF_3}$Ru-(Cl)(Ar) (Ar = arene ligand) were studied to determine the electronic effects of the supporting ligand on the metal and the other coordinating ligands in comparison to analogous complexes with the $L^{Me,m\text{-}Me}$ supporting ligand.[85] Interestingly, there was no clear trend between the Ru^{II}/Ru^{III} redox potentials from the cyclic voltammograms through the series $L^{Me,Me}$, $L^{Me,m\text{-}Me}$, $L^{CF_3,m\text{-}Me}$, and $L^{CF_3,m\text{-}CF_3}$, indicating that other factors also play a role.[86]

Electronic modification can also have an impact on the positions of the maxima in electronic absorption (UV-Vis) spectra. β-Diketiminate complexes typically have a π → π* transition in the 300–400 nm region, which shifts to shorter wavelength with more electron-withdrawing substituents in LCu-(NCCH$_3$).[30] This suggests that electron-withdrawing groups lower the energy of the π orbital more than they do the π* orbital. The positions of d–d transitions was also studied in L$_2$Cu complexes, where the d–d absorption bands shift toward shorter wavelength with electron withdrawing backbone substituents (α-C) and shift to longer wavelength with more electron donating substituents on the N-aryl group.[46] It is proposed that the ligand field was enhanced with electron donating substituents and thus affected the UV-Vis absorptions.

IR and Raman peaks on coordinated diatomic ligands is another traditional method for quantifying the relative electron density of a metal center. The ν(CO) in LCu(CO) complexes and ν(OO) in LCu(O$_2$) each shift to higher frequency when electron withdrawing CF$_3$ groups were installed on the backbone β-C.[68] This is attributable to a less electron rich metal center that has weaker back-donation into ligand antibonding orbitals. The influence of m-CF$_3$ groups on the N-aryl substituents was less, again indicating a smaller influence from N-aryl substitution.

Due to the shielding or deshielding effect of substituents, the chemical shift in NMR spectra also indicates the electron density on metal center. For example, the chemical shift of the backbone (α-C) proton shifted downfield when CF$_3$ was substituted for CH$_3$ on backbone and for meta-positions on the N-aryl.[85] This is correlated to the deshielding effect with more electron withdrawing groups attached directly to the π system.

Though the introduction of electron withdrawing groups hardly affects the metal ligand core structure, it can affect the coordination number as well as bonding properties in some cases. For example, when NO$_2$ was installed on backbone (α-C) of LCu-OAc, one molecule of methanol coordinated to the metal center, but no coordinated methanol was observed with $^{CN}L^{H,iPr}$ and $^{Ph}L^{H,iPr}$. This is consistent with the stronger Lewis acidity of metal center when its supporting ligand has an electron withdrawing NO$_2$ substituent.[90] Ru–Cl bond lengths and Ru–arene distances in LRu(Cl)(η^6-arene) are shorter with $L^{CF_3,m\text{-}CF_3}$ compared with $L^{Me,m\text{-}Me}$, suggesting an increase in Lewis acidity of the metal with more electron-withdrawing substituents.[85]

4.2. Electronic effects on reactivity of metal complexes

Changes of electron density on the metal center can have a significant effect on reactivity of metal complexes. For example, the oxidative addition of methyl iodide to mixed-aryl LCrCp complexes (Scheme 8) is affected by electronic substituents on para-N-aryl (OMe, Me, H, CF$_3$).[97] There was a correlation between the para-substituent and the rate constant, with the rate constant decreasing two-fold from most electron-donating (para-OMe, k_{obs} = (9.80 ± 0.3) × 10^{-1} M^{-1} s^{-1}) to most electron-withdrawing (para-CF$_3$, k_{obs} = (4.96 ± 0.3)×10^{-1} M^{-1} s^{-1}) substituent. Even though the solid-state structures indicate that the N-aryl planes are aligned roughly perpendicular to the metal–ligand plane, the authors noted that the lack of ortho-substituents may allow the N-aryl to rotate closer to the diketiminate plane in solution, enabling some conjugation. In this way, the more electron-donating substituents can stabilize the chromium(III) product, which could lower the barrier if Hammond's postulate holds.

In another example, catalytic oxidation of alkanes to alcohols and ketones was reported with LCu(OAc) as a catalyst (Scheme 14).[90] When LCu(OAc) was supported by a more electron-withdrawing β-diketiminate ligand, the catalytic reactivity was higher. The results were rationalized through a mechanistic model where the reactions proceed through a metal-based oxidant, based on the observed kinetic isotope effect and regioselectivity.[120] Thus, more electron withdrawing groups would give more unstable and energetic high-valent copper intermediates that are more reactive toward the alkane.

Atom transfer radical addition (ATRA) and atom transfer radical cyclization (ATRC) are particularly interesting for organic synthesis (Scheme 15). Using β-diketiminate ruthenium complexes (LRu(Cp*)Cl and LRu(Cp*)), lower conversions were observed with $L^{Me,Me}$, $L^{Me,m\text{-}Me}$, and $L^{Me,m\text{-}CF_3}$, while the addition of electron-withdrawing substituents in $L^{CF_3,m\text{-}Me}$ and $L^{CF_3,m\text{-}CF_3}$ gave higher reactivity.[86] No simple correlation between catalytic reactivity and redox potential of the ruthenium complexes was observed, but the addition of the CF$_3$ groups also rendered the complexes air-stable in solution and solid state. Likewise, in the copper(I) complexes mentioned above, $L^{Me,iPr}$Cu(NCMe) and $L^{CF_3/Me,iPr}$Cu(NCMe) react with O$_2$, but $L^{CF_3,iPr}$Cu(NCMe) does not react with O$_2$. This agrees with the more positive redox potential with an electron-withdrawing group.[68]

The previously mentioned nickel catalyzed polymerization of styrene and norbornene (see section 3.3) showed a strong influence of the β-diketiminate ligand electronic properties.
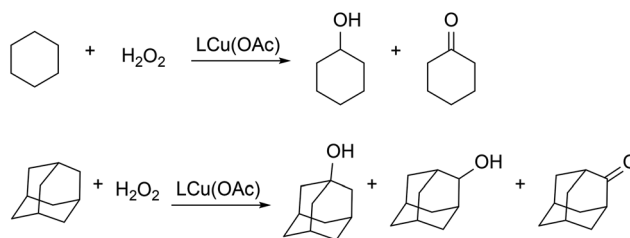
Scheme 14
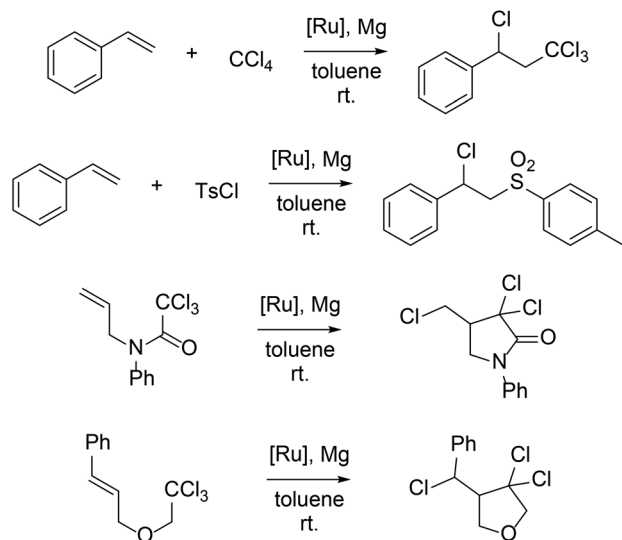
Scheme 15

The substitution of backbone methyl with trifluoromethyl significantly improved the catalytic reactivity.[104,105,121] This can be explained if the more electrophilic nickel center has a lower activation energy for alkene insertion during rate-limiting chain growth.

5. Conclusions

The examples in this Perspective support the idea that β-diketiminate ligands have great tunability in terms of both steric and electronic effects, and they point future chemists in the directions that could benefit their own chemistry. The β-C and N-aryl *ortho* substituents are most important for steric effects, whereas the α-C and β-C positions are most influential for electronic effects. N-Aryl groups can have a small electronic influence, but this has been best documented when there are no *ortho*-substituents and the N-aryl group can rotate closer to planarity with the ligand backbone. In contrast, the steric effects are more varied, because they can change the structure and transition states in different ways depending on the specific coordination number, reaction, and co-ligands. However, the ability of relatively small changes to cause structural, spectroscopic, and reactivity differences suggests that further tuning will uncover multitudes of new chemistry. We note particularly that chiral substituents have only been used in β-diketiminate ligands with N-benzyl substituents,[122–125] and incorporation of chiral anilines should be a fruitful area for preparation of C_1 and C_2 symmetric complexes.

Acknowledgements

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