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Chiral *cis*-iron(II) complexes with metal- and ligand-centered chirality for highly regio- and enantioselective alkylation of N-heteroaromatics†

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Iron-catalyzed highly regio- and enantioselective organic transformations with generality and broad substrate scope have profound applications in modern synthetic chemistry; an example is herein described based on *cis*-Fe^{II} complexes having metal- and ligand-centered chirality. The *cis*-β Fe^{II}(N₄) complex [Fe^{II}(L)(OTf)₂] (L = *N,N'*-bis(2,3-dihydro-1*H*-cyclopenta-[*b*]quinoline-5-yl)-*N,N'*-dimethylcyclohexane-1,2-diamine) is an effective chiral catalyst for highly regio- and enantioselective alkylation of N-heteroaromatics with α,β-unsaturated 2-acyl imidazoles, including asymmetric N1, C2, C3 alkylations of a broad range of indoles (34 examples) and alkylation of pyrroles and anilines (14 examples), all with high product yields (up to 98%), high enantioselectivity (up to >99% ee) and high regioselectivity. DFT calculations revealed that the "chiral-at-metal" *cis*-β configuration of the iron complex and a secondary π–π interaction are responsible for the high enantioselectivity.

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Introduction

The development of highly enantioselective metal-catalyzed reactions usually relies on the design of robust ligand scaffolds containing tetrahedral chiral, axial chiral or planar chiral centers positioned close to the metal ion for effective stereo-differentiation of incoming/coordinated substrate(s). Recently, Meggers and co-workers reported that optically active complexes with metal-centered chirality including iron complexes are potent catalysts for enantioselective organic transformation reactions.¹ A challenge in developing these complexes for asymmetric catalysis, particularly over first-row transition metal complexes that are relatively substitutionally labile, is the potential racemization of the stereogenic metal center.^{2b} In this regard, the use of multidentate ligands such as the tetradentate N₄ ligands used in this work could make the resulting complexes more configurationally stable.^{2a,c} Acyclic tetradentate ligands form stable octahedral complexes with three possible topologies: *trans*, *cis*-α, and *cis*-β configurations.^{2a,b} Complexes with the *cis*-α and *cis*-β configuration feature a stereogenic metal center and a pair of *cis*-oriented

vacant sites for binding the incoming substrate in a η² fashion. Evans and co-workers reported a chiral *cis*-β Al^{III}-salen complex as an effective catalyst for enantioselective aldol reactions of aldehydes and 5-alkoxyoxazoles.³ Yamamoto and co-workers developed a *cis*-β Al^{III} catalyst with a tethered bis(8-quinolinolato) (N₂O₂) ligand for various highly enantioselective organic transformations.⁴ Chiral Fe^{II} complexes supported by tetradentate N₄ ligands have been shown to exhibit remarkable catalytic activities in asymmetric hydrocarbon oxidations.⁵ In a recent study, we showed that the chiral Fe(N₄) complexes with bis(quinolyl)diamine ligands can efficiently catalyze highly enantioselective *cis*-dihydroxylation of alkenes with H₂O₂ (up to 99.8% ee).⁶ These Fe^{II} complexes possess metal- and ligand-centered chirality with a rigid quinoline-based ligand scaffold and therefore a robust chiral pocket, which is in close proximity to the iron center, for discriminative activation/binding of the incoming reagent/substrate and subsequent reactions. We envision that the chiral pocket together with the Lewis acidity of the Fe^{II}/Fe^{III} ion would make chiral Fe(N₄) complexes potent chiral Lewis acid catalysts.^{7,8}

Chiral indoles are ubiquitous structural motifs in bioactive natural products and pharmaceuticals.⁹ Over the past few decades, numerous efforts have been dedicated to developing catalytic asymmetric reactions for the synthesis of these compounds.¹⁰ As the C3 position of indoles is more reactive than the C2 and N1 positions, many elegant methodologies have been reported for the construction of chiral C3-alkylated indoles.¹¹ In contrast, enantioselective alkylations at the C2 and N1 positions of indoles are limited. For asymmetric C2 alkylation of indoles, the literature examples were realized

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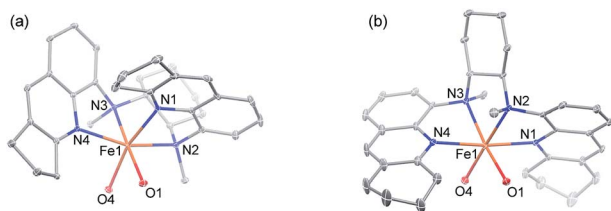


Fig. 1 ORTEP drawings of (a) **1e** and (b) **1f** (30% probability ellipsoids) with the omission of H atoms and OTf[−] ligands except O atoms bound to Fe^{II}.

through the Pictet–Spengler reaction,¹² alkylation/oxidation of 4,7-dihydroindoles,^{11c,13} Michael addition^{14,15} and other methods.¹⁶ So far, examples of highly enantioselective N1 alkylation of indoles have been sparse.¹⁷ In view of the prevalence of chiral indoles in pharmaceuticals and the stringent constraint on trace noble metal levels in the pharmaceutical industry, the development of catalysts based on earth-abundant, biocompatible iron metal is appealing. Herein is described highly regio- and enantioselective C3, C2 and N1 alkylation of indoles, as well as alkylation of pyrroles and anilines, with α,β -unsaturated 2-acyl imidazoles catalyzed by chiral *cis*-iron(II) complexes supported by tetradentate bis(quinolyl)diamine ligands.

Results and discussion

Synthesis and characterization of the catalyst

Following a previously reported method for the synthesis of complexes **1a**,^{6,18a} **1b** and **1c**,⁶ chiral Fe(II) complexes **1a–1f** were synthesized, by the reaction of Fe^{II}(OTf)₂·(CH₃CN)₂ with a series of chiral bis(quinolyl)diamine N₄ ligands **2a–2f**, in 80–95% yields (Scheme 1). The new complexes **1d–1f** were characterized by ESI-MS, elemental analysis and circular dichroism

spectroscopy (ESI[†]). The structures of **1e** and **1f**, featuring fused cyclopentane and cyclohexane rings on the quinoline moiety, respectively, were determined by X-ray crystallography, revealing a *cis*- α configuration for **1f**, similar to that for **1b**·(H₂O)₂,⁶ but, unexpectedly, a *cis*- β configuration for **1e**. As shown in Fig. 1, **1e** adopts a *cis*- β - Δ coordination geometry, where the Fe–N distances of 2.163(2)–2.242(2) Å are comparable to those of *cis*- β -[Fe^{II}(BQCN)(OTf)₂] (BQCN = **2a** in Scheme 1) reported by Nam and co-workers (Fe–N 2.125(2)–2.224(2) Å).^{18b} In contrast, **1f** adopts a *cis*- α - Δ configuration (Fig. 1); its Fe–N distances of 2.220(6)–2.292(5) Å are comparable to those of **1b**·(H₂O)₂ (*cis*- α - Δ ; Fe–N 2.195(5)–2.261(5) Å),⁶ while considerably longer than those of *cis*- α -[Fe^{II}(BQCN)(CH₃CN)₂](ClO₄)₂ (Fe–N 2.112(3)–2.192(3) Å),^{18b} attributed to the steric repulsion between the quinolyl substituents and the *cis*-coordinated ligands. Notably, the two methyl groups on the cyclohexane-1,2-diamine nitrogen atoms are oriented *syn* to each other in **1e** but *anti* to each other in **1b** and **1f**. Consistent with the relatively long Fe–N distances, both **1e** (μ_{eff} = 4.92 μ_{B}) and **1f** (μ_{eff} = 5.44 μ_{B}) are high-spin with *S* = 2 as determined by the Evans method. These complexes are configurationally stable; no change in the circular dichroism (CD) signal was observed upon standing an acetonitrile solution of **1e** or **1f** at room temperature for up to 4 days (Fig. S1, ESI[†]). The ¹H NMR spectrum of **1e** in CD₃CN at room temperature showed no significant change after the solution was left standing for 8 days (Fig. S2, ESI[†]).

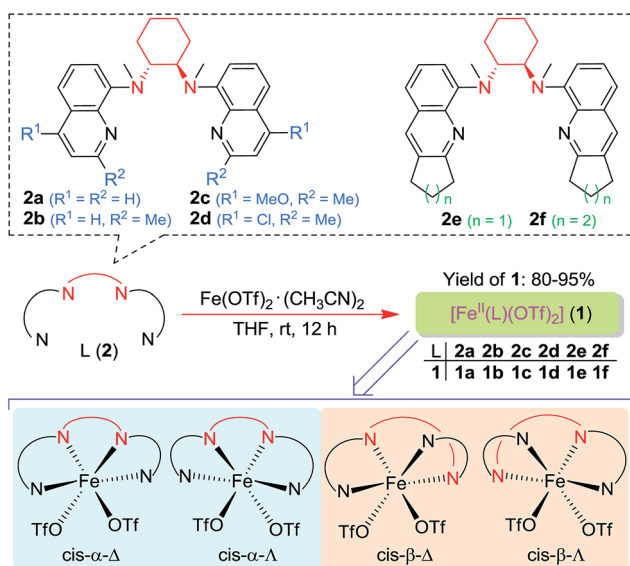
Catalyst screening

At the outset, the catalytic activity of Fe(II) complexes **1a–1f** was examined for C3 alkylation of *N*-methylindole **3a** with α,β -

Table 1 Screening of the catalysts for the asymmetric alkylation of *N*-methylindole **3a** with α,β -unsaturated 2-acyl imidazole **4a**^a

Entry	Cat.	Solvent/ <i>T</i> (°C)	<i>T</i> (h)	% Y ^b	% ee ^c
1	1a	CH ₂ Cl ₂ /25	3	90	65
2 ^d	1a	CH ₂ Cl ₂ /25	4	92	60
3 ^e	1a	CH ₂ Cl ₂ /25	5	88	58
4	1a	CH ₂ Cl ₂ /5	10	91	73
5	1b	CH ₂ Cl ₂ /5	10	98	79
6	1c	CH ₂ Cl ₂ /5	15	90	69
7	1d	CH ₂ Cl ₂ /5	12	87	57
8	1e	CH ₂ Cl ₂ /5	10	99	85
9	1f	CH ₂ Cl ₂ /5	10	94	71
10	1e	CH ₂ Cl ₂ /−15	24	95	90
11	1e	ClCH ₂ CH ₂ Cl/−15	24	97	93
12	1e	CH ₃ Cl/−15	24	96	91
13	1e	CH ₃ CN/−15	20	92	95
14	1e	CH ₃ CN/−25	48	91	97.5

^a Reaction conditions: **3a** (0.45 mmol), **4a** (0.3 mmol), **1** (5 mol%), 4 Å MS (100 mg) and solvent (3 mL), under argon. ^b Isolated yield. ^c Determined by chiral-phase HPLC. ^d Without 4 Å MS. ^e Without 4 Å MS, under air.



Scheme 1 Chiral N₄ ligands **L** (**2a–2f**) and Fe^{II} complexes [Fe^{II}(**L**)(OTf)₂] (**1a–1f**) employed in this work. The bottom shows four possible configurations for such complexes.

unsaturated 2-acyl imidazole **4a** in the presence of molecular sieves (MS) and under argon (under air and/or without MS, the reaction time increased and enantioselectivity reduced, cf. entries 1–3 in Table 1). Complex $[\text{Fe}^{\text{II}}(6\text{-Me}_2\text{-BPMCN})(\text{OTf})_2]$,^{5a} bearing a bis(pyridylmethyl)amine N_4 ligand, was also tested; this complex did not catalyze the alkylation reaction under similar conditions. Revealingly, the reaction of **3a** and **4a** in the presence of 5 mol% $\text{Fe}(\text{II})$ catalysts **1a–1f** in CH_2Cl_2 at 5 °C for 10–15 h (entries 4–9, Table 1) afforded the alkylation product **5a** in high yields (87–99%) and with moderate-to-good enantioselectivity (57–85% ee). Complex **1e** was found to exhibit the highest catalytic activity and enantioselectivity (entry 8). With **1e** as the catalyst, lowering the reaction temperature to –15 °C improved the enantioselectivity to 90% (entry 10). A screening of solvents at this temperature (entries 10–13) revealed that CH_3CN gave the highest enantioselectivity of 95% ee. Further lowering the reaction temperature to –25 °C afforded **5a** in 91% yield and 97.5% ee (entry 14).

Reaction/substrate scope

Under the optimized conditions, we examined the substrate scope of the $\text{Fe}(\text{II})$ -catalyzed C3 alkylation of various indoles with α,β -unsaturated 2-acyl imidazoles. As shown in Table 2, in the presence of 5 mol% of **1e**, a series of substituted indoles reacted with α,β -unsaturated 2-acyl imidazole **4a** and the desired alkylation products **5b–5i** were obtained in high yields (87–98%) and with excellent enantioselectivities (95–99% ee). No C2

alkylation product was observed. The sterically hindered 1,2-dimethylindole **3i** could also undergo C3 alkylation, giving **5i** in 98% yield and 95% ee. The absolute configuration at the stereogenic carbon center of **5i** was determined to be *R* by X-ray crystallography (Fig. 2). The C3 alkylation of *N*-methyl-5-bromoindole proceeded smoothly at –25 °C to give **5f** in 90% yield and 95% ee. However, the presence of the more electron-withdrawing substituent, Cl, on the 5-position of indole was found to deactivate the reaction. This issue was addressed by increasing the temperature to 0 °C, which led to alkylation product **5g** in 87% yield and 95% ee. The scope of α,β -unsaturated 2-acyl imidazoles was also examined. As shown in Table 2, various α,β -unsaturated 2-acyl imidazoles are reactive for the C3 alkylation of indoles to give the corresponding products **5j–5m** in high yields (87–97%) and enantioselectivities (91–96% ee). Neither β - nor *N*-substitution of α,β -unsaturated 2-acyl imidazoles was found to have significant impact on the product enantioselectivity.

We next investigated the C2 alkylation of 3-substituted indoles. The reaction of 3-methylindole and α,β -unsaturated 2-acyl imidazole **4a** with 5 mol% of **1e** in CH_2Cl_2 at room temperature led to the corresponding C2-alkylated product **6a** in 83% yield and 87% ee along with the N1 alkylation product **7a'** (**6a/7a'** = 85 : 15, Table S2, ESI†). CH_3CN as solvent was found to disfavor the C2 alkylation reaction (Table S2, ESI†). Decreasing the reaction temperature to –40 °C further improved the enantioselectivity to 96% ee, although with an extended reaction time of 60 h. Under the optimized conditions, a variety of 3-substituted indoles reacted with α,β -unsaturated 2-acyl imidazoles to give the C2 alkylation products in 76–96% yields and ~72–98% ee (Table 3). It is noteworthy that the reaction of 3,7-dimethylindole with **4b** bearing a bulkier *N*-substituent afforded **6j** in 94% yield and 96.5% ee, without detectable N1 alkylation product. This is attributed to the presence of the 7-methyl substitution which makes N1 alkylation sterically unfavorable. When β -ester substituted α,β -unsaturated 2-acyl imidazole was used as the alkylating agent, in which both the α and β carbons are electrophilic, the nucleophilic addition of indoles took place at the β position (**6f–6h**). X-ray crystallography of **6j** revealed an *R* configuration at the stereogenic carbon center (Fig. 2), which is the same as that of **5i**.

For the reaction of 3-methylindole with α,β -unsaturated 2-acyl imidazoles, the N1 alkylation product was obtained in

Table 2 Asymmetric C3 alkylation of substituted indoles **3** with α,β -unsaturated 2-acyl imidazoles **4** catalyzed by **1e**^a

	5b , 90%, 98% ee	
	5d , 95%, >99% ee	
	5e , 93%, 96.5% ee	
	5g ^b , 87%, 95% ee	
	5h , 96%, >99% ee	
	5j , 97%, 96% ee	
	5k , 92%, 93% ee	
	5m , 87%, 91% ee	

^a Reactions conditions: **3** (0.45 mmol), **4** (0.3 mmol), **1e** (5 mol%), 4 Å MS (100 mg), CH_3CN (3 mL), under argon; yield of the isolated product; ee's determined by chiral-phase HPLC. ^b At 0 °C.

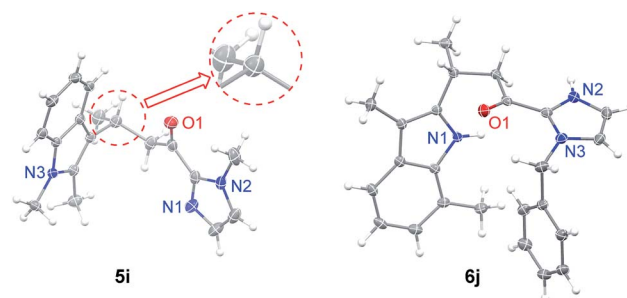


Fig. 2 X-ray crystal structures of **5i** and **6j**. Thermal ellipsoids are drawn at the 50% probability level.



Table 3 Asymmetric C2/N1 alkylation of 3-substituted indoles **3** with α,β -unsaturated 2-acyl imidazoles **4** catalyzed by **1e**^a

C2 Alkylation		
6a , 85%, 96% ee	6b , 86%, 97.5% ee	6c , 92%, 91.5% ee
6d , 87%, 92% ee	6e ^b , 76%, 89% ee	6f , 93%, 91% ee
6g , 91%, 90.5% ee	6h , 92%, 71.5% ee	6i , 93%, 97% ee
6j , 94%, 96.5% ee	6k ^c , 96%, 91.5% ee	6l ^d , 93%, 92% ee
Z = CO ₂ Me		
N1 Alkylation		
7a , 83%, 95.5% ee	7b , 74%, 95% ee	7c , 65%, 88% ee
7d , 85%, 74.5% ee	7e , 93%, 95.5% ee	7f , 76%, 94% ee
7g , 81%, 93.5% ee	7h , 72%, 94% ee	7i ^d , 86%, 76% ee

^a Reaction conditions: **3** (0.45 mmol), **4** (0.3 mmol), **1e** (5 mol%), 4 Å MS (100 mg), CH₂Cl₂ (3 mL), under argon, −40 °C; yield of the isolated product; ee's determined by chiral-phase HPLC. ^b Catalyzed by **1b**. ^c At 0 °C. ^d At −20 °C.

~10% yield, implying a competition between C2 and N1 positions as nucleophiles on the olefinic electrophile. We envisioned that the alkylation of indoles at the nitrogen atom would be favored by increasing the bulkiness of the C3 substituent. To validate this hypothesis, we examined the reaction of 3-ethylindole with α,β -unsaturated 2-acyl imidazole **4b** catalyzed by complex **1e** at −40 °C. Gratifyingly, the reaction afforded the N1 alkylation product in 83% yield and 95.5% ee (Table 3, **7a**). The C2 alkylation product in this case was obtained in less than 10% yield. As shown in Table 3, a range of 3-substituted indoles are reactive for the N1 alkylation, giving the desired products in good yields (65–93%) and with high enantioselectivity (74.5–95.5% ee). It was found that the electron-donating substituent

on the indole makes N1 alkylation more favorable. For example, the reaction of 5-methoxy-3-methylindole with **4b** gave the N1 alkylation product in 93% yield and 95.5% ee (Table 3, **7e**). This is in contrast to the dominating C2 alkylation in the reaction of 3-methylindole and **4b** (Table 3, **6b**), suggesting that the electronic effect, in addition to the steric effect, of the substituent on the indoles also plays a significant role in directing the regioselectivity. The substituents of α,β -unsaturated 2-acyl imidazoles were also found to have a significant impact on the regioselectivity of the Fe(II)-catalyzed reaction. For example, when β -ester substituted α,β -unsaturated 2-acyl imidazole was used as the substrate, C2 alkylation is favored over N1 alkylation irrespective of the choice of indoles (compare **7b** vs. **6g** and **7e** vs. **6h**, Table 3), suggesting that the electronic effect of the β -ester group of α,β -unsaturated 2-acyl imidazoles outweighs the electronic and steric effects of substituents on the indoles on determining the regioselectivity. Sterically hindered 2,3-dimethylindole is also reactive for the N1 alkylation at −20 °C to give **7i** in 86% yield and 76% ee.

The Fe(II)-catalyzed asymmetric addition reaction is also applicable to the alkylation of pyrroles.^{11a,c,e,i,19} The reaction of *N*-phenyl pyrrole with α,β -unsaturated 2-acyl imidazole **4a** and 5 mol% of **1e** in CH₃CN at −25 °C gave the C2 alkylation product **9a** (Table 4) in 89% yield and 93% ee. The ee value reached 99% (**9b**) when a bulkier *N*-substituent is present on the α,β -unsaturated 2-acyl imidazole **4b**. In both cases, the C3 alkylation product was not detected; this is consistent with the C2 position of pyrroles being more nucleophilic than the C3 one. The C3 alkylation of pyrroles was achieved with 2,5-substituted pyrroles, which led to the formation of the C3-alkylated products (**9c–9f**) in high yields (87–94%) and with excellent enantioselectivity (94–99% ee).

Alkylation of electron-rich arenes such as anilines with α,β -unsaturated 2-acyl imidazoles^{11c,l} could also be achieved using the Fe(II) complex **1e** as the catalyst. The **1e**-catalyzed alkylation reaction of **10** with **4** gave *para*-alkylated **11a–11g** in 87–95%

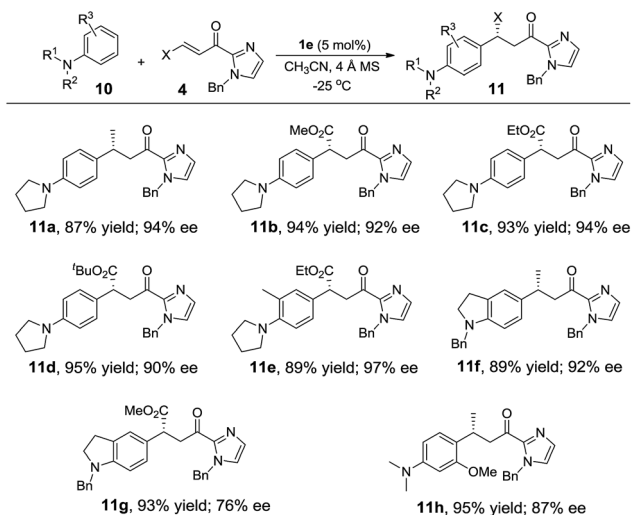
Table 4 Enantioselective alkylation of pyrroles **8** with α,β -unsaturated 2-acyl imidazoles **4** catalyzed by **1e**^a

9a , 89%, 93% ee	9b , 87%, >99% ee	9c , 92%, 94% ee
9d , 94%, >99% ee	9e , 87%, 94% ee	9f , 93%, 94% ee

^a Reaction conditions: **8** (0.45 mmol), **4** (0.3 mmol), **1e** (5 mol%), 4 Å MS (100 mg), CH₃CN (3 mL), under argon.



Table 5 Enantioselective alkylation of *N,N*-disubstituted anilines **10** with α,β -unsaturated 2-acyl imidazoles **4** catalyzed by **1e**^a

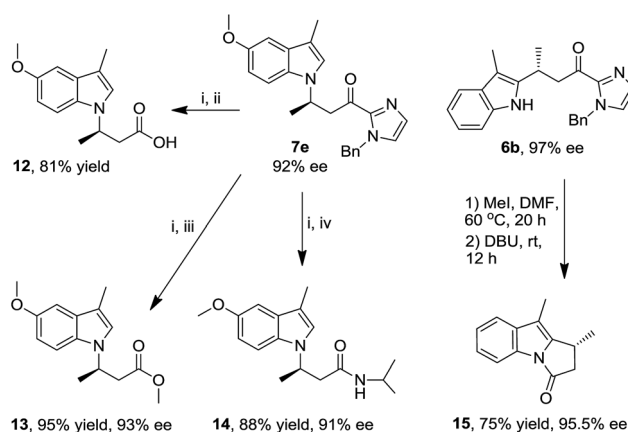


^a Reaction conditions: **10** (0.45 mmol), **4** (0.3 mmol), **1e** (5 mol%), 4 Å MS (100 mg), CH₃CN (3 mL), under argon.

yields and 76–97% ee (Table 5). When 3-methoxy-*N,N*-dimethylaniline was used as the substrate, in which two strongly electron-donating groups are present, the reaction proceeded in high regioselectivity giving **11h** in 95% yield and 87% ee (Table 5).

Synthetic applications

The indolyl 2-acyl imidazoles obtained in this work could be readily converted to a range of indole compounds^{11a,c} which are of interest in medicinal chemistry. For example, methylation of the imidazole moiety of N1 alkylation product **7e** with iodomethane in DMF, followed by hydrolysis with water and DBU, gave the indolyl carboxylic acid **12** in 81% yield (Scheme 2), the analogues of which have been reported to be potent inhibitors



Scheme 2 Synthetic elaboration studies. Reaction conditions: (i) MeI, DMF, 60 °C, 20 h; (ii) DBU, H₂O, rt, 4 h; (iii) DBU, CH₃OH, CH₂Cl₂, rt, 6 h; (iv) DBU, *i*PrNH₂, rt, 12 h.

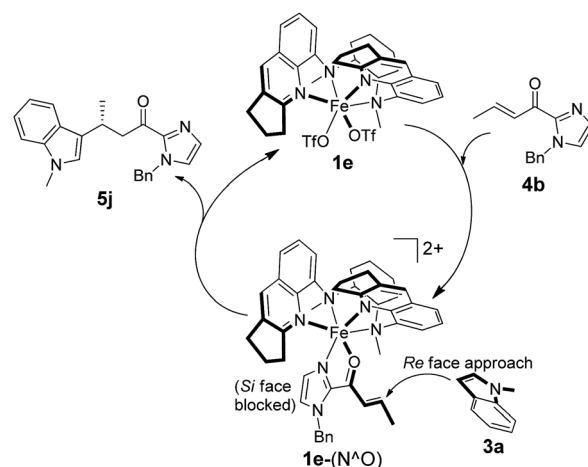
of integrin $\alpha_v\beta_3$.²⁰ In a similar fashion, the ester and amide derivatives **13** and **14** were obtained in 95% and 88% yield, respectively (Scheme 2). The amide **14** can be reduced to an analogue of 3-(1*H*-indol-1-yl)-3-arylpropan-1-amine;²¹ the latter was reported to display inhibition to norepinephrine and serotonin reuptake.²² The pyrrolo[1,2- α]indole scaffold is a basic structure of many bioactive natural products.²³ Methylation and subsequent intramolecular acylation of the C2 alkylation product **6b** afforded pyrrolo[1,2- α]indole **15** in 75% yield (Scheme 2). Notably, these organic transformations are accompanied by retention of ee values, indicating preservation of the stereogenic carbon centers.

Mechanistic studies

A proposed reaction mechanism for the **1e**-catalyzed alkylation of indoles is presented in Scheme 3. Binding of α,β -unsaturated 2-acyl imidazoles (e.g. **4b**) via the imidazolyl nitrogen and carbonyl oxygen atoms to the chiral Fe(II) center generates an intermediate denoted as **1e**-(N⁺O). The Fe(II) catalyst acts as a Lewis acid to activate the C=C bond for nucleophilic attack by indoles (e.g. **3a**) to give the addition product (e.g. **5j**). The enantioselectivity is governed by the chiral pocket environment around the Fe(II) center which shields the *Si* face from indolyl nucleophile attack.

To gain support for the proposed reaction mechanism, the binding between **1e** and **4b** was examined by high-resolution ESI-MS. Treatment of an acetonitrile solution of **1e** with 2 equiv. of **4b** resulted in a predominant ion peak at *m/z* 379.1707 (Fig. 3), where the isotopic distribution pattern (Fig. S3, ESI[†]) and collision-induced dissociation analysis (Fig. S4, ESI[†]) are well consistent with the proposed formulation of **1e**-(N⁺O).

DFT calculations were performed to examine the reaction details of **1e**-catalyzed C3 alkylation of *N*-methylindole (**3a**) with α,β -unsaturated 2-acyl imidazole **4b** and to elucidate the origin of the enantioselectivity (Scheme 4). Since the two *cis*-oriented coordination sites of **1e** are chemically inequivalent, there are two possible binding interactions between **1e** and **4b** (IntA₁ and



Scheme 3 Proposed catalytic cycle for asymmetric alkylation of indoles catalyzed by **1e**.



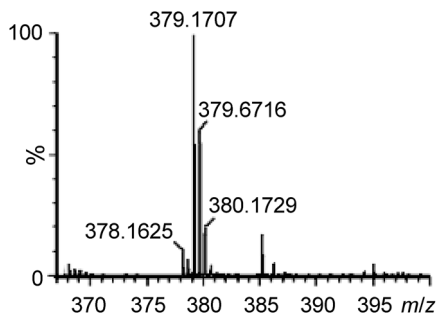


Fig. 3 High-resolution ESI-MS spectrum of a reaction mixture of **1e** with **4b** in CH_3CN .

IntB₁). Both binding modes were calculated to form stable intermediates with a decrease in free energy by more than 10 kcal mol⁻¹ as a result of replacement of two coordinated solvent molecules (acetonitrile) by imidazole **4b** as a bidentate ligand. In addition, **IntB₁** is 3.8 kcal mol⁻¹ more stable than **IntA₁**. Accordingly, there are four possible transition states (**TSA_{2R}**, **TSA_{2S}**, **TSB_{2R}** and **TSB_{2S}**) regarding the two approach directions of **3a** to the C=C bond of the **1e**-(N⁺O) intermediate (Scheme 4). Among the four transition states, **TSA_{2R}** was found to have the lowest transition barrier with a free energy of 17.3 kcal mol⁻¹ (**TSA_{2S}**: 21.4 kcal mol⁻¹; **TSB_{2R}**: 23.4 kcal mol⁻¹; **TSB_{2S}**: 23.1 kcal mol⁻¹). There is π - π stacking interaction between the quinoline ring of **1e** and the indole ring of **3a** in **TSA_{2R}** (Fig. S5, ESI[†]) with a distance of about 3.4 Å, which possibly accounts for the lower energy barrier. Accordingly, the indole ring resides closer to the quinoline ring in **TSA_{2R}** and the additional π - π interaction leads to the comparatively restrained position of the indole, and thus its C₁-C₃ bond (2.08 Å) is slightly longer than those in the other three transition states (2.03 or 2.04 Å). The *R*-enantiomer of the C3 alkylation product is mainly obtained *via* **TSA_{2R}**, which is consistent with the configuration observed experimentally in the isolated products (e.g. **5i**, Table 2). The free energy differences between the four transition states (**TSA/B_{2R/S}**) lead to a calculated ee of 96.3% at -25 °C, which is in excellent agreement with the experimental result of 96% ee. The subsequent steps proceed feasibly with much lower energy barriers or great energy release. When the C₁-C₃ bond is formed (**IntA_{3R}**, **IntA_{3S}**, **IntB_{3R}** and **IntB_{3S}**), the most stable intermediate is **IntA_{3S}**, which probably stems from its strong hydrogen bond between H₁ and O (1.87 Å), compared with other intermediates (2.16, 2.22, and 2.59 Å for **IntA_{3R}**, **IntB_{3R}** and **IntB_{3S}**, respectively). After forming the C₁-C₃ bond, intramolecular proton transfer takes place from C₃ to O by forming a six-membered ring (**TSA_{4R}**, **TSA_{4S}**, **TSB_{4R}** and **TSB_{4S}**). In these transition states, the Fe-O bonds are weakened (from 2.01–2.04 in **IntA/B_{3R/S}** to 2.08–2.17 Å in **TSA/B_{4R/S}**) in association with the shortening of the C₁-C₃ bond (from 1.58–1.61 Å in **IntA/B_{3R/S}** to 1.54 Å in **TSA/B_{4R/S}**). Both **TSA_{4R}** and **TSA_{4S}** hold the π - π stacking between the quinoline and indole, which could stabilize **TSA_{4R}** and **TSA_{4S}** and therefore lead to the free energy difference between **TSA_{4R/S}** and **TSB_{4R/S}**. After proton transfer, four possible enol intermediates are generated (**IntA_{5R}**,

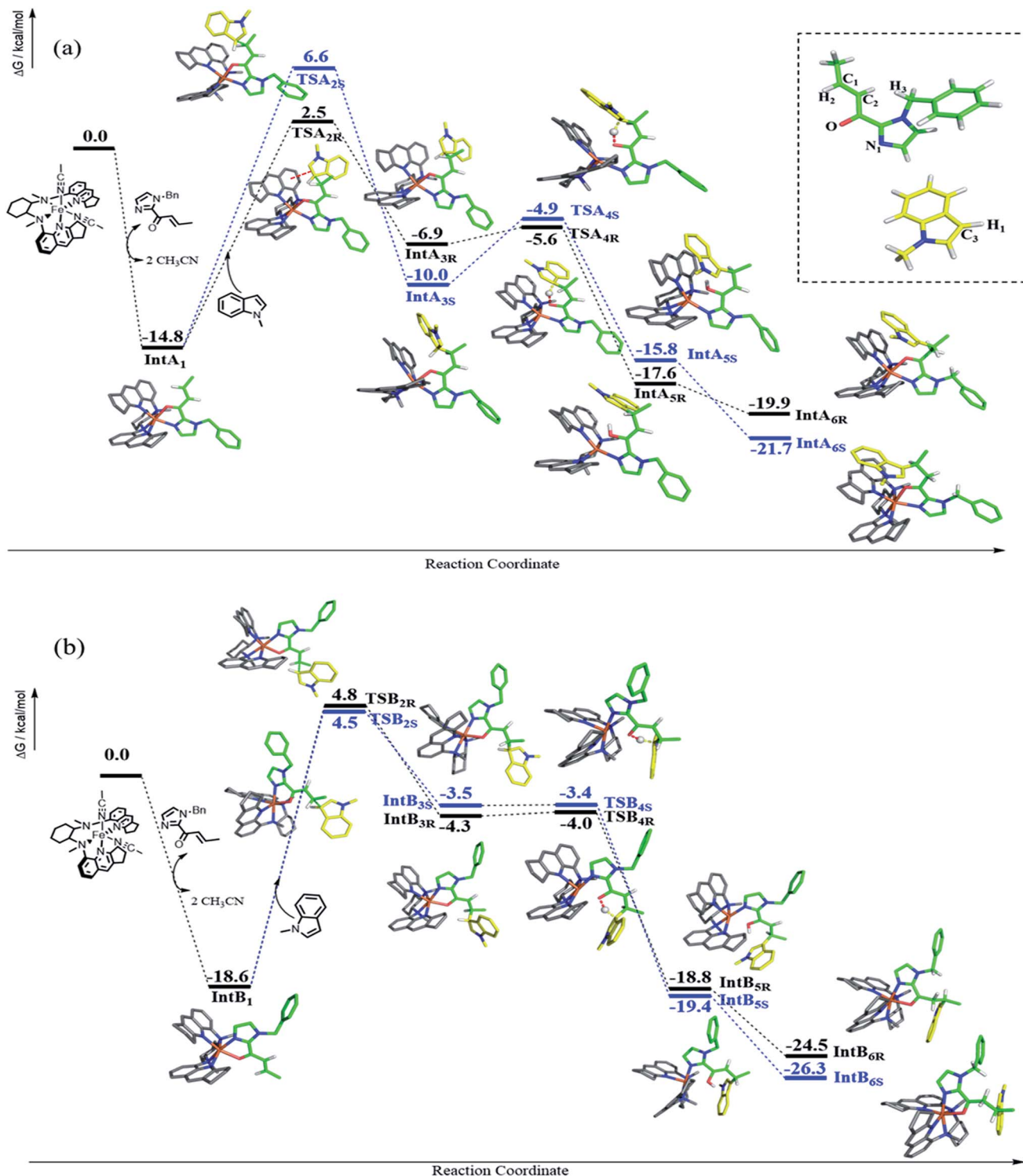
IntA_{5S}, **IntB_{5R}** and **IntB_{5S}**) with significant energy release (5.8–16.0 kcal mol⁻¹). Comparatively, the lower free energy of **IntB_{5R}** and **IntB_{5S}** possibly results from their stronger Fe-N₁ coordination than those in **IntA_{5R}** and **IntA_{5S}**. The subsequent exothermic tautomerization leads to the final alkylation product coordinated to the catalyst (**IntA_{6R}**, **IntA_{6S}**, **IntB_{6R}** and **IntB_{6S}**), which would possibly result in some extent of product inhibition of the catalyst. Indeed, time-course studies revealed that addition of 25 mol% of product **5j** to the initial reaction mixture of the **1e**-catalyzed reaction of **3a** and **4b** appreciably lowered the product yield (Fig. S6, ESI[†]).

General remarks

The Fe(II) catalysis reported herein is reminiscent of an Ir(III) system developed by Meggers and co-workers for the same type of reaction,^{1a} where the “chiral-at-metal” property of the catalyst is responsible for the excellent enantioselectivity as the reaction occurs in close proximity to the metal bearing “intense” chiral information. These Ir(III) catalytic systems composed of achiral ligands are very potent in effecting a wide range of enantioselective reactions and could be accessed in relatively facile pathways.^{1b} However, it should be noted that such systems rely heavily on specific heavy metal ions, *i.e.* Ir(III) and Rh(III), and/or specific ligand sets, *i.e.*, cyclometalating ligands, to maintain a high coordination stability to prevent catalyst racemization. These features are generally not compatible with the development of base metal catalysts, *e.g.* Fe complexes, when it comes to harnessing their non-toxicity and earth-abundance for chemical synthesis. In this regard, we demonstrate here, along with other reported examples,^{2,3} that efficient “chiral-at-metal” catalysis could be achieved with base metals through rational ligand design, *e.g.* using chiral tetradentate N₄ ligands, installing rigidity, and fine-tuning the sterics on the ligand fragments.

In recent years, chiral Fe(N₄) complexes have been extensively studied in oxidative organic catalysis such as asymmetric alkene epoxidation,^{5c-g} asymmetric alkene *cis*-dihydroxylation^{5a,b,6} and asymmetric C-H hydroxylation.^{5h} The class of [Fe^{II}(N₄)(OTf)₂] complexes presented in this work, as well as those in the literature,^{5,6} features some of the fundamental defining properties of an appealing chiral transition-metal asymmetric catalyst, including (1) ease of preparation and modification, (2) configurationally stable chirality, (3) possession of labile coordination sites for reagent/substrate activation, and (4) high sustainability. The chiral, linear N₄ ligands could be prepared on the multigram scale with good yields from commercially available optically active diamine backbones. The steric and electronic properties of the pyridyl/quinolyl donors are highly flexible for manipulation, as have been demonstrated by us⁶ and by Costas and co-workers.^{5e} Upon coordination to Fe(II) ions, under mild conditions, the N₄ ligands form stable *cis*- α or *cis*- β chelates depending on the ligand structure. These complexes possess metal-centered chirality without further needs of chiral resolution or purification. Thanks to the large barrier possibly resulting from the different orientations of the *N*-methyl groups in the *cis*- α and *cis*- β isomers, these chelates do





Scheme 4 The reaction details of the Fe(II) complex **1e**-catalyzed C3 alkylation of *N*-methylindole **3a** by α,β -unsaturated 2-acyl imidazole **4b**. Owing to the inequivalent *cis*-oriented binding sites of **1e**, two pathways are presented in panel a and panel b, representing coordination pattern A and B, respectively. Because of the stereoselectivity of the attacked alkene, the reaction pathways of both A and B possess both *R* and *S* configurations, which are displayed in black and blue, respectively. In each transition state, the breaking and forming bonds are highlighted by dashed lines. Color code: iron, orange; oxygen, red; nitrogen, blue; carbon, gray for **1e**, yellow for **3a** and green for **4b**; hydrogen (white). For simplicity, only critical hydrogen atoms are displayed. Atomic labels referred to in the text are denoted in the inset of panel a.

not undergo isomerization in solution;^{6,18b} thus the stereogenic configuration is robust. The weakly coordinating triflate (OTf⁻) ligands, in addition to the fast ligand substitution rate of Fe(II),

are favorable for substrate binding/activation. Beyond that, iron is readily available, relatively inexpensive and non-toxic which makes the utilization of iron complexes in catalysis particularly

attractive.²⁴ In our continuing efforts to develop synthetically useful catalysis mediated by chiral Fe(N₄) complexes, we were interested to examine their potential as Lewis acid catalysts which has not been documented to the best of our knowledge.

As there are many organic reactions of synthetic interest that can be catalyzed by Lewis acids, the development of effective chiral Lewis acid catalysts for asymmetric organic transformations has been receiving considerable interest in organic chemistry. The N₄ ligands employed in this work are quinolylamines. Due to the conjugation with the quinoline moiety, the amino group of bis(quinolyl)diamines is a less effective donor to the metal ion than that of bis(pyridylmethyl)diamine ligands, thereby making the corresponding Fe(II) complexes more Lewis acidic and capable of catalyzing a wide range of addition reactions, even at low temperature. In addition, due to the lack of a methylene moiety, tetradentate quinolylamine ligands have a more rigid scaffold than pyridylmethylamine analogues, which would provide a robust chiral pocket around the reaction center for reactions.

In the literature, reports on asymmetric C3 addition of indoles to α,β -unsaturated 2-acyl imidazoles are confined to the use of the Sc(III) complex,^{11a} Cu(II)-DNA/protein adducts,^{11e,f} and Ir(III) complex^{1a} as Lewis acid catalysts. In this work, we report for the first time that chiral Fe^{II}(N₄) complexes could catalyze similar reactions.

Due to the inherently low nucleophilicity of the nitrogen atom, asymmetric N1 alkylation of indoles with α,β -unsaturated 2-acyl imidazoles remains a formidable challenge. There are only a few examples of highly enantioselective N1 alkylation of indoles reported in the literature, including intramolecular aza-Michael additions,^{17b,e} cinchona alkaloid-catalyzed reaction with Morita-Baylis-Hillman carbonates,^{17c} Pd(II) and Ir(I)-catalyzed *N*-allylations^{17d,f,i} and Ir(I)-catalyzed allylation/oxidation of indolines.^{17h} Our present method provides an alternative access to *N*-alkyl indoles in good yields and high enantiomeric excess *via* Fe(II)-catalyzed intermolecular alkylation which is unprecedented in the literature. To the best of our knowledge, **1e** also represents the first example of a single chiral catalyst capable of effecting intermolecular alkylation of various indoles at the C3, C2 and N1 positions, respectively, with high enantio- and regioselectivities.

DFT calculations provide some insight into the origin of high enantioselectivities in the **1e**-catalyzed reactions. The two possible binding interactions between **1e** and α,β -unsaturated 2-acyl imidazole **4b**, in addition to the two possible approach faces (*Si* or *Re*) of indole **3a** to the **1e**-(N⁺O) adduct, generate four possible reaction pathways. In all the four pathways, nucleophilic attack of **3a** on **1e**-(N⁺O) was found to have the largest barrier (17.3–23.4 kcal mol⁻¹) and logically this is the enantioselectivity-determining step. Among them, **TSA**_{2R} was computed to have the lowest transition barrier (17.3 kcal mol⁻¹), being 4.1 kcal mol⁻¹ lower than that of the second-lowest **TSA**_{2S}; the calculated ee at -25 °C derived from this energy difference is in excellent agreement with the experimental value. In the DFT-optimized structure of **TSA**_{2R}, interestingly, there is substantial π - π stacking between the quinoline moiety of **1e** and indole **3a**. This suggests that the

enantioselectivity is not only determined by how well the steric environment around the *cis*- β Fe(II) center can guide/shield the approach of indoles, but also assisted by secondary π - π interaction provided by the quinolyl ligand. The DFT-calculation findings are insightful for the rational design of future generation of Fe^{II}(N₄) Lewis acid catalysts with more effective chiral pockets for the enantio-differentiation of aromatic substrates.

Conclusions

In summary, we have developed C3, C2 and N1 alkylations of indoles catalyzed by a chiral *cis*- β Fe(II) complex supported by a (*R,R*)-tetradentate bis(quinolyl)diamine ligand. A range of indoles reacted with various α,β -unsaturated 2-acyl imidazoles to give diverse chiral indoles in high yields with high enantio- and regioselectivity. The catalysis is also applicable to the alkylation of pyrroles and anilines. The thus obtained products can be converted to a variety of chiral indole derivatives with potential synthetic and biological values. The stable configuration of the iron(II) complex and a secondary π - π interaction in the transition state are suggested to be the origin of the high enantioselectivity. Our results suggest that chiral Fe^{II}(N₄) complexes are potentially a new and efficient class of Lewis acid catalysts.

Conflicts of interest

There are no conflicts to declare.

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