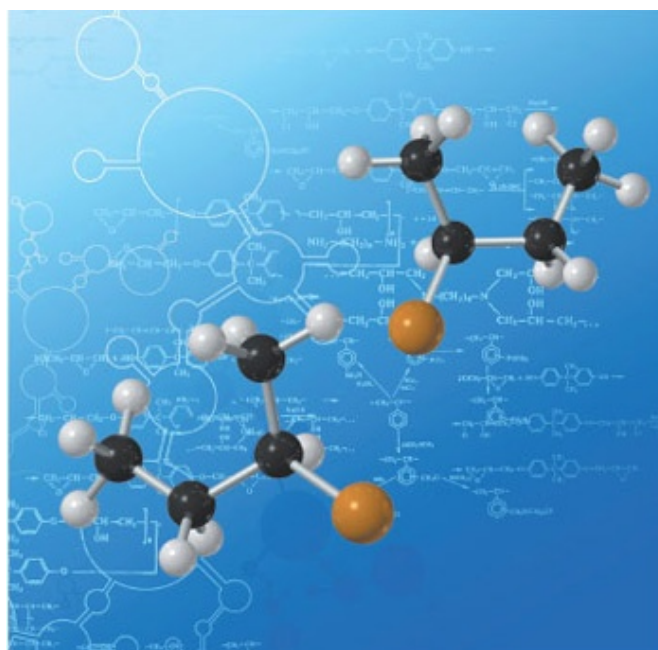This article is part of the

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Kinetics and mechanism of organocatalytic aza-Michael additions: direct observation of enamine intermediates†‡

Sami Lakhdar, Mahiuddin Baidya and Herbert Mayr*

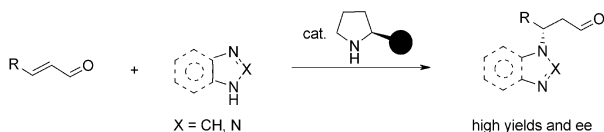**The imidazoles 1a–g add to the CC-double bond of the iminium ion 2 with rate constants as predicted by the equation log k = $s_N(N + E)$. Unfavourable proton shifts from the imidazolium unit to the enamine fragment in the adduct 3 account for the failure of imidazoles to take part in iminium-activated aza-Michael additions to enals.**

Since MacMillan's pioneering work in 2000,[1] the so-called iminium activation has become one of the most attractive methods in asymmetric synthesis.[2] In his seminal paper, MacMillan showed that α,β-unsaturated aldehydes can be activated by the addition of catalytic amounts of chiral secondary amines; the initially generated iminium ions undergo fast Diels–Alder reactions with dienes to give cycloadducts, which release the chiral catalyst upon hydrolysis.[1] By using this strategy it has become possible to realise a large variety of enantioselective organic reactions.[3]

In 2007, Jørgensen *et al.*[4] and Vicario *et al.*[5] independently reported the first enantioselective aza-Michael additions of nitrogen heterocycles to aliphatic unsaturated aldehydes (Scheme 1), using diarylprolinol silyl ethers or imidazolidinones as catalysts.[6]

While these reactions proceeded readily with tetrazoles and triazoles, they generally failed with imidazoles and benzimidazoles.

Only 4,5-dicyano-imidazole was found to react with moderate yield and low enantioselectivity with aliphatic enals utilizing MacMillan's second generation imidazolidinone as a catalyst.[5] Under the same conditions, the parent imidazole **1c** gave only traces of the product.

We now report the kinetics and mechanism of the reactions of imidazoles with the iminium ion **2** and rationalise why imidazoles, in contrast to triazoles and tetrazoles, do not undergo organocatalytic aza-Michael additions.

In previous work we have shown that the rates of the reactions of carbocations and Michael acceptors with n, π, and σ nucleophiles can be described by eqn (1),[7] where k_2 is a second-order rate constant in M^{-1} s^{-1}, s_N is a nucleophile-specific sensitivity parameter, N is a nucleophilicity parameter, and E is an electrophilicity parameter.

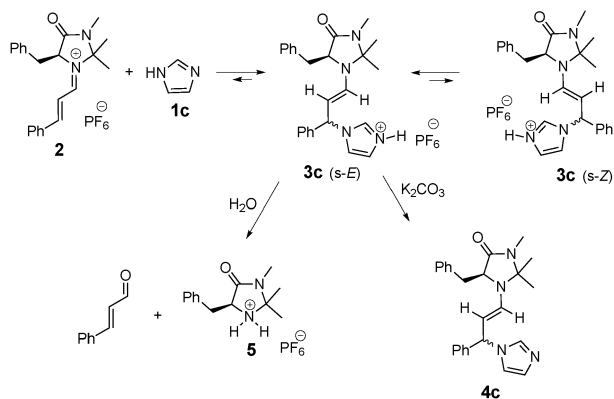$$\log k_2 \, (20 \, °C) = s_N(N + E) \qquad (1)$$

Using the known reactivity parameters N and s_N of the imidazoles **1a–g**[8] (Table 1) and the electrophilicity parameter of the cinnamaldehyde-derived iminium ion **2**[9] ($E = -7.37$), we had calculated second-order rate constants of 300–3000 M^{-1} s^{-1} by eqn (1) indicating that the reactions of **2** with **1a–g** should proceed readily.

Accordingly, treatment of the iminium salt **2-PF₆** with 4 equivalents of imidazole **1c** leads to the formation of the enamine **3c**, which bears a protonated imidazole ring. Adduct **3c**, which incorporates two stereocenters, was formed as a 1 : 1 mixture of two diastereoisomers, as revealed by ^{1}H and [13]C

Scheme 1 Aza-Michael additions of *N*-heterocycles to enals catalysed by chiral secondary amines.[4–6]

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‡ Electronic supplementary information (ESI) available: Details of the kinetic experiments, synthetic procedures and product characterisation. See DOI: 10.1039/c2cc31224g

Table 1 Nucleophile-specific reactivity parameters N and s_N for azoles in acetonitrile[8]

Azoles	N	s_N	Azoles	N	s_N
1a	10.50^a	0.79^a	**1e**	11.74	0.76
1b	11.43	0.79	**1f**	11.79	0.77
1c	11.47	0.79	**1g**	11.90	0.73
1d	11.51	0.84			

^{a} N and s_N refer to DMSO, as 4-(dimethylamino)pyridine was reported to have identical reactivity parameters in DMSO and acetonitrile ($\Delta N = 0.15$)[10] the variation of solvent does not affect the analysis in Table 2.
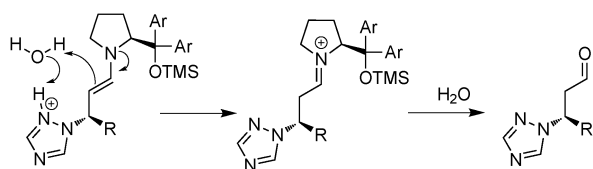
Scheme 2

NMR spectroscopy. NOE analysis showed the preference of the (s-E)-conformation of the C(sp^2)–N bond in both diastereoisomers as depicted in Scheme 2. As kinetically controlled additions of nucleophiles to **2** often proceed with high diastereoselectivity, the formation of **3c** as a 1 : 1 mixture of diastereomers may either be explained by a reversible reaction of the iminium ion **2** with the imidazole **1c** or by deprotonation of the imidazolidinone moiety in the intermediate iminium ion in the α-position to the carbonyl group, as suggested by Seebach *et al.*[11] The latter mechanism can be excluded under the conditions of this work because we have recovered the enantiopure imidazolidinone **5** after hydrolysis of **3c**, confirming that the configuration of the asymmetric center of the imidazolidinone **5** has not been affected.
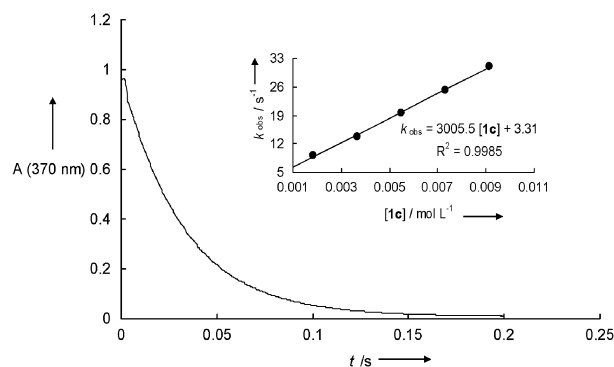
In acetonitrile solution, the enamine intermediates **3c** (or **4c** in the presence of excess imidazole) are stable for more than 8 hours but decompose during several days. Addition of water to the enamine **3c** leads to the formation of cinnamaldehyde. However, stirring of a solution of **3c-PF$_6$** in CD$_3$CN with dry K$_2$CO$_3$ led to the formation of **4c**, which was characterised by ^{1}H and ^{13}C NMR spectroscopy (see ESI‡).[12,13]

Jørgensen's DFT calculations on the organocatalytic conjugate addition of 1,2,4-triazole to α,β-unsaturated aldehydes showed that the addition of the triazole to the iminium ion is followed by a water-assisted proton transfer from the triazolium ring to the enamine as depicted in Scheme 3.[4] In the case of the reactions of the azoles **1** with the iminium ion **2**, the enamines **3** or their conjugate bases **4** were observed by NMR spectroscopy, and we did not observe an analogous proton transfer which may be explained by the lower acidities of the imidazolium ions compared to triazolium ions.

The different UV-absorbances of the iminium ions **2** and the adducts **3** allowed us to follow the kinetics of the reactions of **2** with the azoles **1a–g** photometrically at the absorption maximum of the iminium ion **2** (370 nm). All kinetic experiments were performed under first-order conditions using a high excess of the

Fig. 1 Exponential decay of the absorbance at 370 nm during the reaction of **2-OTf** (5.10×10^{-5} M) with imidazole **1c** (5.47×10^{-3} M). Inset: determination of the second-order rate constant k_2 from the dependence of the first-order rate constant k_{obs} on the concentration of imidazole **1c** (20 °C in CH$_3$CN).

nucleophiles **1a–g**. From the exponential decays of the UV-absorbances of the electrophile **2**, the first-order rate constants k_{obs} were obtained.

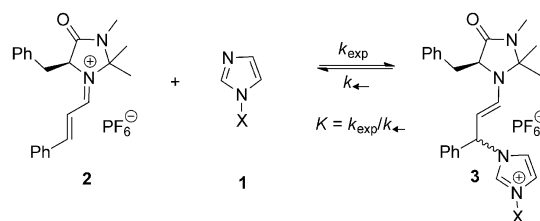Plots of k_{obs} (s^{-1}) against the concentrations of the nucleophiles **1a–g** were linear (Fig. 1) and their slopes gave the second-order rate constants k_2 (M^{-1} s^{-1}) which are summarised in Table 2.

The second-order rate constants thus obtained (Table 2) have been compared with those calculated by eqn (1) from the electrophilicity parameter $E = -7.37$ of the iminium ion **2**[9] and the N and s_N parameters of the azoles.[8] Table 2 shows that all calculated (k_{calc}) and experimental rate constants (k_{exp}) match within a factor of two. This good agreement is impressive, as E(**2**) has been derived from reactions with C-nucleophiles and N and s_N for the azoles **1a–g** have been derived from their reactions with benzhydrylium ions.[7]

Some of the reactions of the iminium ion **2** with imidazoles proceeded incompletely, and for the reactions with **1b** and **1g** the equilibrium constants K (Scheme 4) have been determined photometrically as described in the ESI.‡

Table 2 Comparison of experimental (k_{exp}) and calculated rate constants (k_{calc}, using eqn (1)) for the reactions of the azoles **1a–g** with the iminium ion **2** ($E = -7.37$) in CH$_3$CN at 20 °C

Azoles	k_{exp}/M^{-1} s^{-1}	k_{calc}/M^{-1} s^{-1}	k_{exp}/k_{calc}
1a	2.75×10^2	2.97×10^2	0.93
1b	2.88×10^3	1.61×10^3	1.8
1c	3.01×10^3	1.73×10^3	1.7
1d	3.00×10^3	3.00×10^3	1.0
1e	2.84×10^3	2.10×10^3	1.4
1f	4.62×10^3	2.53×10^3	1.8
1g	4.15×10^3	2.03×10^3	2.0

Scheme 3

Scheme 4 Reversible additions of the imidazoles **1b** and **1g** to the iminium ion **2** in CH$_3$CN.

Table 3 Equilibrium constants (K) for the reactions of azoles **1b** and **1g** with the iminium ion **2** (counterion PF_6^-) in CH_3CN at 20 °C

Azoles	K/M^{-1}	$\Delta G^{\neq}/$ kJ mol^{-1}	$\Delta G^0/$ kJ mol^{-1}	$\Delta G_0^{\neq}/$ kJ mol^{-1}	$k_{\leftarrow}/s^{-1}$
1b	1.61×10^3	52.3	−18.0	61.0	1.79
1g	1.84×10^2	51.4	−12.7	57.6	22.6

Table 3 shows that 1-(trimethylsilyl)-imidazole **1b**, which is 1.4 times less nucleophilic than 1-methylimidazole **1g** (Table 2), is a nine-fold stronger Lewis base than **1g**. Substitution of the rate and equilibrium constants into the Marcus equation (2) yields the intrinsic barriers $\Delta G_0^{\neq}$, which are defined as the activation energies of processes with $\Delta G^0 = 0$.[14]

$$\Delta G^{\neq} = \Delta G_0^{\neq} + 0.5\,\Delta G^0 + ((\Delta G^0)^2/16\Delta G_0^{\neq}) \qquad (2)$$

Remarkably, the intrinsic barrier for the addition of the azole **1g** to the iminium ion **2** is about 10 kJ mol^{-1} lower than that for its reaction with diarylcarbenium ions.[8] This difference reflects that more reorganisation energy is needed for the reactions of nucleophiles with diarylcarbenium ions than with unsaturated iminium ions due to the more extensive delocalisation of the positive charge in diarylcarbenium ions.

In conclusion, we have shown that the reactions of the iminium ion **2** with imidazoles proceed readily with formation of stable enamines which have been fully characterised by NMR spectroscopy. The failure of the azoles **1a–g** to act as nucleophiles in iminium-activated processes is rationalised by the low acidities of the initially generated azolium species which do not undergo proton shifts. The rate constants determined for the reactions of the iminium ion **2** with the azoles **1a–g** are in good agreement with those calculated by eqn (1), showing the suitability of the benzhydrylium-based reactivity parameters N and s_N for predicting reactivities toward iminium ions.

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