

Squaramide-Catalyzed Enantioselective Michael Addition of Masked Acyl Cyanides to Substituted Enones

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S Supporting Information

ABSTRACT: Masked acyl cyanide (MAC) reagents are shown to be effective umpolung synthons for enantioselective Michael addition to substituted enones. The reactions are catalyzed by chiral squaramides and afford adducts in high yields (90–99%) and with excellent enantioselectivities (85–98%). The addition products are unmasked to produce dicyanohydrins that, upon treatment with a variety of nucleophiles, provide γ -keto acids, esters, and amides. The use of this umpolung synthon has enabled, in enantiomerically enriched form, the first total synthesis of the prenylated phenol (+)-fornicin C.

The importance of umpolung chemistry stems from its capacity to provide access to functional group arrangements that are difficult to realize through normal chemical reactivity considerations.¹ For example, whereas 1,3-dicarbonyl compounds are readily synthesized through classical reactivity (nucleophilic enolate plus an electrophilic carbonyl reactant), 1,4-dicarbonyl compounds, having dissonant connectivity,² can prove challenging. An efficient route to such compounds is through an umpolung strategy, the reaction of an enone with a masked acyl anion. Given their value in synthesis, many acyl anion equivalents have been developed and utilized for a variety of synthesis problems.³ Among useful and versatile umpolung synthons are protected hydroxyl malononitriles, known as masked acyl cyanide (MAC) reagents, developed by Yamamoto and Nemoto.⁴ Treatment of a MAC reagent with base generates a nucleophilic acyl anion equivalent that will react with a variety of electrophilic units. Subsequent unmasking of the MAC unit produces an acyl cyanide, which can be intercepted to afford carboxylic acids, esters, or amides.⁵ Thus, unlike typical acyl anion equivalents, MAC reagents also harbor masked acid chloride-like reactivity. This dual capability, found in few acyl anion equivalents, allows MAC reagents to function as carbon monoxide equivalents.⁶ The usefulness of these reagents is demonstrated by their use in the total syntheses of (–)-bestatin, cyclotheonamide C, and oseltamivir (tamiflu).^{7–9} Despite their importance in synthesis, there appears to be no report of a catalytic enantioselective reaction of a MAC reagent.¹⁰ The ease of deprotonation of MAC reagents suggested that their reactions could be rendered enantioselective using a bifunctional chiral base. In this report, we disclose the first enantioselective reactions of the MAC family of umpolung synthons.^{11,12} Specifically, we show that Michael additions of MAC reagents to enones are efficiently catalyzed by chiral squaramides, affording adducts in high yields and excellent enantioselectivities

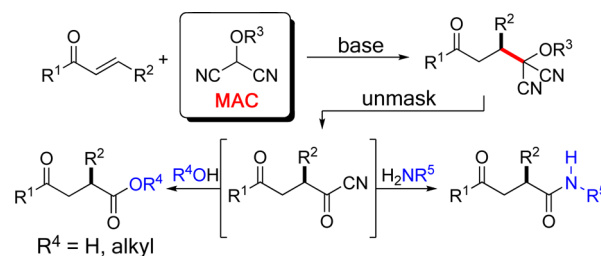


Figure 1. Synthesis of 1,4-dicarbonyls via addition of MAC reagents.

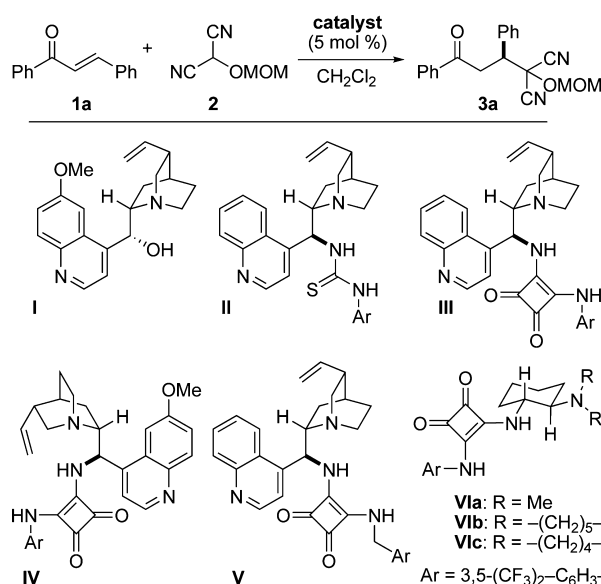


Figure 2. A selection of chiral catalysts examined.

and, upon unmasking, provide ready access to chiral γ -keto-carboxylic acids, -esters, and -amides (Figure 1).

Our initial efforts focused on the identification of effective catalysts and conditions for the enantioselective Michael reaction between *trans*-chalcone (**1a**) and MOM-MAC (**2**) to afford adduct **3a** (Figure 2).^{13,14} Although quinine (**I**) promoted this reaction, it afforded the adduct with poor enantioselectivity. The cinchonidine-derived thiourea catalyst (**II**) gave better enantioselectivity but low conversion. On the other hand, the corresponding squaramide (**III**) provided both superior conversion and enantioselectivity. As anticipated, the pseudo-

Received: August 30, 2013

Published: October 3, 2013

Table 1. Substrate Scope of Squaramide-Catalyzed MAC Addition Reactions^a

Reaction scheme: Enone **1** (R¹-C(=O)-CH=CH-R²) reacts with squaramide catalyst **2** (R³ = MOM) or **4** (R³ = TBS) in CH₂Cl₂ using **VIc** (5 mol %) as a catalyst to form product **3** (R³ = MOM) or **5** (R³ = TBS).

entry	enone	R ³	product	temp (°C)	time (h)	yield ^c (%)	ee ^d (%)	entry	enone	R ³	product	temp (°C)	time (h)	yield ^c (%)	ee ^d (%)
1		MOM	3a	-30	24	98	97	10		MOM	3j	-10	28	98	95
		TBS	5a	-20	36	96	96			TBS	5j	0	48	96	94
2		MOM	3b	-30	20	98	97	11		MOM	3k	23	24	97	88
3		MOM	3c	-30	24	98	97	12		MOM	3l	23	24	96	92
4		MOM	3d	0	28	99	92	13 ^e		MOM	3m	-10	72	96	89
		TBS	5d	0	72	96	90			TBS	5m				
5		MOM	3e	-30	24	99	98	14		MOM	3n	-30	20	96	96
		TBS	5e	-20	36	95	97	TBS	5n	-20	36	98	94		
6		TBS	5f	0	24	93	90	15		MOM	3o	-30	28	96	97
7		TBS	5g	-10	48	94	90	16		MOM	3p	-30	24	98	96
8		MOM	3h	-30	30	98	91	17		MOM	3q	23	24	94	89
		TBS	5h							TBS	5q	-20	36	92	94
9		MOM	3i	0	36	97	92	18		MOM	3r	-30	24	98	97
		TBS	5i	0	72	90	89	TBS	5r						
								19		MOM	3s	-30	24	99	94
										TBS	5s	-20	36	93	85

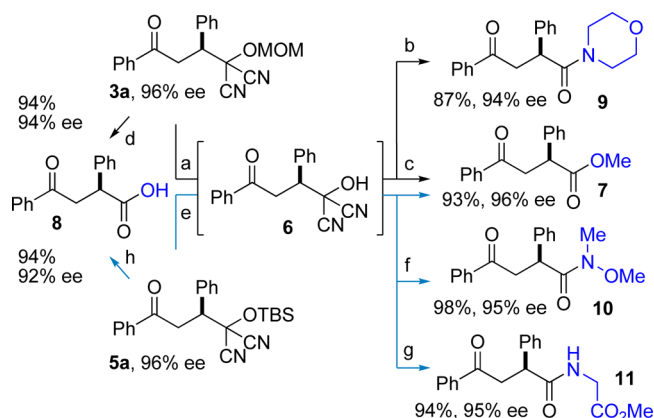
^aConditions: reactions performed with **1** (0.36 mmol), **2** or **4** (0.30 mmol), **VIc** (5 mol %), CH₂Cl₂ (0.6 mL). ^cIsolated yield. ^dee determined by chiral stationary phase HPLC. ^e4 Å MS was used as an additive to avoid hydrolysis.

enantiomeric, quinidine-derived squaramide (**IV**) provided the product with comparable selectivity and conversion but enriched in the opposite enantiomer. Among the other squaramide catalysts examined, those based on 1,2-diaminocyclohexane proved most effective, particularly the pyrrolidine catalyst **VIc**, which formed **3a** at a high rate and with 90% ee.^{15,16} The enantiomeric excess improved significantly, to 97%, when the reaction was carried out at -30 °C. It is noteworthy that comparable enantioselectivity was observed even at 1 mol % catalyst loading, albeit with lower conversion. Moreover, increasing the catalyst loading to 10 mol % did not significantly increase the conversion over the 5 mol % reaction, presumably due to the limited solubility of the catalyst.¹⁷ Additional solvents were also investigated for the reaction, but they did not lead to an improved outcome.

We next examined the practical utility and substrate scope of squaramide-catalyzed enantioselective Michael reactions of MAC reagents (Table 1). In general, high yields and enantioselectivities were observed for a broad range of enones. Importantly, the related TBS-MAC reagent (**4**)^{4b} was also found to be effective for the enantioselective Michael reaction. Although **4** displays lower reactivity compared to **2**, requiring higher reaction temperatures and/or longer times for complete reaction, it provides comparable enantioselectivity while offering an alternative deprotection protocol (vide infra). Enones having

electron-deficient aryl groups at the β -position gave excellent results (entries 2–3). The more electron-rich substrate, *p*-methoxyphenyl enone **1d**, required higher reaction temperatures and provided reduced enantioselectivity (entry 4). The Michael addition was also effective with naphthyl enone **1e** and various heteroaryl-substituted enones (entries 5–9). Conjugated enone **1j** reacted selectively at the β -position to afford the addition products in excellent yields and enantioselectivities (entry 10). The related substituted dienone **1k** delivered the addition product with a notable drop in enantioselectivity, coinciding with its reduced reactivity (entry 11). The reaction of enone **1l**, which contains a free phenol, is noteworthy as it demonstrates the tolerance for acidic hydrogen-bond donor groups in the substrate (entry 12). The successful use of benzotriazole-containing enone **1m** (entry 13) is noteworthy.

Enones having β -alkyl-substituents were also found to be effective substrates for the MAC-Michael reaction.¹⁸ The reaction of both MAC reagents with enones having primary alkyl substituents proceeded well, affording the addition products in near-quantitative yields and with high enantioselectivities (entries 14–16). The more sterically hindered cyclohexyl-enone **1q** (entry 17) required room temperature for the reaction to proceed but still gave the product with 89% ee. Cyclopropyl enone **1r**, on the other hand, was more reactive and gave the addition product in excellent yield and selectivity (entry

Scheme 1. Functionalization of MAC Adducts^a

^aConditions: (a) 1:1 AcOH/DME (0.5 M), (R)-CSA (50 mol %), 60 °C, 2 h; (b) morpholine, Et₃N, -40 °C; (c) MeOH, Et₃N, -40 °C; (d) 1:1 AcOH/H₂O (0.5 M), (R)-CSA (50 mol %), 15 h; (e) 3HF·Et₃N, THF, -30, 2 h; (f) HN(OMe)Me·HCl, Et₃N, -40 °C; (g) H₂NCH₂CO₂Me·HCl, Et₃N, -40 °C; (h) TBAF, H₂O:THF:AcOH (1:1:2), 24 h.

18). The Michael reaction of MOM-MAC and trifluoromethyl-substituted enone **1s** proceeded nicely, providing straightforward access to chiral units containing the trifluoromethyl group. Curiously, for reasons that are unclear, the same reaction with TBS-MAC gave significantly lower selectivity with this substrate.

The optimized conditions are suitable for the preparation of gram quantities of the Michael adducts. For example, the reaction with chalcone **1a** was scaled up to produce 1 g of product, formed in the same yield and enantioselectivity as noted in entry 1. Additionally, in large-scale reactions the low solubility of the catalyst makes it convenient to recover most of it (84%) from the ether extracts by simple filtration. Such recovered catalyst can be used multiple times without measurable loss in efficacy.

We recognized that the above reactions would be of value only if the MAC group in the Michael adducts could be transformed into useful functional groups without eroding enantioselectivity. A careful examination of hydrolysis conditions allowed us to identify mild, efficacious conditions for unmasking the MAC moiety. For the MOM-MAC adduct **3a**, excellent results were obtained if the intermediate acyl cyanide, which is prone to racemization, was intercepted at a relatively low temperature (ca. -40 °C).¹⁹ Thus, treatment of the labile dicyanohydrin (**6**), formed upon removal of the MOM group, at -40 °C with MeOH/Et₃N or morpholine/Et₃N, gave ketoester **7** or ketoamide **9**, respectively, in high yields and with little or no enantiomeric excess loss (Scheme 1). Ketoacid (**8**) can be formed directly by treating the MOM-adduct with CSA in aqueous acetic acid. Conditions were also developed for the efficient transformation of the TBS-MAC adduct **5a** to methyl ester **7**, Weinreb amide **10**, and glycine-derived amide **11** as well as for its direct transformation to ketoacid **8**.²⁰

The absolute stereochemistry induced during the Michael reaction was determined by taking advantage of the unique reactivity of acyl benzotriazoles.²¹ The MAC addition product of cinnamyl-benzotriazole (**1m**) catalyzed by (R,R)-**Vic** was first converted to methyl ester **12**, unmasking of which generated the known succinate ester **13**, enriched in the *S* enantiomer (Scheme 2). The observed asymmetric induction is consistent with a pretransition state assembly shown in Figure 3, wherein the ammonium salt directs nucleophilic addition of MAC anion to

Scheme 2. Determination of Absolute Stereochemistry

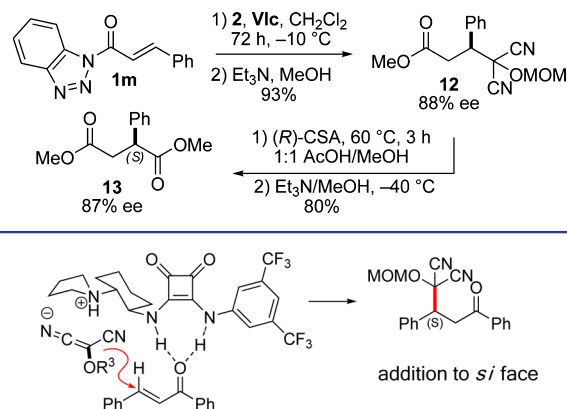
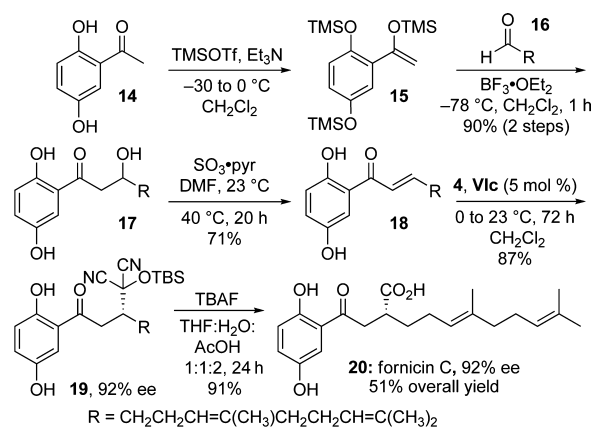


Figure 3. Model for asymmetric induction.

Scheme 3. Total synthesis of Fornicin C



chalcone, which in turn is activated through hydrogen bonding. Analogous models have been posited for thiourea- and squaramide-catalyzed asymmetric reactions.^{14b,h}

The utility of the above enantioselective methodology is demonstrated through the total synthesis of prenylated phenol (+)-fornicin C (Scheme 3). Isolated from *Ganoderma fornicatum*, extracts of which are reported in Chinese medicine to have health-promoting properties,²² fornicin C displays moderate in vitro cytotoxic activity against the human larynx carcinoma cell line Hep-2 (23 μg/mL). Triol **17** can be prepared directly from the aldol reaction of the trianion of 2,5-dihydroxyacetophenone (**14**) with geraniol-derived aldehyde **16** in ca. 50% yield²³ or in a higher yield via the Mukaiyama aldol reaction of silyl enol ether **15** with **16**. Dehydration to the enone was an unforeseen challenge since basic conditions promoted cyclization to a chromanone, while strongly acidic conditions introduced complications through reactions with the geranyl side chain. Furthermore, with dehydrating agents such as Martin's sulfuranol or Burgess' reagent, the phenols competed with the secondary alcohol. We found that sulfur trioxide-pyridine complex was effective in achieving the dehydration of triol **17** to enone **18**.²⁴ Addition of TBS-MAC (**4**) to enone **18** provided adduct **19** with 92% ee.²⁵ When subjected to AcOH-buffered TBAF, **19** was successfully unmasked to reveal (+)-fornicin C in 51% overall yield, with complete retention of ee.

To summarize, we have developed the first catalytic, enantioselective reaction of the MAC family of umpolung synthons. We have shown that chiral squaramide **Vic** catalyzes

the conjugate addition reactions of MAC reagents **2** and **4** to a collection of electronically and structurally diverse enones to afford adducts in high yields and with excellent enantioselectivities. The addition products can be unmasked to afford γ -keto-carboxylic acids, -esters, and -amides, all formed in excellent yields and with near complete retention of enantiomeric excesses. We have utilized this methodology for the total synthesis of (+)-fornicin C, prepared in five steps from known aldehyde **16**, in 51% overall yield and 92% ee. The development of other enantioselective reactions of MAC and related umpolung reagents is expected to be of great value in organic synthesis.

■ ASSOCIATED CONTENT

Supporting Information

Experimental procedures and characterization data. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Notes

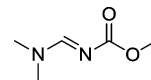
The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

Financial support from the National Institutes of Health (R01GM069990) is gratefully acknowledged.

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- (17) Catalyst does not dissolve fully even at 5 mol % loading.
- (18) Benzylideneacetone reacted very slowly under the standard conditions, giving the product in low yield (<10%).
- (19) Standard deprotection conditions (e.g., TsOH, TFA, LiBF₄, TMSBr, ZnBr₂, BF₃OEt₂, MgBr₂, TMSOTf, Amberlyst 15, PPTS, Sc(OTf)₃, Bi(OTf)₃, and ZrCl₄) gave low yields and racemization.
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- (25) The MOM-MAC reagent also reacted well with the free dihydroxyenone **18** (>90% ee).