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Stereoselective reaction of 2-carboxythioesters-1,3-dithiane with nitroalkenes: an organocatalytic strategy for the asymmetric addition of a glyoxylate anion equivalent†

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An efficient organocatalytic methodology has been developed to perform the stereoselective addition of 2-carboxythioesters-1,3-dithiane to nitroalkenes. Under mild reaction conditions γ -nitro- β -aryl- α -keto esters with up to 92% ee were obtained, realizing a formal catalytic stereoselective conjugate addition of the glyoxylate anion synthon. The reaction products are versatile starting materials for further synthetic transformations; for example, the simultaneous reduction of the nitro group and removal of the dithiane ring was accomplished, allowing the preparation of a GABA_B receptor agonist baclofen.

Introduction

Chiral α -keto esters are considered products of great importance, as starting materials for the preparation of highly functionalized compounds *via* synthetic manipulation of their functional groups.¹ They also find successful applications in biochemistry and are an integral part of several biologically active natural compounds.² Therefore, it is not surprising that the stereoselective synthesis of chiral α -keto esters is still the object of intense studies. Recently a novel approach was proposed by Johnson, who described a copper(II)-catalyzed aerobic oxidation reaction as a key step for the preparation of β -substituted α -keto esters, in a procedure where acetoacetate esters were used as glyoxylate anion synthons.³

However, most of the reported synthesis of chiral α -keto esters rely on the umpolung strategy that allows the introduc-

tion of a nucleophilic acyl anion synthon,⁴ *N*-heterocyclic carbenes are known for their ability to reverse the polarity of aldehydes, to generate acyl anion equivalents that have been widely employed in benzoin and Stetter-type reactions.⁵ More traditional methods involve the use of metalated amino-nitriles⁶ and 1,3-dithiane-protected glyoxylates,⁷ prepared by deprotonation with stoichiometric amounts of a strong base.⁸ It must be noted that all of these methods suffer from intrinsic limitations and problems (limited general applicability, low yields, modest stereoselectivity, and strongly depending on reactant substrates); in addition, examples of direct synthesis of enantiomerically enriched keto esters are still very rare, especially those related to the preparation of enantiopure β -aryl- α -keto esters, where the strong acidity of the proton in the β position represents a real issue for any successful stereoselective methodology.

With the goal to develop a metal-free catalytic strategy for the synthesis of chiral β -substituted- α -keto esters, we decided to study the addition of 1,3-dithiane-2-carboxy derivatives to nitroalkenes. Surprisingly, a catalytic stereoselective version of this powerful transformation seems to be unknown.⁹ Here we wish to report the first enantioselective organocatalytic conjugate addition of 2-carboxythioesters-1,3-dithiane to nitroalkenes.

At the beginning of our investigation the chemical reactivity of different 2-carboxyester 1,3-dithianes was investigated in the presence of a cinchona-derived bifunctional catalyst **A** (Scheme 1)¹⁰.

Commercially available 1,3-dithiane-2-ethylcarboxylate **2** in the presence of 20 mol% of quinine-thiourea derivative **A** did not react at all with nitrostyrene. The corresponding *S*-phenyl thioester **4** showed poor reactivity, affording the corresponding addition product **5** in low yields, even after prolonged reaction times, although in a remarkable 89% enantioselectivity (entries 2 and 3, Table 1). As expected 2-pyridylthioester **6** afforded adduct **7** in higher yields; unfortunately the reaction product turned out to be not very stable, thus preventing the determination of its enantiomeric excess. Finally the use of

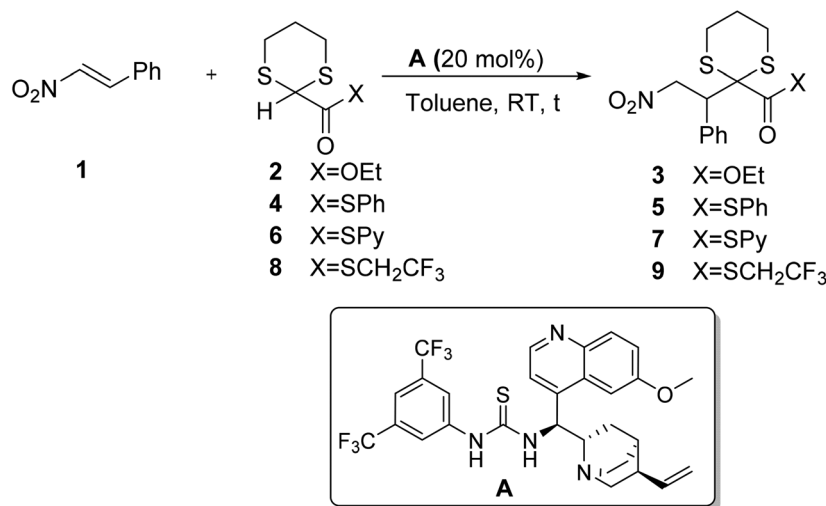
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† Electronic supplementary information (ESI) available: Experimental details for nitrostyrene derivatives synthesis, catalysts preparation and organocatalytic reactions. Characterization details for Michael addition products and Baclofen synthesis. Determination of absolute configuration of the reaction products.

¹H-NMR, ¹³C-NMR and ¹⁹F-NMR spectra and HPLC chromatograms on the chiral stationary phase of addition products. See DOI: 10.1039/c5ob00492f





Scheme 1 Preliminary studies of organocatalytic addition of dithianes to nitrostyrene.

Table 1 Chemical activity of 2-substituted 1,3-dithianes

Entry ^a	X	Product	<i>t</i> (h)	Yield ^b (%)	ee ^c (%)
1	OEt	3	24	—	—
2	SPh	5	24	16	89
3	SPh	5	72	25	87
4	SPy	7	24	41	n.d.
5	SCH ₂ CF ₃	9	24	71	87

^a Typical reaction conditions: 0.1 mmol of nitroalkene, 0.2 mmol of dithiane, 0.02 mmol of catalyst **A**, 1 mL of toluene. ^b Yields were determined after chromatographic purification. ^c Enantiomeric excess determined by HPLC on a chiral stationary phase.

2-*S*-trifluoroethyl carboxy-thioester-1,3-dithiane **8** gave the best results; after 24 hours at RT the stable conjugate addition product **9** was obtained in 71% yield and 87% ee.¹¹

The use of 1,1,1-trifluoroethyl thioester was documented for the first time by Barbas in an organocatalytic Michael reaction.¹² Later, our group took advantage of low *pK_a* of the protons in the α position of the carboxy group in the trifluoroethyl thioesters to realize the first organocatalytic direct aldol reaction between a thioester and an aromatic aldehyde.¹³ Due to its favorable reactivity profile (entry 5, Table 1), thioester **8** was selected as a reagent of choice and its addition to nitrostyrene was studied in the presence of different chiral bifunctional organocatalysts (Scheme 2).

In the model reaction performed in toluene for 24 hours at room temperature 2 mol eq. of thioester **8** were reacted with 1 mol eq. of nitrostyrene in the presence of 0.2 mol eq. of a catalyst. Based on the good results obtained with catalyst **A**, another successful thiourea-based bifunctional catalyst, Take-moto catalyst **B**,¹⁴ was tested; the product was isolated in higher yield but lower enantioselectivity than the cinchona-derived catalyst **A** (entries 1 and 2 of Table 2, 67% ee for catalyst **B** vs. 87% ee for **A**).

Therefore we turned our attention to the variation of different structural elements of quinine-derived catalyst **A**; for example with the cinchonidine derivative **C**, lacking the methoxy group on the quinolone ring, a little decrease in enantioselection was observed (75% vs. 87%). A modification of the same alkoxy group was deleterious for enantioselectivity (see entries 4 and 5, catalysts **D** and **E**). Compounds **F–I** clearly demonstrated the crucial role exerted by the thiourea moiety which cannot be replaced by alternative hydrogen bond donor groups. In addition, when squaramides **L–N**,¹⁵ derived from quinine or *trans*-diamine-cyclohexane, were tested, very low enantioselectivities were obtained.

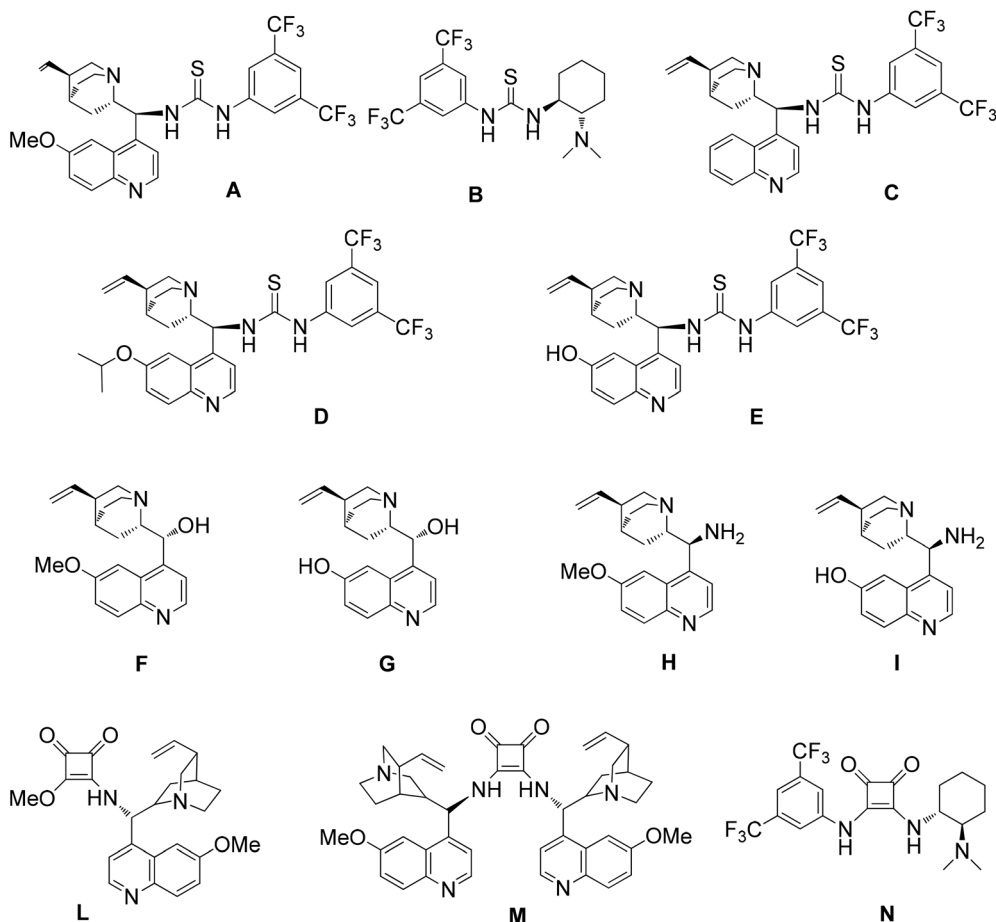
Once compound **A** was selected as a catalyst of choice, a few reaction parameters were optimized (Table 3).

The reaction tolerates different solvents, although only dichloromethane guaranteed the same level of ee as toluene (80% ee at RT). Lowering the reaction temperature to 0 °C had a positive effect, allowing isolating the product in 92% ee; lower temperatures did not give satisfactory results. It is noteworthy that the loading of the catalyst could be decreased to 10 mol% and even to 5 mol% with only a marginal loss of enantioselectivity, albeit with diminished chemical yield. The general applicability of the methodology was then investigated (Scheme 3, eqn (a)).

Differently substituted nitro alkenes were easily prepared¹¹ and reacted with dithiane **8** in the presence of catalyst **A** (Table 4).

The reaction works with nitrostyrenes substituted both in *ortho* and in *para* positions, with electron rich or poor substituents, leading to products with enantioselectivities ranging typically between 70% and 92%. Typically less reactive alkyl nitro alkenes¹⁶ did not react under the present conditions (entries 10 and 11); however, 4-nitro-1-phenyl butadiene reacted smoothly to afford product **17**, bearing a styryl moiety in 61% yield and 71% ee (eqn (b), Scheme 3).





Scheme 2 Chiral organocatalysts investigated in the 1,3-dithiane addition to nitrostyrene.

Table 2 Screening of chiral organocatalysts in the 1,3-dithiane addition to nitrostyrene

Entry ^a	Catalyst	Yield ^b (%)	ee ^c (%)
1	A	71	87
2	B	81	67
3	C	55	75
4	D	53	51
5	E	41	37
6	F	53	<5
7	G	55	31
8	H	73	37
9	I	15	<5
10	L	51	21
11	M	47	7
12	N	57	<5

^aTypical reaction conditions: 0.1 mmol of nitroalkene, 0.2 mmol of dithiane, 0.02 mmol of catalyst **A**, 1 mL of toluene. ^bYields were determined after chromatographic purification. ^cEnantiomeric excess determined by HPLC on a chiral stationary phase.

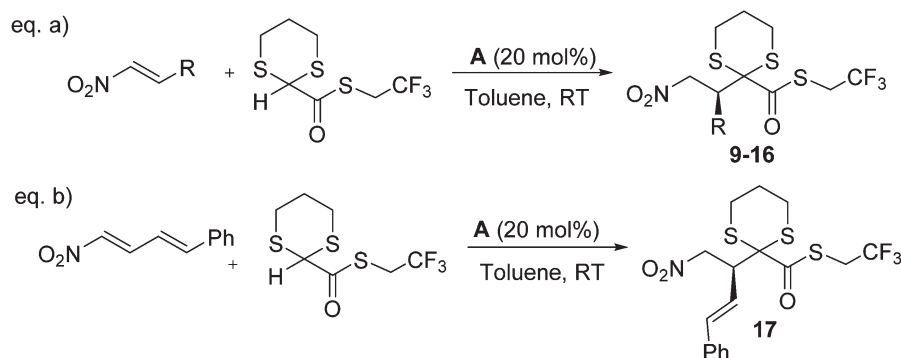
Table 3 Solvent screening for the organocatalyzed addition of dithiane **8** to nitrostyrene

Entry ^a	Solvent	Temp. (°C)	Yield ^b (%)	ee ^c (%)
1	Toluene	25	71	87
2	DCM	25	55	80
3	CH ₃ CN	25	80	41
4	Hexane	25	75	57
5	Et ₂ O	25	73	67
6	Toluene	0	60	92
7	Toluene	−20	61	66
8 ^d	Toluene	25	55	81
9 ^e	Toluene	25	47	80

^aTypical reaction conditions: 0.1 mmol of nitroalkene, 0.2 mmol of dithiane, 0.02 mmol of catalyst **A**, 1 mL of toluene. ^bYields were determined after chromatographic purification. ^cEnantiomeric excess determined by HPLC on a chiral stationary phase. ^dReaction run with 10 mol% of a catalyst. ^eReaction run with 5 mol% of a catalyst.

In order to establish the absolute configuration of the addition products, compound **11** was treated with powdered zinc and 6 M HCl¹⁷ to afford enantiomerically enriched lactam





Scheme 3 Organocatalytic 1,3-dithiane addition to different nitroalkenes.

Table 4 General applicability of the catalytic methodology

Entry ^a	R	Product	Yield ^b (%)	ee ^c (%)
1	Ph	9	71	87
2 ^d	Ph	9	60	92
3	4-CF ₃ Ph	10	41	53
4	4-ClPh	11	81	86
5	4-CH ₃ Ph	12	71	73
6	2-CH ₃ Ph	13	43	67
7	4-OCH ₃ Ph	14	75	73
8	2-OCH ₃ Ph	15	80	71
9	2-OAcPh	16	55	85
10	i-Pr	n.r.	—	—
11	CH ₂ CH ₂ Ph	n.r.	—	—

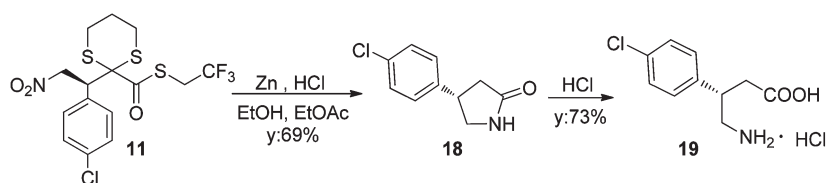
^a Typical reaction conditions: 0.1 mmol of nitroalkene, 0.2 mmol of dithiane, 0.02 mmol of catalyst **A**, 1 mL of toluene at 25 °C. ^b Yields were determined after chromatographic purification. ^c Enantiomeric excess determined by HPLC on a chiral stationary phase. ^d Reaction run at 0 °C.

18, which was determined to be in the (*S*) configuration by comparison of optical rotation power values. Indeed final hydrolysis with hydrochloridric acid afforded (*S*)-baclofen **19** (Scheme 4).¹⁸

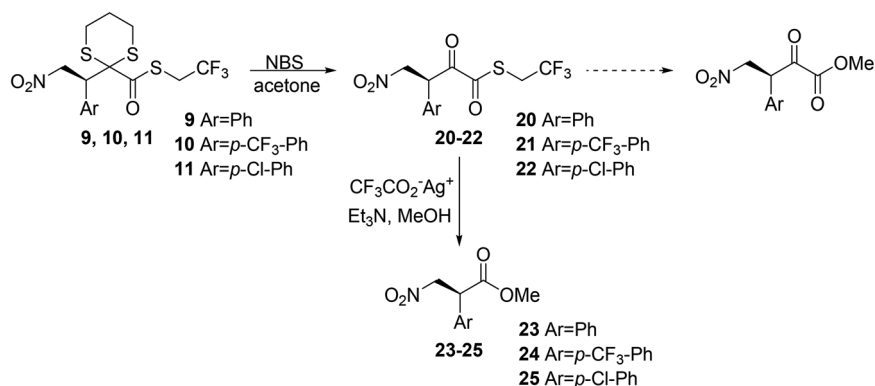
Therefore, the method offers a straightforward access to a direct precursor of baclofen-type derivatives, GABA_B receptor agonists used in the treatment of spasticity and alcohol dependence.¹⁹

Finally, conversion of the addition products into the corresponding α-ketothioesters was achieved by reaction with *N*-bromosuccinimide in acetone (Scheme 5).²⁰

For example, compounds **9–11** were quantitatively converted to α-ketothioesters **20–22**. Interestingly, upon treatment of **20–22** with silver trifluoroacetate, β-nitro esters **23–25** were obtained (Scheme 5).²¹



Scheme 4 Synthesis of baclofen.



Scheme 5 Synthesis of β-aryl-α-keto thioesters.



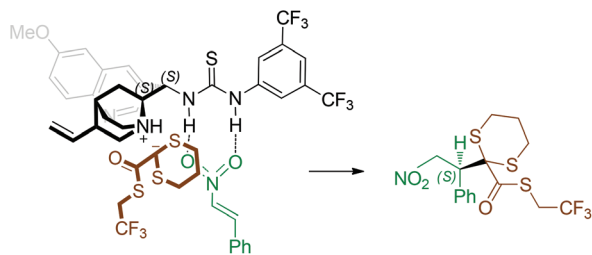


Fig. 1 Proposed stereoselection model.

The procedure afforded compounds **20–22** without affecting the stereochemical integrity of the starting materials. For instance, a sample of (*R*)-**9** (53% ee) was quantitatively transformed into α -ketothioester **20** through a 4 hour reaction with NBS in acetone at 0 °C, and then reacted with $\text{CF}_3\text{CO}_2\text{Ag}$ in methanol to give the known (*R*)- β -nitro methylester **23**²² in 70% yield, its absolute configuration reflecting that of the starting compound **9** without appreciable variation of the enantiomeric excess.²³

In proposing a stereoselection model, the coordination of both reaction partners to the bifunctional catalyst may be envisaged, the substrate nitro group being engaged in hydrogen bonds by the thiourea moiety, and the charged quinuclidine nitrogen possibly forming an ion pair with the nucleophile (Fig. 1).

Accordingly, with the quinine-derived catalyst, the thioester attack occurs on the nitroalkene *Si* face, affording the experimentally observed (*S*)-configured product.

In conclusion, the enantioselective organocatalyzed conjugate addition of a newly activated thioester acting as an acyl anion mimic has been performed. The use of chiral bifunctional catalysts to control the stereochemical outcome through a hydrogen bond network imposed by the thiourea moiety afforded good yields and enantioselectivities up to 92%. This methodology represents an entry to highly functionalized, enantiomerically enriched products, such as γ -nitro- β -aryl- α -keto thioesters, valuable precursors of a wide variety of chiral organic compounds.

Acknowledgements

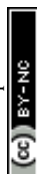
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