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Organocatalytic sulfa-Michael/aldol cascade: constructing functionalized 2,5-dihydrothiophenes bearing a quaternary carbon stereocenter†

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A practical sulfa-Michael/aldol cascade reaction of 1,4-dithiane-2,5-diol and α -aryl- β -nitroacrylates has been developed, which allows efficient access to functionalized 2,5-dihydrothiophenes bearing a quaternary carbon stereocenter in moderate to good yields with high enantioselectivities.

Among the various classes of heterocycles, members of the thiophene family have received particular attention from the chemical community because of their widespread occurrence as ubiquitous motifs in natural products, pharmaceuticals, agrochemicals as well as materials.¹ In this context, the 2,5-dihydrothiophene ring is a common structural feature of many bioactive compounds and a potential intermediate for various synthetic applications.² Over the decades, only a few examples of the assembly of optically active 2,5-dihydrothiophenes have been documented.³ For instance, the Spino^{3b} group successfully prepared non-racemic dihydrothiophenes using an efficient chiral auxiliary. The first gold-catalyzed cycloisomerization of α -hydroxyallenes to 2,5-dihydrothiophenes was reported by Krause^{3c} and co-workers. Then in 2010, the Xu^{3e} group developed a highly stereoselective domino thia-Michael/aldol reaction between 1,4-dithiane-2,5-diol and α,β -unsaturated aldehyde catalyzed by a chiral diphenylproline TMS ether, which provided a new avenue for the synthesis of functionalized 2,5-dihydrothiophenes.

Quaternary carbon stereocenters are often contained in natural products and pharmaceuticals.⁴ Compared with the chiral pool synthesis,⁵ the procedure of chiral materials or catalysts to construct such sterically congested stereogenic centers is more challenging because of the difficulty of orbital overlap.⁶ To date, a lot of progresses have been made in the construction of chiral quaternary carbon centers in cyclic compounds,⁷ which are greatly accelerated by the advancement of transition metal catalysis,⁸ and organocatalysis,⁹ including methods beyond radical initiation.¹⁰ However, only few examples are about the construction of a quaternary carbon in 2,5-dihydrothiophene ring.¹¹ Inspired by the previous work of the Xu group,^{3e} we describe herein an elegant organocatalytic

asymmetric cascade sulfa-Michael/aldol reaction, providing a convenient way for the synthesis of 2,5-dihydrothiophenes bearing a chiral quaternary carbon center.

Results and discussion

At the outset of our pursuit, a series of organocatalysts **I–VII** were examined in the cascade reaction of 1,4-dithiane-2,5-diol **1a** and α -phenyl- β -nitroacrylate **2a** in chloroform at room temperature (Fig. 1). The desired 2,5-dihydrothiophene **3a** could be successfully obtained in the presence of cinchona alkaloid **I** within 24 hours in 40% yield with only 12% ee (Table 1, entry 1). A substantial increase of yield and enantioselectivity were observed when thiourea-tertiary amine catalysts **II–V** were employed (Table 1, entries 2–5). Further optimization indicated that cinchona-based squaramide **VI** and **VII** with stronger H-bonding donors gave the products with comparable results

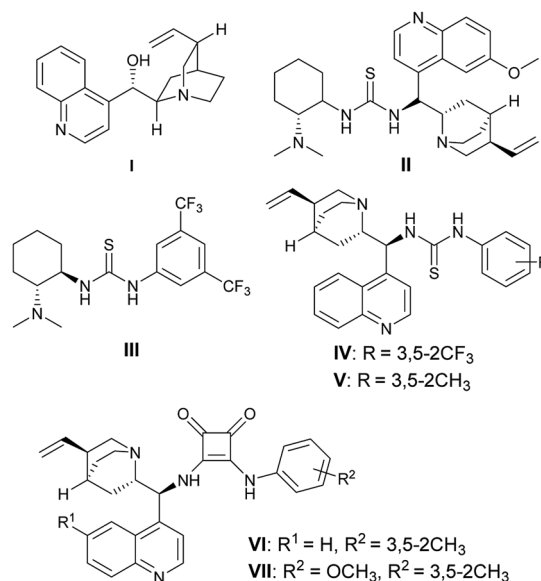
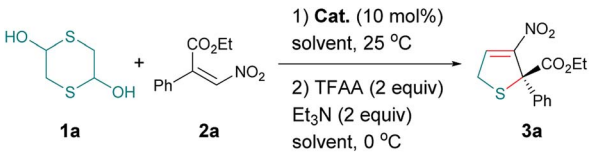


Fig. 1 The organocatalysts tested in this study.

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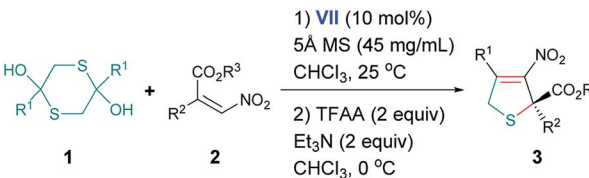
Table 1 Optimization of the reaction conditions^a


Entry	Catalyst	Solvent	Time (h)	Yield ^b (%)	ee ^c (%)
1	I	CHCl ₃	24	40	12
2	II	CHCl ₃	24	55	33
3	III	CHCl ₃	24	62	−56
4	IV	CHCl ₃	24	65	45
5	V	CHCl ₃	24	50	66
6	VI	CHCl ₃	24	66	70
7	VII	CHCl ₃	24	69	71
8	VII	CH ₂ Cl ₂	24	64	60
9	VII	Toluene	24	56	62
10	VII	THF	24	35	48
11	VII	Et ₂ O	24	68	66
12	VII	CH ₃ OH	24	44	3
13	VII	CH ₃ CN	24	43	12
14 ^d	VII	CHCl ₃	24	63	73
15 ^e	VII	CHCl ₃	24	68	76
16 ^f	VII	CHCl ₃	24	70	77
17 ^{f,g}	VII	CHCl ₃	55	68	73
18 ^{f,h}	VII	CHCl ₃	120	Trace	ND

^a Reaction conditions (unless otherwise stated): **1a** (0.18 mmol), **2a** (0.3 mmol), catalyst (10 mol%), solvent (2 mL), $T_1 = 25\text{ }^\circ\text{C}$; then trifluoroacetic anhydride (TFAA, 0.6 mmol), Et₃N (0.6 mmol), $T_2 = 0\text{ }^\circ\text{C}$. ^b Isolated yield based on **2a**. ^c Determined by chiral HPLC analysis. ^d MgSO₄ (90 mg) was added. ^e 4 Å MS (90 mg) was added. ^f 5 Å MS (90 mg) was added. ^g $T_1 = 15\text{ }^\circ\text{C}$. ^h $T_1 = 0\text{ }^\circ\text{C}$, ND = not determined.

(Table 1, entries 6 and 7), then we selected **VII** as the optimal catalyst. Next, changing chloroform to other solvents resulted in an obvious decrease both in yield and stereoselectivity (Table 1, entries 8–13). To our delight, the addition of MgSO₄ or molecular sieve (MS) into the reaction mixture could bring out a slight increase of the enantioselectivity, while the 5 Å MS gave the best results (Table 1, entry 16). We also tried to set up the reaction under a lower temperature, the cycloadduct **3a** could maintain the yield and enantioselectivity within a longer time at $15\text{ }^\circ\text{C}$ (Table 1, entry 17). When the temperature was further decreased to $0\text{ }^\circ\text{C}$, the reaction became very slow, and we only observed trace amount of product after 120 hours (Table 1, entry 18).

With the optimal conditions in hand, the cascade reaction of 1,4-dithiane-2,5-diol and diverse α -aryl- β -nitroacrylates was investigated. The results were summarized in Table 2. We found that α -phenyl- β -nitroacrylates with different ester groups, including benzyl, isopropyl, *tert*-butyl, could react smoothly to give the desired chiral dihydrothiophenes in 67–73% yields with 66–83% ee (entries 1–3), while bulky *tert*-butyl group gave the best results (entry 3). The position and nature of substituents on the α -phenyl ring had slight influence on the stereocontrol, 65–75% yields and 82–86% ee were observed (entries 4–9). The reaction was also applicable to α -naphthyl and furanyl nitroacrylates, producing the desired adducts with 83% and 85% ee,

Table 2 Scope for the synthesis of 2,5-dihydrothiophenes bearing a quaternary carbon stereocenter^a


Entry	Product	R ¹	R ²	R ³	t (h)	Yield ^b (%)	ee ^c (%)
1	3b	H	Ph	Bn	24	68	66
2	3c	H	Ph	<i>i</i> -Pr	24	67	80
3	3d	H	Ph	<i>t</i> -Bu	24	73	83
4	3e	H	4-CH ₃ C ₆ H ₄	<i>t</i> -Bu	36	65	86
5	3f	H	4-OCH ₃ C ₆ H ₄	<i>t</i> -Bu	24	71	82
6	3g	H	4-FC ₆ H ₄	<i>t</i> -Bu	24	68	85
7	3h	H	4-ClC ₆ H ₄	<i>t</i> -Bu	24	75	86
8	3i	H	3-OCH ₃ C ₆ H ₄	<i>t</i> -Bu	24	74	84
9	3j	H	3,5-2CH ₃ C ₆ H ₃	<i>t</i> -Bu	30	67	83
10	3k	H	2-Naphthyl	<i>t</i> -Bu	24	72	86
11	3l	H	2-Furanyl	<i>t</i> -Bu	24	62	83
12	3m	H	<i>i</i> -Pr	<i>t</i> -Bu	24	69	85
13 ^d	3n	CO ₂ Et	Ph	<i>t</i> -Bu	36	65	87
14 ^d	3o	CO ₂ Et	3,5-2CH ₃ C ₆ H ₃	<i>t</i> -Bu	36	64	88

^a Reaction conditions (unless otherwise stated): **1** (0.18 mmol), **2** (0.3 mmol), **VII** (10 mol%), 5 Å MS (90 mg), CHCl₃ (2 mL), $T_1 = 25\text{ }^\circ\text{C}$; then TFAA (0.6 mmol), Et₃N (0.6 mmol), $T_2 = 0\text{ }^\circ\text{C}$. ^b Isolated yield based on **2**. ^c Determined by chiral HPLC analysis. ^d $T_1 = -40\text{ }^\circ\text{C}$.

respectively, the yields were also satisfactory (entries 10 and 11). Gratifyingly, branched as well as aliphatic nitroacrylate is equally suitable substrate (entry 12). In addition, we tried the reaction of 2,5-dihydroxy-1,4-dithiane-2,5-dicarboxylate with α -aryl- β -nitroacrylates **2n** and **2o**, which exhibited promising results, leading to 64–65% yields and 87–88% ee at $-40\text{ }^\circ\text{C}$ within 36 hours (entries 13 and 14).

To determine the absolute configuration of the products, we carried the cascade reaction in a large amount, and a single crystal of the optically pure product **3a** was obtained after crystallization,¹² which was then determined to be *S* by X-ray crystallographic analysis.¹³ Based on the experimental result and previous studies,¹⁴ the transition state is shown in Scheme 1. Mechanistically, the β -nitroacrylate was activated by the hydrogen bonds of the squaramide, while the tertiary amine of the catalyst would enhance the nucleophilicity of mercaptoacetaldehyde based on its basicity, the nucleophile attacked the Michael acceptor from the *Si* face, which favoured the formation of the C2 (*S*) stereogenic center. It is worth noting that the prediction of possible interaction in the transition state was in agreement with the DFT calculation (Fig. 2).¹⁵

Because of the highly functionalization of the product, we also tried the synthetic transformation of 2,5-dihydrothiophene. Taking the reaction of **1a** and **2d** as example, when the addition and dehydration steps completed at the optimal conditions, the benzotriazole was added directly to the mixture as a nucleophile. Interestingly, the adduct **4** was



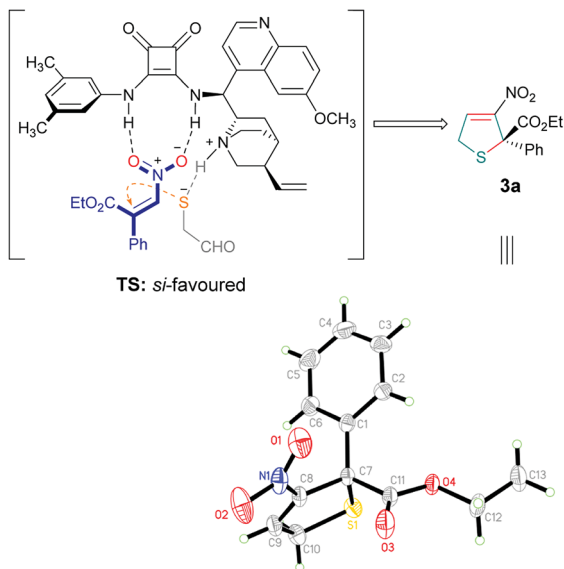
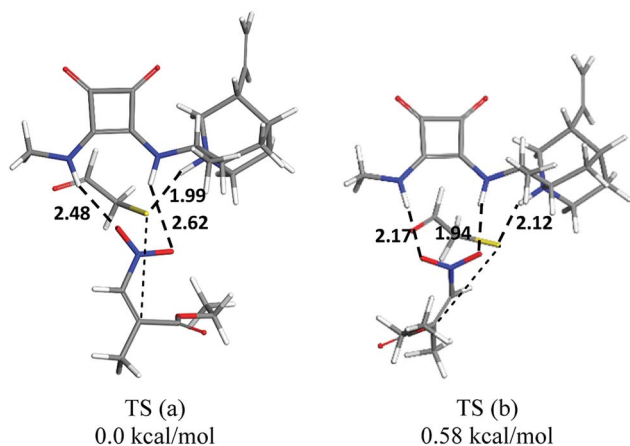
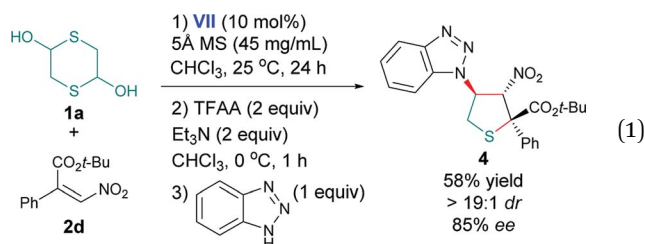
Scheme 1 Proposed transition state and the X-ray structure of **3a**.

Fig. 2 Optimized structures and free energy gap of the transition states.

obtained with moderate yield and high stereoselectivities (eqn (1), 58% yield, >19 : 1 dr, 85% ee).¹⁶



Conclusions

In summary, we have developed a practical sulfa-Michael/aldol cascade reaction of 1,4-dithiane-2,5-diol and α -aryl- β -

nitroacrylates, which allows an efficient access to functionalized 2,5-dihydrothiophenes bearing a quaternary carbon center in moderate to good yields with high enantioselectivities. Moreover, the products can be easily derivated to more complex compound. Further application of this strategy to the enantioselective synthesis of other heterocyclic systems is underway in our laboratory.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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