

## REVIEW

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# Recent advances in organocatalytic asymmetric oxa-Michael addition triggered cascade reactions

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The oxa-Michael cascade reaction, as an important part of the Michael reaction, has also been significantly developed over the recent decade. This is because the problem of the reactivity and selectivity of the addition reactions of oxygen nucleophiles to conjugated systems has been solved by different organocatalysts and reaction conditions. Therefore, using various efficient strategies, many novel and potentially bioactive chiral compounds with excellent yields and stereoselectivities have been synthesized. In this review, we summarize the recent advances in organocatalytic asymmetric oxa-Michael addition triggered cascade reactions for the stereoselective synthesis of heterocyclic compounds.

## 1 Introduction

The Michael addition reaction is one of the most powerful and valuable tools for the formation of C–X (X = C, O, N, S) bonds, which plays a vital role in organic synthetic chemistry.<sup>1–3</sup> It is

one of the most typical reactions between a nucleophile called a Michael donor and an activated Michael acceptor, and a huge number of optically functional structures and natural products have been prepared by this important transformation over the years. Meanwhile, the nucleophilicity and reactivity of alcoholic hydroxyl are relatively poor, but the oxa-Michael reaction, as an important part of the Michael reaction for the construction of the C–O bond, has also received considerably

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more attention from researchers after solving the problems involved in recent years.<sup>4,5</sup>

Small organic molecules capable of catalyzing reactions have been known for many decades, with the result that asymmetric organocatalysis has become a hot field of research. Moreover, because of the high efficiency, selectivity, low cost, environmental benignness and other characteristics of organocatalysts, more and more research groups are engaged in the development of organocatalytic asymmetric reactions to obtain the corresponding enantiomer-enriched products. In past decades, the rapid development of organocatalysis has greatly changed the profile of asymmetric catalysis.<sup>6,7</sup> Many types of chiral organocatalysts, such as chiral prolinol analogues, chiral bifunctional squaramides and thioureas, and other chiral compounds, have been applied to oxa-Michael cascade reactions to synthesize many diverse and versatile oxygen-containing heterocyclic, bicyclic, spiro, and polycyclic structures with multiple stereocenters. In addition, these reaction protocols greatly decrease the number of laboratory operations, save time, cost and work steps,<sup>8</sup> and have a high synthetic efficiency, thereby substantially broadening the scope of oxa-Michael reactions.

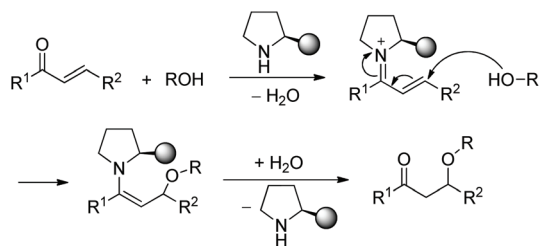
Therefore, this review aims to illustrate the developments in asymmetric oxa-Michael addition triggered cascade reactions over the last decade. Particular attention will be paid to organocatalytic reaction strategies and their applications in the synthesis of chiral oxygen-containing heterocycles.

## 2 Organocatalytic enantioselective oxa-Michael initiated cascade reactions

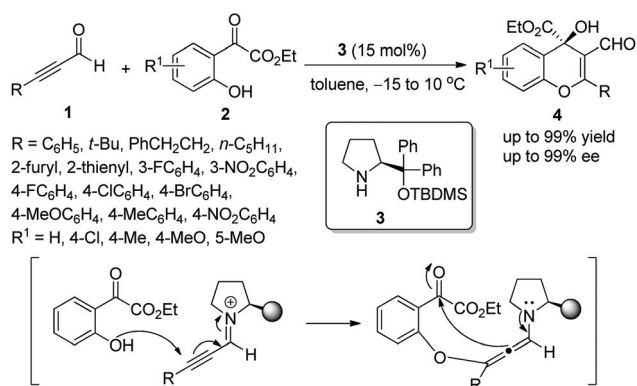
### 2.1 Reactions catalyzed by chiral prolinol analogues

In this sort of catalytic process, the chiral secondary amine will activate the electrophile and generate an iminium-enamine intermediate, which then undergoes conjugate addition and hydrolysis steps to release the product and amine catalyst (Scheme 1). In general, some additives will be used to facilitate this process. It turns out that the proline-derived chiral catalysts have performed with outstanding results in oxa-Michael addition cascade reactions.<sup>9</sup>

**2.1.1 Oxa-Michael cascade reactions for the synthesis of chiral chromenes.** In 2010, Wang's group<sup>10</sup> explored a novel



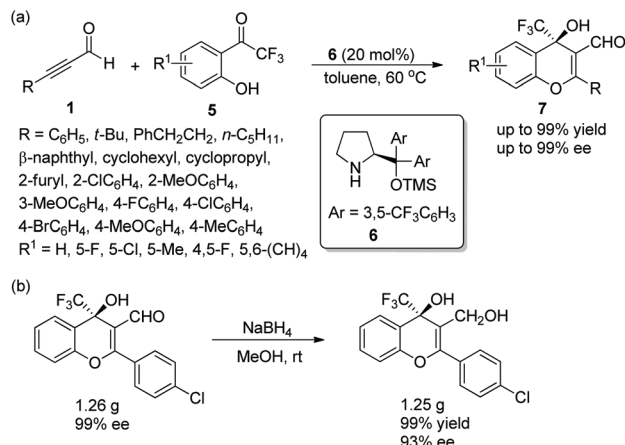
**Scheme 1** Chiral prolinol analogue catalytic general mechanism of the oxa-Michael addition reaction.



**Scheme 2** Oxa-Michael/aldol cascade reaction for the synthesis of chromenes.

mode of generating diverse 4*H*-chromenes **4** bearing a versatile  $\alpha$ -hydroxy carboxylate moiety by using chiral diarylprolinol TBDMS ether **3** as a catalyst. The oxa-Michael/aldol cascade reactions of alkynals **1** with *o*-hydroxyacetophenone esters **2** tolerated a wide range of substrates and afforded the desired products with excellent yields (up to 99%) and enantioselectivities (up to 99% ee) under mild reaction conditions (Scheme 2). It was noteworthy in the scope of substrates that the less reactive alkyl-substituted alkynals ( $R = \text{PhCH}_2\text{CH}_2$  or  $n\text{-C}_5\text{H}_{11}$ ) also could take part effectively in this cascade reaction and provided corresponding products with satisfactory enantioselectivities (97 and 99% ee, respectively), even though this process needed more reaction time and a higher catalyst loading (25 mol%) than other aryl substituents. Additionally, when the substituent of the aromatic structures was electron-withdrawing Cl, the high enantioselectivity would be ensured by using quinine as a cocatalyst which possibly interacted with the carbonyl moiety of the ketoester. In this cascade process, the iminium-allenamine activation mode is a quite efficient way to generate functionalized chromenes and successfully expand the scope of the organocatalytic oxa-Michael-aldol cascade sequence.

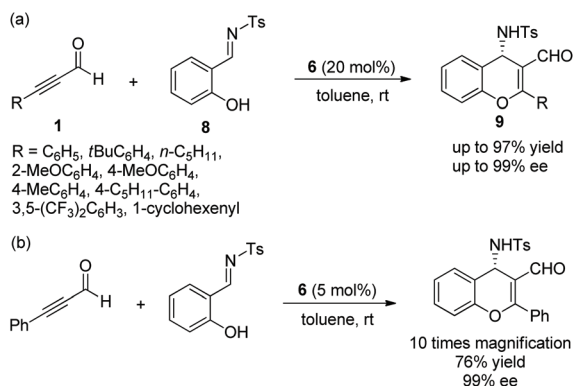
In consideration of the special properties of trifluoromethylated compounds in the pharmaceutical chemistry, it is very important to synthesize biologically active derivatives containing trifluoromethyl substituents. Then, there was the significant work in which Huang and coworker<sup>11</sup> first synthesized chiral 4*H*-chromenes **7** bearing a trifluoromethyl group in 2015. The alkynals **1** and *o*-hydroxy trifluoroacetophenones **5** underwent an oxa-Michael/aldol cascade process catalyzed by catalyst **6** in toluene at 60 °C, when the desired products **7** were obtained with high yields (up to 99%) and enantioselectivities (up to 99% ee) (Scheme 3a). In the scope of the reaction, both electron-donating and electron-withdrawing group substituted aromatic alkynals were revealed to be efficient reactants and afforded satisfactory results, while furyl-substituted alkynal showed lower yield (40%) and enantioselectivity (84% ee) than other groups. The position of the substituents on the aromatic ring of aromatic alkynals had an



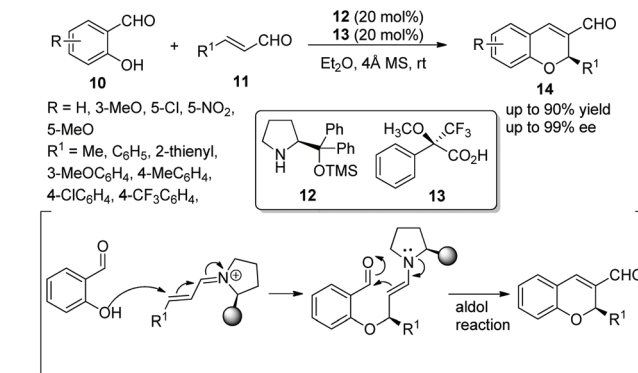
**Scheme 3** Oxa-Michael/aldol cascade reaction for the synthesis of 4*H*-chromenes bearing a trifluoromethyl and the derivatization of chiral chromene.

effect on the reaction: for example, *meta*-substituted alkyne afforded better yield and enantioselectivity than *ortho*-substituted alkyne; and electron-donating substituents on the aromatic ring of **5** provided better reactivity than electron-withdrawing substituents. Furthermore, the reaction product could be further reduced to other valuable building blocks on a gram-scale without sacrificing reaction yield or enantioselectivity (Scheme 3b).

Alemán and co-workers<sup>12</sup> also reported a similar method for the highly enantioselective synthesis of 4-amino-4*H*-chromenes **9** in 2010. Alkynes **1** and salicyl *N*-tosylimine **8**, as the starting materials, could undergo an oxa-Michael/aza-Baylis-Hillman (aza-BH) cascade process catalyzed by (*S*)-diarylprolinol TMS ether **6** to generate the corresponding products with excellent yields (up to 97%) and enantioselectivities (up to 99% ee) in toluene at room temperature (Scheme 4a). In the scope of substrates, almost all substituents maintained high stereoselectivities; compared with aliphatic substituents, aromatic substituents afforded corresponding products with better yields; a *para*-substituent on the aromatic ring provided a better result than an *ortho*-substituent. However, the reactiv-



**Scheme 4** Oxa-Michael/aza-BH cascade reaction for the synthesis of 4-amino-4*H*-chromenes and the large-scale preparation reaction.

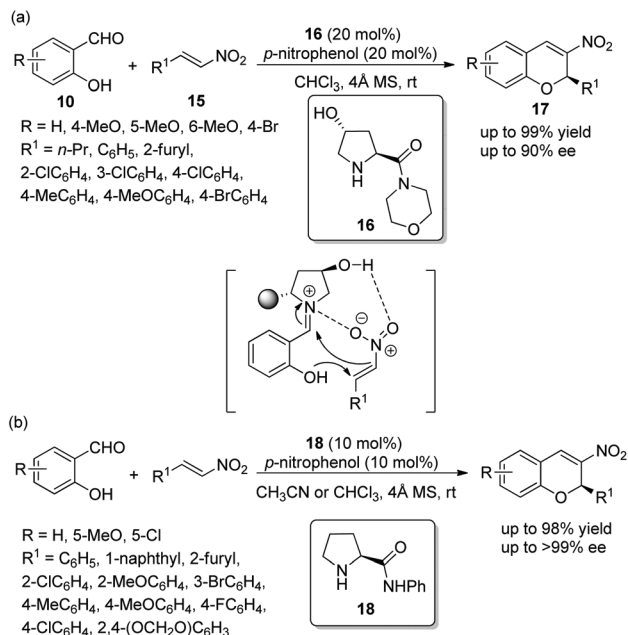


**Scheme 5** Oxa-Michael/aldol cascade reaction for the synthesis of chromene derivatives.

ity of the strong electron-poor aromatic ring 3,5-(CF<sub>3</sub>)<sub>2</sub>C<sub>6</sub>H<sub>3</sub> of alkyne was substantially decreased (only 55% yield), even when the catalyst loading had been increased to 40 mol% and the reaction time had been prolonged. Additionally, in the large-scale reaction, this process provided the corresponding product in moderate yield without losing enantiopurity (Scheme 4b).

Xu's group<sup>13</sup> investigated an oxa-Michael/aldol cascade reaction for the synthesis of chiral 2*H*-chromenes catalyzed by a chiral amine/chiral acid organocatalytic system in 2009. Salicylic aldehydes **10** and α,β-unsaturated aldehydes **11** in diethyl ether at room temperature, under the catalysis of catalyst **12** and (*S*)-Mosher's acid **13**, afforded the desired products **14** in good yields (up to 90%) with excellent enantioselectivities (up to 99% ee) (Scheme 5). Compared with a single organocatalyst, the chiral amine/chiral acid organocatalytic system efficiently increased the enantioselectivity under the same conditions. In the scope of substrates, compared with aliphatic-substituted unsaturated aldehydes, aromatic-substituted unsaturated aldehydes provided the corresponding products with better yields and enantioselectivities. The electronic property and position of the substituents on the aromatic ring of salicylaldehydes had an effect on the yields and enantioselectivities of the products: for example, compared with 5-methoxy and 5-nitro groups, the electron-donating substituent 3-methoxy provided the corresponding product with the best yield and enantiopurity.

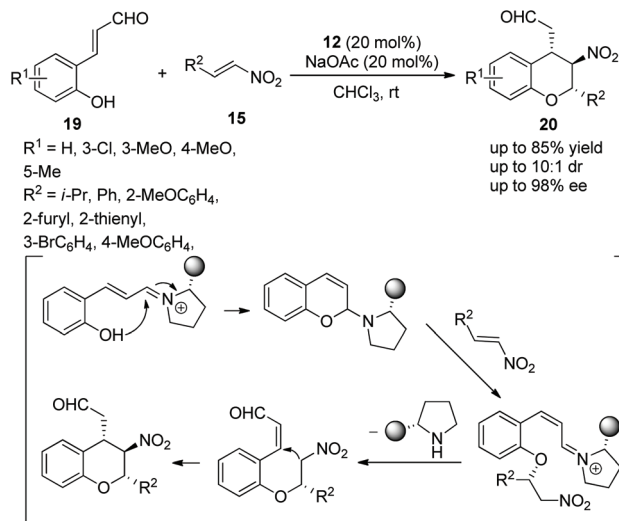
An efficient method for the synthesis of chiral chromenes *via* an oxa-Michael/Henry cascade reaction was reported by Chen's group<sup>14</sup> in 2013. In this cascade procedure, salicylaldehydes **10** and nitroalkenes **15** were employed as starting materials catalyzed by a 4-hydroxy-*L*-proline derived chiral aminocatalyst **16** and the additive *p*-nitrophenol, which afforded the desired chromenes **17** in excellent yields (up to 99%) with high enantioselectivities (up to 90% ee) under mild conditions (Scheme 6a). In the scope of substrates, the reactivity of the electron-donating substituents on the aromatic ring was better than that of electron-withdrawing substituents. The position of the substituents on the aromatic ring had a significant influence on the enantioselectivity of the desired products: for



**Scheme 6** Oxa-Michael/Henry cascade reaction for the synthesis of chromene derivatives.

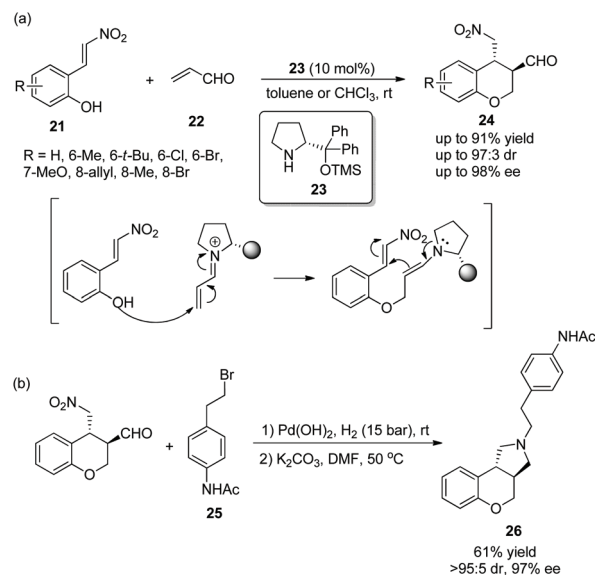
example, 4-methoxy-substituted salicylaldehyde only afforded the desired product with 54% ee, but 5-methoxy-substituted salicylaldehyde provided the corresponding product with 89% ee. Additionally, it should be noted that, besides the iminium ion, the formation of hydrogen-bonding also took part in the transition state of the cascade process. A few years later, similar work was performed by Bez's group.<sup>15</sup> Another catalyst **18** was employed in this cascade process (Scheme 6b). In general, in the substrate scope of this cascade reaction, regardless of electron-withdrawing or electron-donating substituents on the  $\beta$ -nitrostyrene derivatives, all of them had little effect on the yield or enantioselectivity of the products. However, it should be noted that the solvent still affected the result under optimal conditions. For example, the reaction of salicylaldehyde with *p*-fluoro- $\beta$ -nitrostyrene afforded the corresponding product with 86% ee in CHCl<sub>3</sub> and 99% ee in acetonitrile.

**2.1.2 Oxa-Michael/Michael cascade reaction for the synthesis of chiral chromans.** Wang's group<sup>16</sup> developed a valuable and efficient strategy for the highly enantioselective synthesis of chiral chromans *via* an oxa-Michael/Michael cascade reaction in 2009. 2-Hydroxy cinnamaldehydes **19** and nitroolefins **15** catalyzed by chiral (*S*)-diarylprolinol TMS ether **12** afforded the desired products **20** in good yields (up to 85%) with excellent enantioselectivities (up to 98% ee) in chloroform at room temperature (Scheme 7). In the scope of substrates, all the substituents, regardless of whether they were electron-rich or electron-poor groups, provided the corresponding products with excellent enantiopurity; the reactivity of unsubstituted 2-hydroxy cinnamaldehyde was better than 2-hydroxy cinnamaldehyde bearing substituents; and compared with alkyl-substituted nitroolefins, aryl-substituted nitroolefins gave a better result.



**Scheme 7** Oxa-Michael/Michael cascade reaction for the synthesis of trisubstituted chromans.

A short and convenient strategy for the synthesis of 3,4-di-substituted chromans *via* an oxa-Michael/Michael cascade reaction was reported by Enders' group<sup>17</sup> in 2012. Nitrovinylphenols **21** and acrolein **22** were employed to undergo the cascade process catalyzed by (*R*)-diphenylprolinol TMS ether **23** and provided the desired products **24** in good yields (up to 91%) with great stereoselectivities (up to 97:3 dr and 98% ee) in toluene or chloroform at room temperature (Scheme 8a). The reason why there were two kinds of solvents is that some substrates **21** bearing substituents, such as 6-Me, 6-Br, 8-Br, or 8-allyl, have poor solubility in toluene and thus the corresponding reactions were conducted in chloroform.



**Scheme 8** Oxa-Michael/Michael cascade reaction for the synthesis of 3,4-*trans*-disubstituted chromans and the derivatization of chiral chroman.



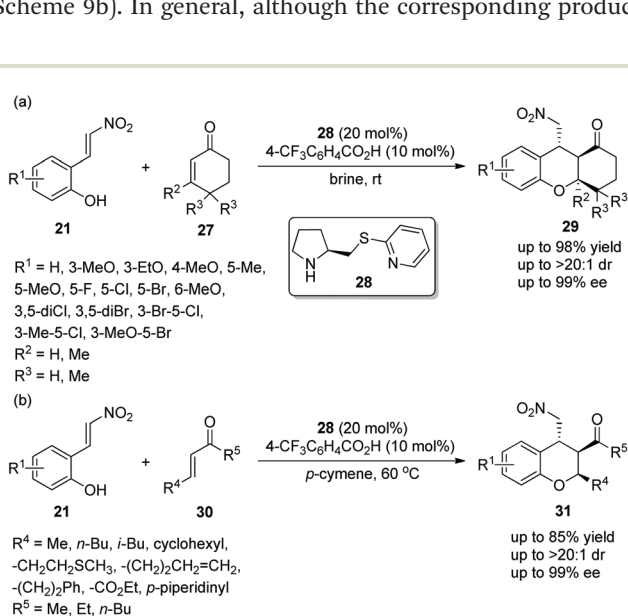
Then, in the limited scope of substrates, all of them afforded the corresponding products with excellent stereoselectivities, except the nitrovinylphenol-bearing an electron-rich methoxy-substituent which did not generate the desired product; electron-donating substituents showed better reactivity than electron-withdrawing substituents; and compared with other positions, the reactivity of the methyl substituent at the C-7 position was the best. Furthermore, through a two-step reductive amination/*N*-alkylation sequence, they successfully transformed the cascade product to *trans*-*N*-alkylated benzopyrano [3,4-*c*]pyrrolidine **26**, an antagonist of the dopamine-D3-receptor and potential antipsychotic medicament, without losing much enantiopurity (Scheme 8b).

In view of the good result reported by Enders' group, similar work for the construction of functionalized tetrahydroxanthenones and chromans was finished by Xu's group<sup>18</sup> in 2014. Under the catalysis of prolinol thioether catalyst **28** and 4-trifluoromethylbenzoic acid additive, nitrovinylphenols **21** and cyclohexenones **27** gave the products tetrahydroxanthenones **29** with excellent yields (up to 98%) and stereoselectivities (up to >20:1 dr and 99% ee) in brine at room temperature (Scheme 9a). Interestingly, while maintaining the stereoselectivity of this cascade reaction, brine took a shorter reaction time than organic solvents. In the scope of substrates, the electronic property and position of the substituents had little effect on the stereoselectivities of the reaction, and all of them afforded the corresponding products with excellent stereoselectivities. Compared with other positions on the aromatic ring, the reactivity of a methoxy substituent at the C-6 position was lowest, and the reason could be that methoxy is too close to the carbon-carbon double bond to affect the electrophilic site of the reaction. Furthermore, nitrovinylphenols **21** and linear unsaturated ketones **30** were also employed to generate chiral chromans **31** *via* the same catalytic cascade process (Scheme 9b). In general, although the corresponding products

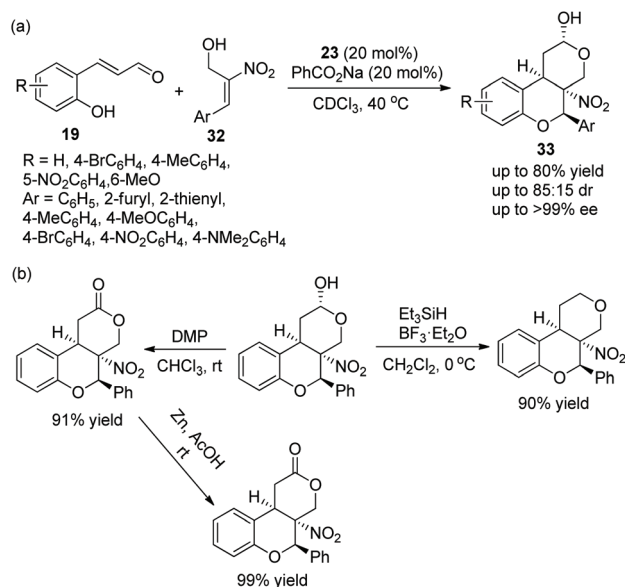
were obtained with excellent stereoselectivities, the reactivity of linear unsaturated ketones **30** with nitrovinylphenols **21** in *p*-cymene was not as good as the reactivity of cyclohexenones **27** with nitrovinylphenols **21** in brine.

Tricyclic chroman, as an important structure, has attracted the attention of many researchers, especially in the synthesis of enantiomerically enriched chroman derivatives with this structure. A study for synthesizing functionalized tricyclic chromans with four stereocenters *via* an oxa-Michael/Michael/nucleophilic ring-closing cascade process was reported by Jørgensen's group<sup>19</sup> in 2014. They applied *o*-hydroxycinnamic aldehydes **19** and 2-nitroallylic alcohols **32** to this cascade process and, at the same time, PhCO<sub>2</sub>Na was added to improve the reactivity and yield. Under the catalysis of chiral diarylprolinol TMS ether **23**, the cycloadducts **33** were obtained in good yields (up to 80%) with excellent enantioselectivities (up to >99% ee) in CDCl<sub>3</sub> at 40 °C (Scheme 10a). In the substrate scope of this cascade reaction, heteroaryl-substituted 2-nitroallylic alcohols afforded the corresponding products with the best enantioselectivity. The electron-donating groups on the phenyl ring of the *o*-hydroxycinnamic aldehydes gave better yields and stereoselectivities than the electron-withdrawing groups. Furthermore, they demonstrated that the cascade product could be selectively transformed into other structures, such as the dehydroxylation reaction of chromans with Et<sub>3</sub>SiH and BF<sub>3</sub>·Et<sub>2</sub>O, the oxidation of the hydroxyl group under Dess–Martin conditions and further selective reduction of the nitro group. Each reaction provided the corresponding product with high isolated yield (Scheme 10b).

In 2014, Wang's group<sup>20</sup> conceived an effective strategy to synthesize tricyclic chroman derivatives under the catalysis of chiral diarylprolinol TMS ether **12** under adjustable reaction

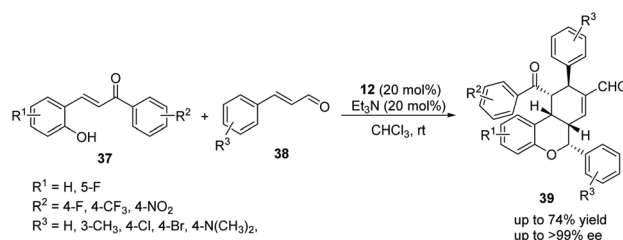


**Scheme 9** Oxa-Michael/Michael cascade reaction for the synthesis of tetrahydroxanthenones and chromans.



**Scheme 10** Oxa-Michael/Michael/nucleophilic ring-closing cascade reaction for the synthesis of chromans and the derivatizations of chiral chroman.

conditions. What is noteworthy is that the reaction temperature is a vitally important factor. When the cascade reaction of (*E*)-2-hydroxyaryl-2-oxa-but-3-enoates **34** with enals **11** catalyzed by catalyst **12** occurred in PhCl at  $-5^{\circ}\text{C}$ , the corresponding products **35** were obtained in high yields with excellent stereoselectivities, but when the reaction temperature was continuously elevated to  $30^{\circ}\text{C}$ , products **36** would be generated. Obviously, if the cascade reaction catalyzed by catalyst **12** was performed in PhCl at  $30^{\circ}\text{C}$ , the chroman derivatives **36**, with great yields and stereoselectivities, would be synthesized exclusively (Scheme 11a). It is amazing that, irrespective of whether there are electron-withdrawing or electron-donating groups in the substrate scope of the cascade reaction, all of them offered the corresponding products with excellent stereoselectivities ( $>10:1$  dr and  $>98\%$  ee); electron-withdrawing substituents on the aromatic ring of **34** showed better reactivity than electron-donating substituents; compared with aliphatic-substituted enals, aromatic-substituted enals displayed better reactivity. Furthermore, it was demonstrated that the chiral product could be converted to the corresponding hydrazine by reaction with *p*-toluenesulfonylhydrazide in a mixed solvent of metha-

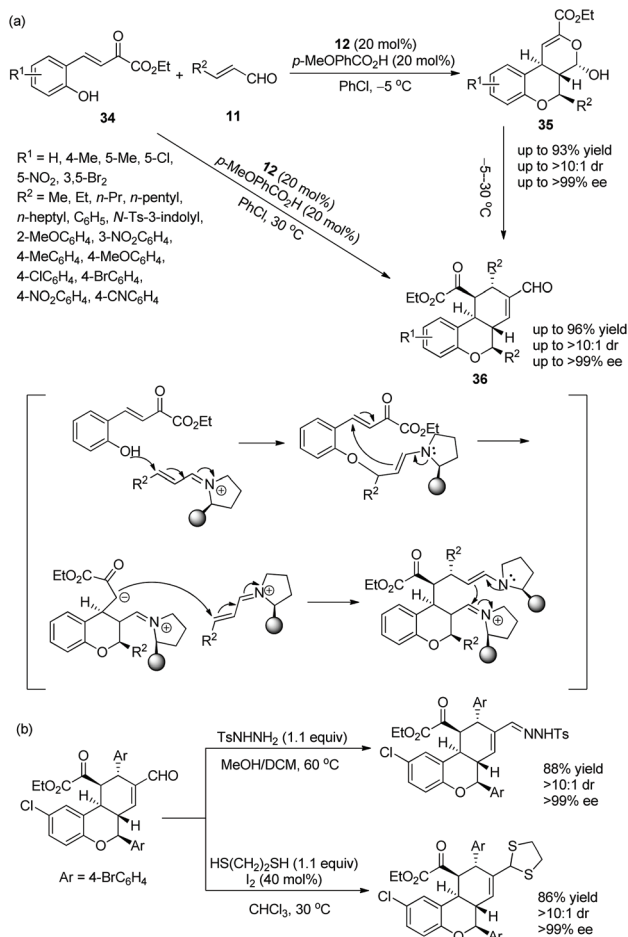


**Scheme 12** Oxa-Michael cascade reaction for the synthesis of chiral tricyclic chromans.

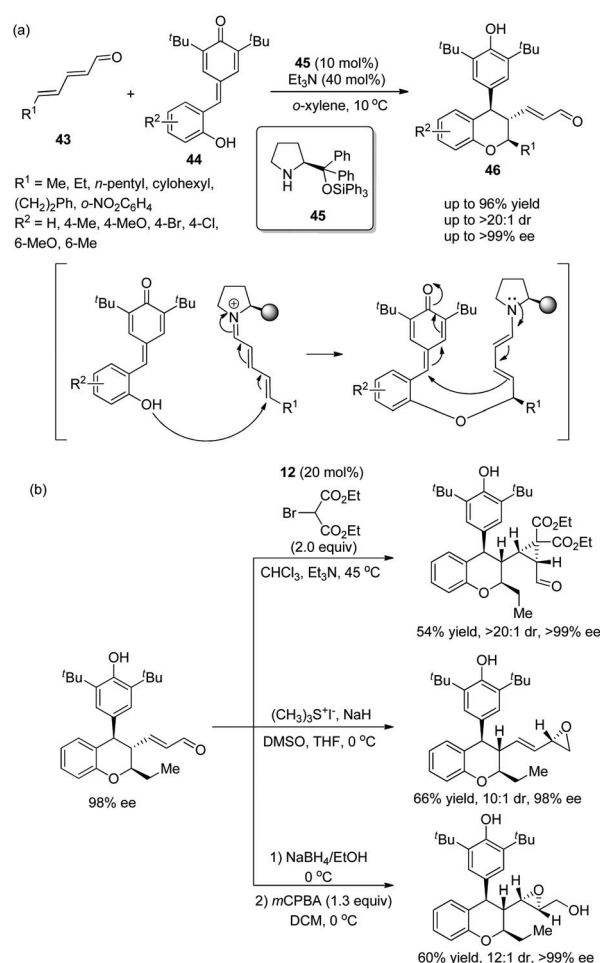
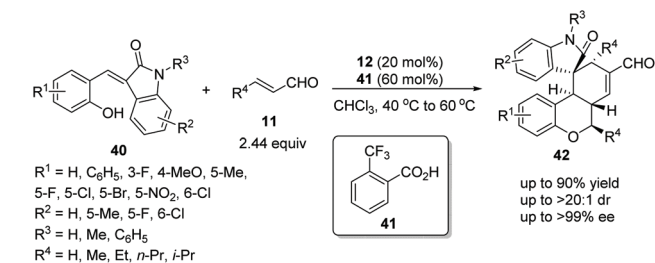
nol and dichloromethane at  $60^{\circ}\text{C}$ , but it could also be transformed to the corresponding mercaptal by reaction with 1,2-ethanedithiol in  $\text{CHCl}_3$  at  $30^{\circ}\text{C}$ , both without losing enantiomeric purity (Scheme 11b).

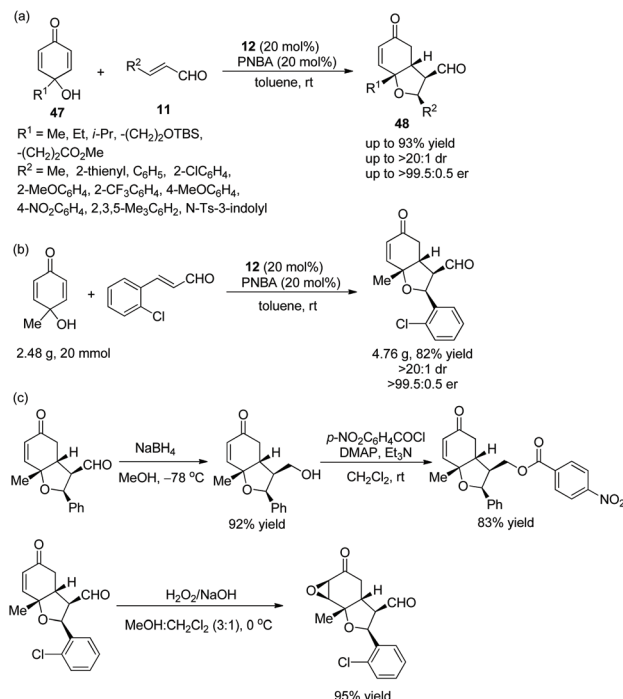
In view of the satisfactory study by Wang's group, Li's group<sup>21</sup> also reported some similar work in 2014. *o*-Hydroxychalcones **37** and cinnamaldehydes **38** were used as the starting materials to synthesize chiral chroman derivatives **39** catalyzed by the same organocatalyst **12**. In this procedure, they added  $\text{Et}_3\text{N}$  to activate the cascade reaction, and eventually gained a satisfying result (up to 74% yield and  $>99\%$  ee) (Scheme 12). Nevertheless, the scope of substrates for this cascade reaction is relatively limited owing to the fine-tuning of the electronic properties of the chalcone substrates, and the positions of the substituents have a great influence on the reaction. For instance, the reaction would hardly proceed when  $\text{R}^2$  was a strong electron-withdrawing nitro group, and no reaction would be observed if the position of the electron-withdrawing group ( $\text{R}^1 = \text{F}$ ) was changed from the 5-position to the 3-position. From the perspective of the mechanism, the carbon anion of the intermediate does not have sufficient nucleophilic ability to attack the cinnamaldehyde intermediate activated by the organocatalyst under the influence of these factors.

Wang's group<sup>22</sup> in 2017 reported a straightforward and efficient protocol for the construction of highly stereoselective polycyclic spiro-fused carbocyclic oxindoles *via* an oxa-Michael addition triggered cascade reaction. The introduction of a hydroxy group on the alkylideneoxindoles **40**, successively performed as nucleophiles and electrophiles under the catalysis of catalyst **12** and additive *o*-trifluoromethylbenzoic acid **41** in chloroform at  $40^{\circ}\text{C}$  to  $60^{\circ}\text{C}$ , and effectively facilitated the cascade reaction with unsaturated aldehydes **11**. The desired products **42** were obtained in good yields (up to 90%) with excellent stereoselectivities (up to  $>20:1$  dr and  $>99\%$  ee) (Scheme 13). It is noteworthy that it was actually a three-component cascade process. After an oxa-Michael/Michael addition reaction had occurred between alkylideneoxindoles **40** and unsaturated aldehydes **11**, the intermediate product bearing a new nucleophilic site at the 3-position of oxindole would continue to undergo Michael/aldol condensation with unsaturated aldehydes **11** to generate the desired products **42**. Additionally, in the scope of substrates, the worst result (55% yield,  $1:1.25$  dr and 64% ee) was obtained when there was no

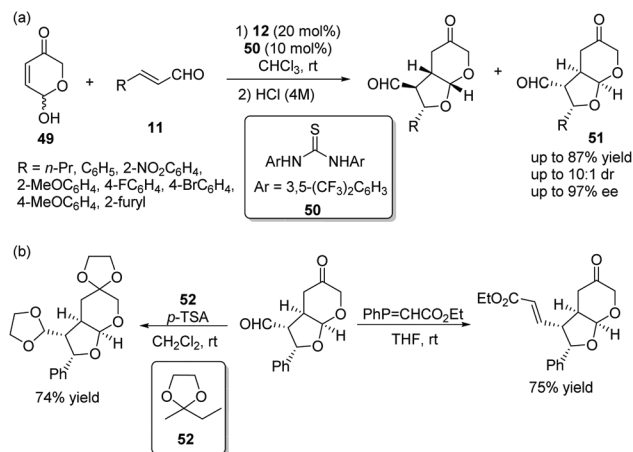


**Scheme 11** Oxa-Michael-IED/HDA-Michael-aldol cascade reaction for the synthesis of chroman derivatives and the derivatizations of chiral chroman.





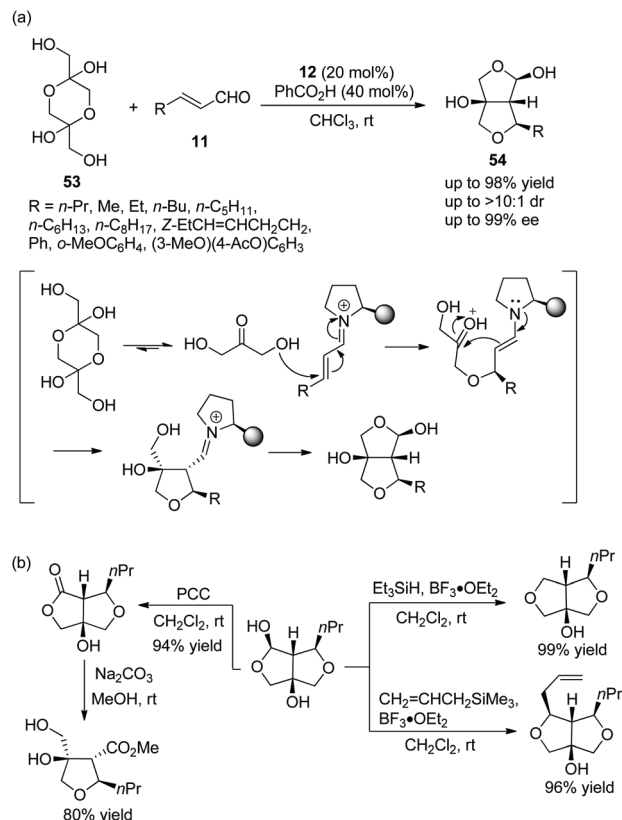
**Scheme 15** Oxa-Michael/Michael cascade reaction for the synthesis of cyclic dialkyl ethers and the derivatization of the chiral adduct.



**Scheme 16** Oxa-Michael/Michael cascade reaction for the synthesis of bicyclic furo[2,3-*b*]pyrans and the derivatization of the chiral adduct.

compared with aromatic substituents, aliphatic substituents showed higher reactivity and lower stereoselectivity. The electronic property and position of the substituents on the aromatic ring had little effect on the cascade reaction, but electron-withdrawing substituents also showed better reactivity than electron-donating substituents. In addition, some transformations were performed to construct functionalized chiral building blocks, which successfully demonstrated the application of this method (Scheme 16b).

**2.1.4 Oxa-Michael cascade reaction for the synthesis of chiral alicyclic compounds.** An unprecedented and efficient

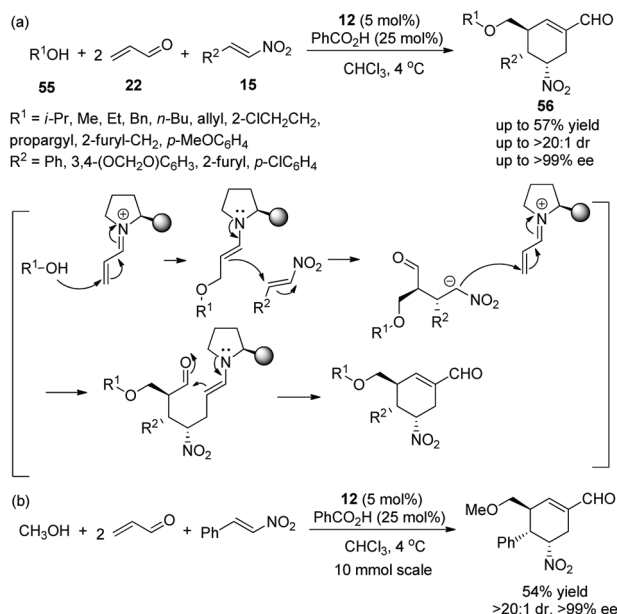


**Scheme 17** Oxa-Michael/aldol/hemiacetalization cascade reaction for the synthesis of furofurans and the derivatization of the chiral adduct.

method for the construction of polysubstituted furofurans *via* an oxa-Michael cascade reaction was reported by Vicario and coworkers.<sup>26</sup> Dihydroxyacetone dimer 53 as an oxygen nucleophile was reacted with  $\alpha,\beta$ -unsaturated aldehydes 11 under the catalysis of catalyst 12 and benzoic acid in chloroform at room temperature. After an initial oxa-Michael addition reaction, a subsequent intramolecular aldol reaction, and lastly a hemiacetalization step, the desired products 54 were generated with excellent yields (up to 98%) and stereoselectivities (up to >10:1 dr and 99% ee) (Scheme 17a). In the scope of substrates, all of the substituents afforded the corresponding products with high stereoselectivities; and the reactivity of aliphatic-substituted unsaturated aldehydes was better than that of aromatic-substituted unsaturated aldehydes. In order to illustrate the applicability of this strategy, some further transformations were performed, such as replacement of functionalized groups, oxidation of hydroxyl, and ring-opening reactions, which afforded the corresponding products with high yields (Scheme 17b).

An asymmetric multicomponent cascade reaction catalyzed by organocatalysts for the construction of functionalized structures always receives considerable attention. Gong's group<sup>27</sup> in 2009 reported a valuable and efficient protocol for the synthesis of chiral polysubstituted cyclohexenes *via* a multicomponent cascade reaction. Under the catalysis of catalyst 12 and benzoic acid, simple alcohol 55, a double equivalent of acro-

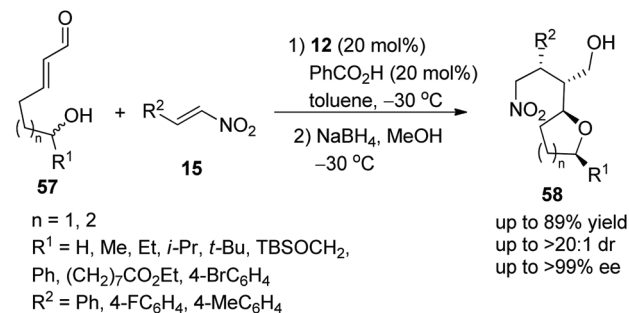




**Scheme 18** Multicomponent cascade reaction for the synthesis of cyclohexene carbaldehydes.

lein 22, and nitroalkenes 15 underwent an oxa-Michael/Michael/Michael/aldol condensation cascade process to generate the desired products 56 in moderate yields (up to 57%) with excellent stereoselectivities (up to >20:1 dr and >99% ee) in chloroform at 4 °C (Scheme 18a). In the scope of substrates, regardless of whether the substituents were aliphatic or aromatic, all of them afforded the corresponding products with excellent stereoselectivities; the longer the carbon chain of the alcohol, the lower the yield of the corresponding products. In addition, in a 10-times scale reaction, the desired chiral poly-substituted cyclohexene was obtained without losing any enantiopurity (Scheme 18b).

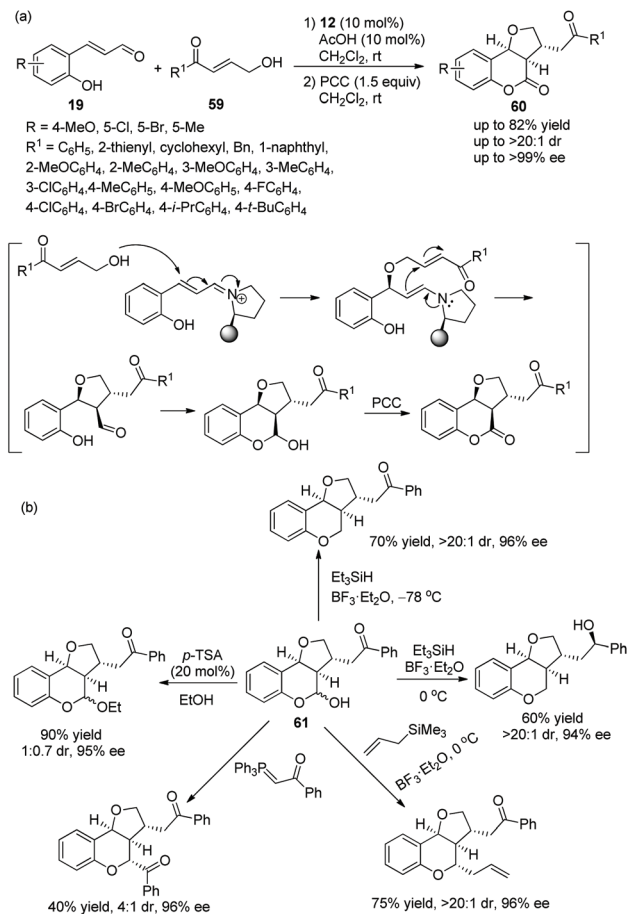
Brenner-Moyer and co-workers<sup>28</sup> reported a new strategy for the highly enantioselective construction of functionalized tetrahydropyrans and tetrahydrofurans *via* an oxa-Michael/Michael cascade reaction in 2012. Under the catalysis of catalyst 12 and benzoic acid, linear unsaturated aldehyde 57 bearing hydroxyl underwent an intramolecular oxa-Michael addition followed by an intermolecular Michael addition with nitroalkenes 15; then the desired products 58 were obtained in good yields (up to 89%) with excellent stereoselectivities (up to >20:1 dr and >99% ee) in toluene at −30 °C (Scheme 19). In the scope of the tetrahydropyran-forming substrates, although the reactivity of substituted 57 was low, the enantiopurity of the corresponding products was well maintained. Then in the scope of tetrahydrofuran-forming substrates, the substituents  $\text{R}^1$  had little influence on the enantioselectivity of this reaction, substrate 57 with aliphatic-substituted  $\text{R}^1$  afforded the corresponding products with better yield than substrate 57 with aromatic-substituted  $\text{R}^1$ ; and unsubstituted 57 gave a single cascade product in moderate yield (62%) with excellent enantioselectivity (>99% ee). Furthermore, some compu-



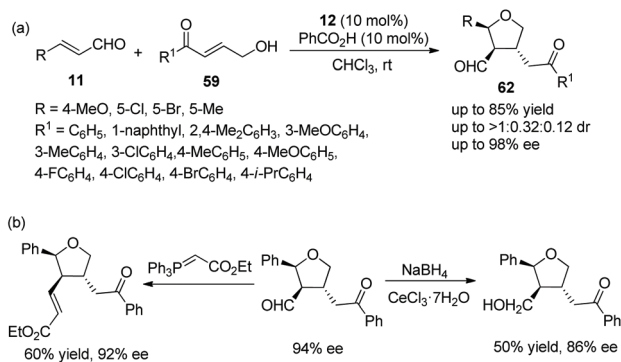
**Scheme 19** Oxa-Michael/Michael cascade reaction for the synthesis of chiral tetrahydropyrans and tetrahydrofurans.

tational studies were performed to illustrate this catalytic cascade process.

**2.1.5 Oxa-Michael cascade reaction of  $\gamma$ -hydroxyl enones.** Pan and coworker<sup>29</sup> developed a convenient strategy for the asymmetric synthesis of dihydrocoumarin derivatives bearing a tetrahydrofuranylated carbon stereocenter *via* an oxa-Michael cascade reaction in 2018. In this cascade reaction between *o*-hydroxy cinnamaldehydes 19 and  $\gamma$ -hydroxy enones 59, the desired products 60 were obtained in good yields (up to 82%) with satisfactory diastereoselectivities (up to >20:1 dr) and enantioselectivities (up to >99% ee) under the combined action of chiral catalyst 12 and acetic acid (Scheme 20a). Nevertheless, in the scope of  $\gamma$ -hydroxy enones, though the thienyl-substituted  $\gamma$ -hydroxy enone could participate in the cascade reaction, the corresponding product was obtained with the lowest yield (62%) and enantioselectivity (46% ee); the electron-donating group on the aromatic substituents of 59 showed better reactivity than the electron-withdrawing group; and *ortho*-substituents on the aromatic ring of 59 provided the corresponding products with lower yields than other positions. No reaction occurred with  $\gamma$ -hydroxy enones bearing aliphatic substituents. In addition, they converted the unoxidized hemiacetal 61 to some valuable organic structures, such as the formation of acetal, the introduction of an allyl group and the reaction with Wittig reagent. Interestingly, the same reaction of dehydroxylation treated with triethyl silane and  $\text{BF}_3\cdot\text{OEt}_2$  at different temperatures would generate different products, and all their enantioselectivities were satisfactory (Scheme 20b). In the same year,<sup>30</sup> they again explored the synthesis of tetrahydrofuran derivatives *via* this cascade reaction using  $\alpha,\beta$ -unsaturated aldehydes 11 and  $\gamma$ -hydroxy enones 59. They used another additive, benzoic acid, to promote the process in  $\text{CHCl}_3$  solvent, and the corresponding products 62 were obtained with increased yields and enantioselectivities (Scheme 21a). In the scope of enals, when the methyl-substituted enal was employed, the corresponding product was obtained with the lowest enantioselectivity (57% ee). At the same time, in the scope of  $\gamma$ -hydroxy enones, no reaction was observed when the alkyl-substituted  $\gamma$ -hydroxy enone was utilized. Additionally, two further transformations were established subsequently, which illustrated the applicability of this



**Scheme 20** Oxa-Michael cascade reaction for the synthesis of chiral dihydrocoumarin derivatives and the derivatization of hemiacetal.

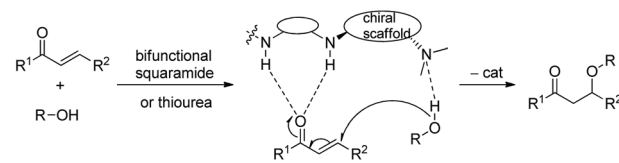


**Scheme 21** Oxa-Michael cascade reaction for the synthesis of chiral tetrahydrofuran derivatives and the derivatization of chiral tetrahydrofuran.

strategy and afforded the corresponding products without losing much enantiomeric purity (Scheme 21b).

## 2.2 Reactions catalyzed by chiral bifunctional squaramides

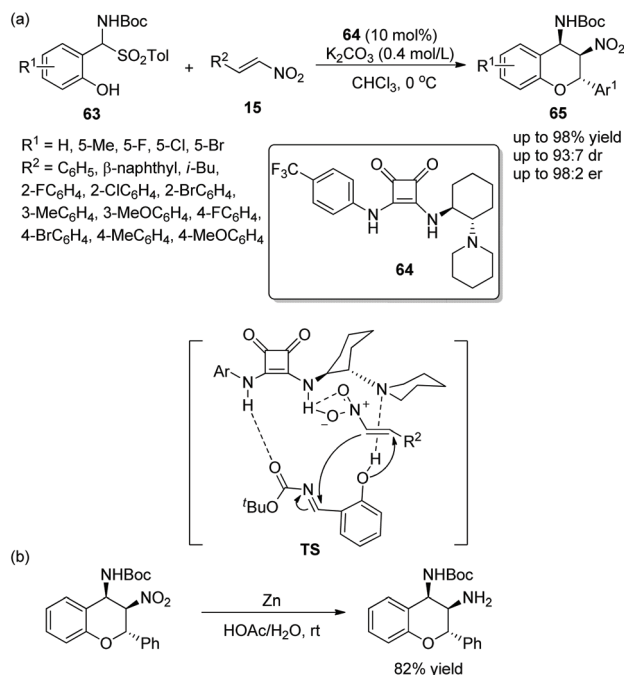
As we know, chiral squaramide, as a type of bifunctional chiral catalyst with good catalytic activity, has been of interest to a



**Scheme 22** Chiral bifunctional squaramide-catalyzed general mechanism of the oxa-Michael reaction.

wide range of organic researchers and has been successfully applied in many asymmetric catalytic cascade reactions, including the oxa-Michael cascade reaction, which has achieved high enantioselectivities.<sup>31</sup> The catalytic process of bifunctional squaramides is similar to bifunctional thioureas, which catalyze conjugate addition by forming hydrogen bonds with the reactants (Scheme 22).

**2.2.1 Oxa-Michael cascade reaction for the synthesis of chiral chromans.** Due to the significance of chromans in pharmaceutical structures and natural products, many novel methods for the construction of these chiral structures *via* an oxa-Michael cascade reaction catalyzed by various chiral bifunctional squaramides have been developed. Peng's group<sup>32</sup> developed an effective approach to synthesize the chroman derivatives with high enantioselectivities in 2014. The *o*-hydroxyaryl-substituted  $\alpha$ -amido sulfones **63** and nitroolefins **15** catalyzed by squaramide **64** bearing chiral cyclohexane underwent an oxa-Michael/aza-Henry cascade process and afforded the products **65** with excellent yields (up to 98%) and stereo-selectivities (up to 93:7 dr and 98:2 er) (Scheme 23a). It should be noticed that all the reaction times were relatively

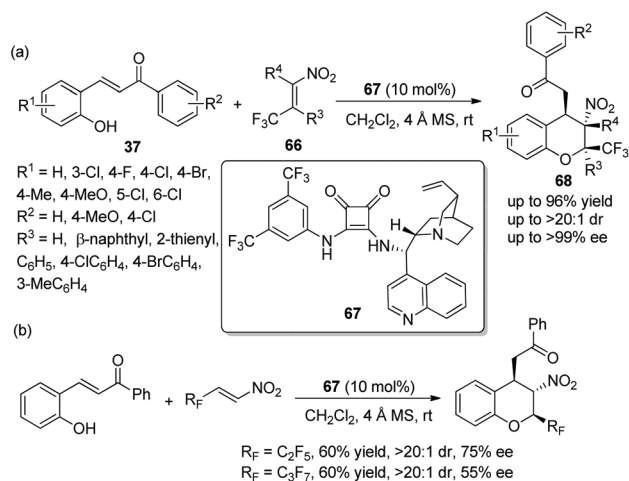


**Scheme 23** Oxa-Michael/aza-Henry cascade reaction for the synthesis of benzopyran derivatives and the derivatization of the chiral product.

long in the scope of this cascade reaction; in particular, isobutyl-substituted nitroolefin generated the lowest yield (46%), but other substituted nitroolefins gave satisfactory results; and the position and electronic property of substituents on the aromatic ring of **63** had little effect on the reaction. Moreover, in order to demonstrate the applicability of this strategy, they tried a reduction reaction of a chiral product using Zn/HOAc at room temperature, which transformed the nitro group to an amino group (Scheme 23b).

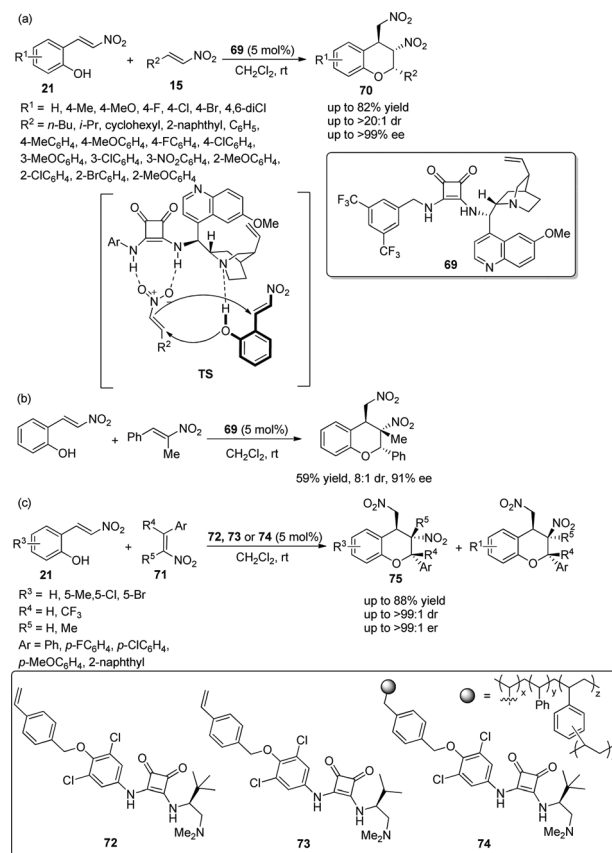
An admirable piece of work was reported by Wang's group<sup>33</sup> for the synthesis of 2-CF<sub>3</sub> chromans in 2015. In this cascade reaction, β-CF<sub>3</sub>-nitroalkenes **66** and *o*-hydroxychalcones **37** were selected as the starting materials, which were catalyzed by a chiral squaramide **67** for the construction of optically active trifluoromethylated chromans **68**. The products **68** bearing three contiguous stereocenters were obtained in high yields (up to 96%) with excellent stereoselectivities (up to >20:1 dr and >99% ee) (Scheme 24a). It has to be said that in the scope of substrates, all the substituents, whether electron-withdrawing or electron-donating groups, afforded the corresponding products with excellent enantioselectivities, even though the reaction times were different. The position of the substituents had an effect on the reactivity: for example, if the Cl was moved from the 6-position to the 3-position of *o*-hydroxychalcone, the relevant reaction time would be prolonged. Compared with unsubstituted R<sup>4</sup>, methyl-substituted R<sup>4</sup> gave the corresponding product with better yield. In addition, the authors further performed this cascade reaction by utilizing perfluoroalkyl substituted β-nitroalkenes (Scheme 24b). The result displayed that pentafluoroethyl- and heptafluoropropyl-substituted nitroalkenes showed lower reactivity and afforded the products with lower yields and enantioselectivities than β-CF<sub>3</sub> nitroalkenes, but it is still of great significance for the construction of chiral polyfluorine-substituted chromans.

Xu's group<sup>34</sup> in 2018 developed an efficient strategy for the construction of chiral chroman derivatives *via* an oxa-Michael cascade reaction catalyzed by chiral bifunctional squaramide



**Scheme 24** Oxa-Michael cascade reaction for the synthesis of chromans bearing trifluoromethyl.

**69**. The desired products **70** containing two nitro groups were generated in good yields with excellent diastereo- and enantioselectivities by the oxa-Michael/Michael cascade process of 2-(*E*)-(2-nitrovinyl)phenols **21** and nitroolefins **15** under mild conditions (Scheme 25a). In the scope of this cascade reaction, the position of the substituents on the aromatic ring of **21** had little effect on the reaction, but the electron-donating substituents provided the corresponding products with better yields and enantioselectivities than electron-withdrawing substituents; and aliphatic-substituted nitroolefins generally provided adducts in higher yields than did aromatic substituents. In aromatic-substituted nitroolefins, compared with electron-withdrawing groups, electron-donating groups afforded better enantioselectivities. Additionally, they further examined the synthetic applicability of this cascade reaction by using *trans*-α-Me-β-nitroolefin, which showed a good result with moderate yield and stereoselectivity (Scheme 25b). Later, Pedrosa's group<sup>35</sup> developed a similar strategy for the construction of chiral trisubstituted chroman derivatives *via* an oxa-Michael triggered cascade reaction catalyzed by 4-vinylphenyl-substituted squaramides. Under the catalysis of 4-vinylphenyl-substituted squaramides **72** or **73**, nitrovinylphenols **21** and multi-substituted nitroalkenes **71** afforded desired products **75** with good yields (up to 88%) and stereoselectivities (up to >99:1 dr and >99:1 er) in dichloromethane at room temperature



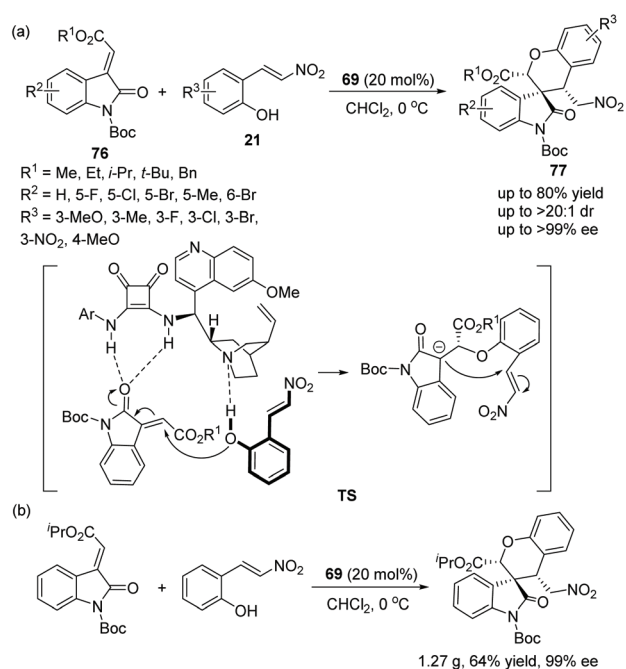
**Scheme 25** Oxa-Michael cascade cyclization reaction for the synthesis of chiral dinitro chromans.

(Scheme 25c). In the scope of substrates, compared with nitroolefins with electron-donating groups, nitroolefins with electron-withdrawing groups showed better reactivity, but afforded the corresponding products with lower stereoselectivities; and if trisubstituted nitroolefins were applied to the reaction, such as  $R^4$  being replaced by trifluoromethyl or  $R^5$  being replaced by methyl, worse results were obtained. Additionally, another catalyst **74** as the polymeric homolog of **72** was also employed, but no better results were obtained. In general, catalyst **72** showed the best catalytic ability.

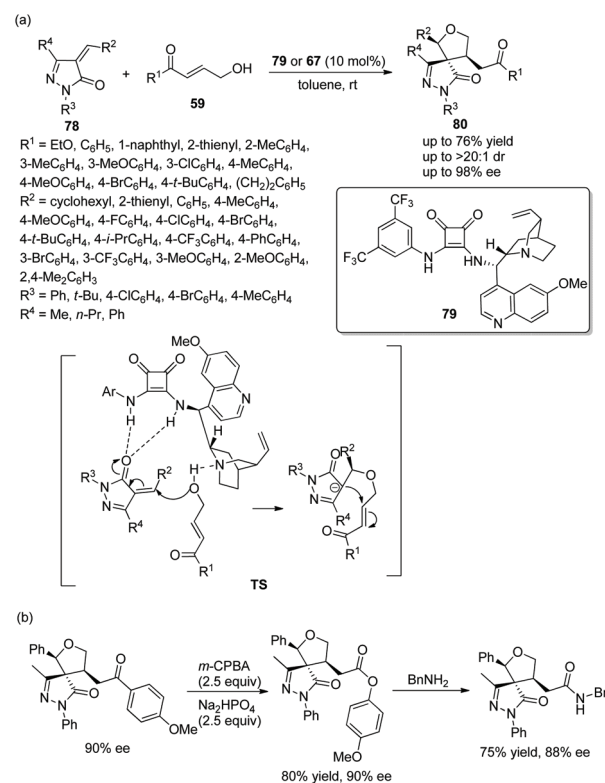
**2.2.2 Oxa-Michael cascade reaction for the synthesis of chiral spiro cyclic compounds.** Zhu's group<sup>36</sup> described a novel, efficient and valuable methodology for the synthesis of spirooxindoles **77** bearing benzopyran structures *via* oxa-Michael cascade reactions in 2013. It is worth noting that the cascade reaction of methyleneindolinones **76** with nitrovinylphenols **21** catalyzed by chiral bifunctional squaramide **69** was highly determined by the *N*-protecting group. Only when the *N*-protecting group of the oxindoles was a Boc group could the desired products be obtained. Satisfactorily, the reactions afforded good yields and great stereocontrol (>20:1 dr, >99% ee) under mild conditions (Scheme 26a). In an exploration of the substrate scope, the position and electronic property of the substituents on the aromatic ring of **21** and **76** had little effect on the reaction; it is noted that reactant **21**, in which  $R^3$  was substituted by the electron-withdrawing nitro group, needed the longest reaction time to generate the desired product in the lowest yield (53%), even though the enantioselectivity achieved was 98% ee. Furthermore, they successfully performed this cascade reaction on a gram-scale without losing

enantioselectivity and demonstrated the applicability of this strategy (Scheme 26b).

Given the pharmaceutical and medicinal applications of pyrazolone derivatives, an effective method to prepare chiral spiro pyrazolones with three consecutive stereocenters was reported by Pan's group<sup>37</sup> in 2018. They employed unsaturated pyrazolones **78** and  $\gamma$ -hydroxyenones **59** to catalytically synthesize chiral spiro tetrahydrofuran-pyrazolones **80**. Whether chiral bifunctional squaramide **79** or **67** was employed, the desired products were attained in good yields with excellent stereoselectivities (up to >20:1 dr and 98% ee) (Scheme 27a). The reason why two catalysts were used is that, when examining the generality of unsaturated pyrazolones in this cascade reaction, it was found that cinchonidine-derived catalyst **67** showed better enantioselective control than quinine-derived catalyst **79**. In the scope of substrates, aryl-substituted  $\gamma$ -hydroxyenones showed better results than alkyl-substituted  $\gamma$ -hydroxyenones; in the scope of pyrazolones with different benzylidene substituents, all the substituents provided the corresponding products with high enantioselectivities, and the reactivity of the electron-donating groups was better than that of the electron-withdrawing groups; in the scope of *N*-substituents, no desired product was observed when the *N*-substituent was *tert*-butyl. In addition, a few further transformations, selective oxidation and a substitution reaction with benzylamine, were performed and afforded the corres-



**Scheme 26** Oxa-Michael cascade reaction for the synthesis of benzo-pyran-spirooxindoles.



**Scheme 27** Oxa-Michael/Michael cascade reaction for the synthesis of spiro tetrahydrofuran-pyrazolones and the derivatization of the chiral product.

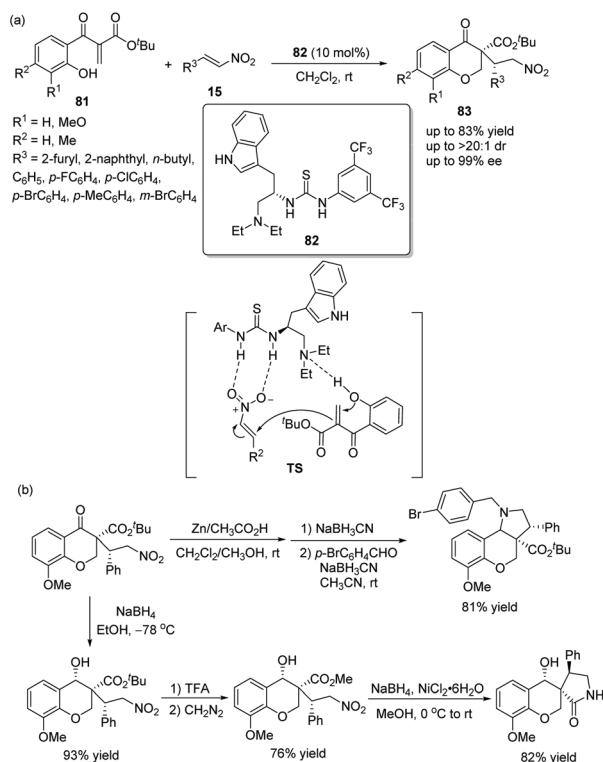


ponding products without losing much enantiopurity, which proved the synthetic utility of the cascade strategy (Scheme 27b).

### 2.3 Reactions catalyzed by chiral bifunctional thioureas

Since chiral thioureas have been shown to have an ability to participate in molecular recognition processes by selectively forming hydrogen bonds, they have been employed as comparatively effective organocatalysts in many conversions besides oxa-Michael reactions. The chiral bifunctional thiourea catalytic general mechanism of oxa-Michael is similar to that of chiral squaramides (Scheme 22).

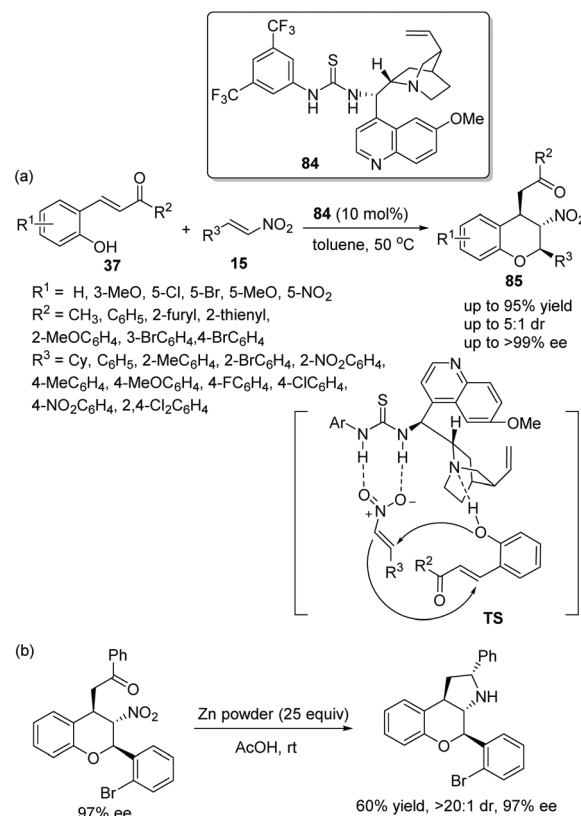
**2.3.1 Oxa-Michael/Michael cascade reaction of ethylene  $\beta$ -keto esters with nitroolefins.** A novel and efficient strategy for the construction of highly functionalized chiral chromanones *via* an oxa-Michael/Michael cascade reaction was reported by Lu's group<sup>38</sup> in 2011. Ethylene  $\beta$ -keto esters **81**, as starting materials would first be employed in a cascade reaction; then reaction with nitroolefins **15** in dichloromethane could smoothly generate the desired adducts **83** in good yields (up to 83%) with excellent stereoselectivities (up to >20:1 dr and 99% ee) under the catalysis of a tryptophan-derived tertiary amine-thiourea catalyst **82** (Scheme 28a). In the scope of substrates, *n*-butyl substituted nitroolefin showed the worst reactivity (41% yield) and enantioselectivity (89% ee), but other substituted nitroolefins provided satisfactory results; and electron-rich substituents on the aryl ring of ethylene  $\beta$ -keto esters **81** promoted the reaction more efficiently.



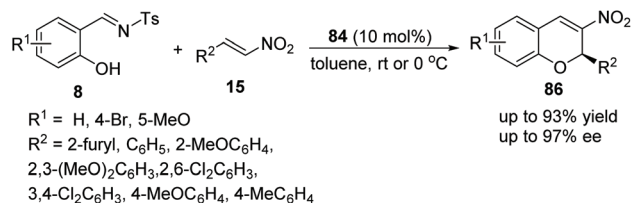
**Scheme 28** Oxa-Michael/Michael cascade reaction for the synthesis of chiral chromanones and the derivatization of the chiral product.

Furthermore, the 3,3-disubstituted 4-chromanone products could be further functionalized, such as reductive amination of the nitro group and the formation of lactam (Scheme 28b), which successfully demonstrated the applicability of this methodology.

**2.3.2 Oxa-Michael cascade reactions for the synthesis of chiral chromans.** Another effective and simple strategy for the construction of chromans bearing multiple stereocenters was proposed by Singh's group<sup>39</sup> in 2015, which expanded the field of synthesis of chiral chromans. The desired products **85** were obtained with excellent yields (up to 95%) and stereoselectivities (up to 5:1 dr and >99% ee) *via* an oxa-Michael/Michael cascade reaction catalyzed by a chiral bifunctional amino-thiourea **84** (Scheme 29a). And in the substrate scope of this cascade reaction, the  $\beta$ -nitrostyrenes **15** bearing electron-donating groups, such as methyl and methoxy, would take a longer reaction time than the  $\beta$ -nitrostyrenes bearing electron-withdrawing groups to generate the corresponding products. Moreover, in the scope of the substrates **37**, irrespective of whether substituents on the aromatic ring were electron-donating or electron-withdrawing, all of them provided the corresponding products with high enantioselectivities. However, for substrate **37** with a nitro group on the aromatic ring, no product was obtained, which might be due to the hydrogen bond formed by the nitro group and the bifunctional catalyst hindering the cascade process. Additionally, to display the syn-



**Scheme 29** Oxa-Michael/Michael cascade reaction for the synthesis of chiral chromans and the derivatization of the chiral product.



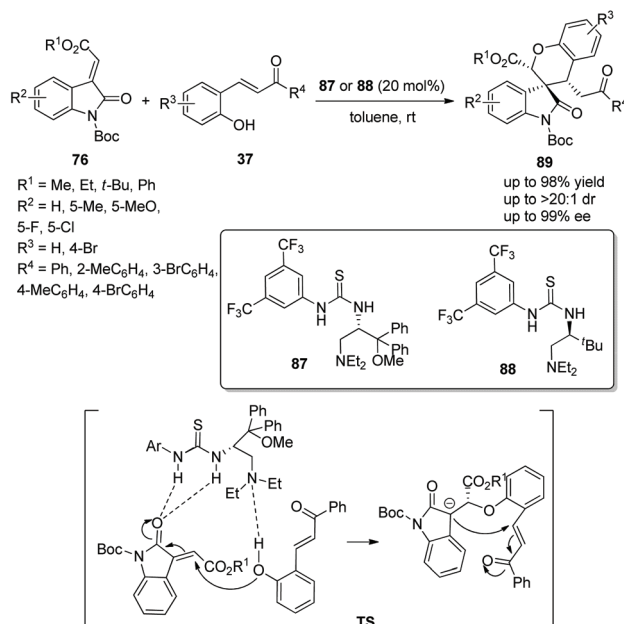
**Scheme 30** Oxa-Michael/aza-Henry/desulfonamidation cascade reaction for the synthesis of chiral 2H-chromenes.

thetic utility of this cascade method, a transformation to generate tricyclic chroman was performed, and the corresponding product was obtained without losing enantioselectivity (Scheme 29b).

Schreiner's group<sup>40</sup> reported a simple and feasible protocol for the construction of 2H-nitrochromenes *via* an oxa-Michael cascade reaction. Under the catalysis of bifunctional thiourea **84** in toluene, salicyl *N*-tosylimines **8** and nitrostyrenes **15** underwent an oxa-Michael/aza-Henry/desulfonamidation cascade process and afforded the desired products **86** with high yields and enantioselectivities (Scheme 30). In the scope of substrates, compared with electron-withdrawing substituents, electron-donating-group-substituted salicyl *N*-tosylimines provided the corresponding products with better enantiopurity; when the reaction temperature was decreased to 0 °C, the enantioselectivity of the corresponding products was promoted but the yield was decreased.

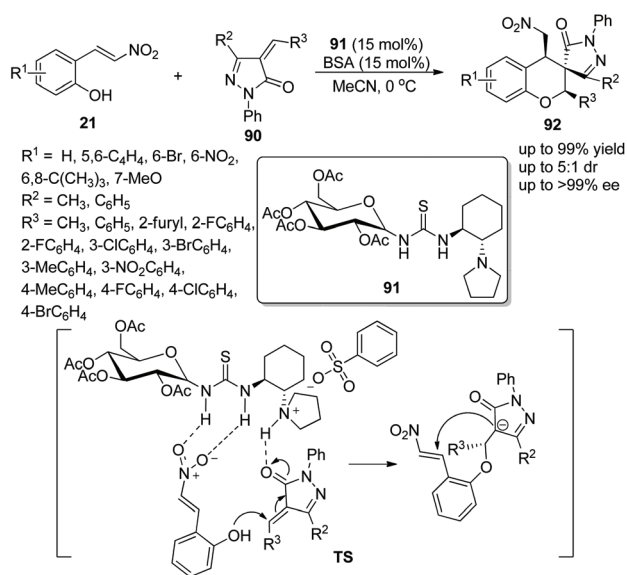
**2.3.3 Oxa-Michael/Michael cascade reaction for the synthesis of chiral spirocyclic compounds.** Spirocyclic compounds as a vital structure in natural products and organic synthesis has always attracted the attention of many researchers, which has resulted in various functionalized spiro compounds being synthesized. Zhao's group<sup>41</sup> in 2014 reported an available and efficient strategy for the construction of chiral spirooxindole-chromans *via* an oxa-Michael/Michael cascade reaction. Under the catalysis of amino-acid-derived thioureas **87** or **88**, methyleneindolinones **76** reacted with *o*-hydroxychalcones **37** and provided the corresponding products **89** with excellent yields (up to 98%) and stereoselectivities (up to >20:1 dr and 99% ee) in toluene at room temperature (Scheme 31). In the scope of substrates catalyzed by organocatalyst **87**, the electronic property and position of substituents on the aromatic ring of **76** had little effect on the stereoselectivity, but electron-withdrawing substituents showed better reactivity than electron-donating substituents; ester and ketone groups at the carbon-carbon double bond could also be tolerated well. Noticeably, except for the *N*-protecting group Boc, other *N*-protecting groups could not participate in the process. When organocatalyst **88** was used, the corresponding products were obtained with higher yields but lower enantioselectivities.

A suitable and effective proposal for the construction of functional spiro chroman-pyrazolones *via* a bifunctional amino-thiourea **91** catalyzed oxa-Michael/Michael addition cascade reaction was presented by Miao's group<sup>42</sup> in 2015. The nitrovinyl phenols **21** and 4-alkenyl pyrazolin-3-ones **90** afforded



**Scheme 31** Oxa-Michael/Michael cascade reaction for the synthesis of chiral spiro benzohydropyran-oxindoles.

the target products **92** bearing three contiguous stereocenters with excellent yields (up to 99%) and enantioselectivities (up to >99% ee) under the catalysis of catalyst **91** and benzene sulfonic acid (BSA) additive in MeCN at 0 °C (Scheme 32). In the substrate scope of this cascade reaction, the good reactivity of all the pyrazolones with electron-donating or electron-withdrawing groups at any position of the benzene ring furnished the corresponding products with high yields and enantioselectivities; a methyl substituent at  $\text{R}^2$  showed better reactivity



**Scheme 32** Oxa-Michael/Michael cascade reaction for the synthesis of chiral spiro benzohydropyran-pyrazolones.

and stereoselectivity than a phenyl substituent. Moreover, an alkyl substituted at  $R^3$  of unsaturated pyrazolone provided the corresponding product with the best enantioselectivity in a comparably short reaction time.

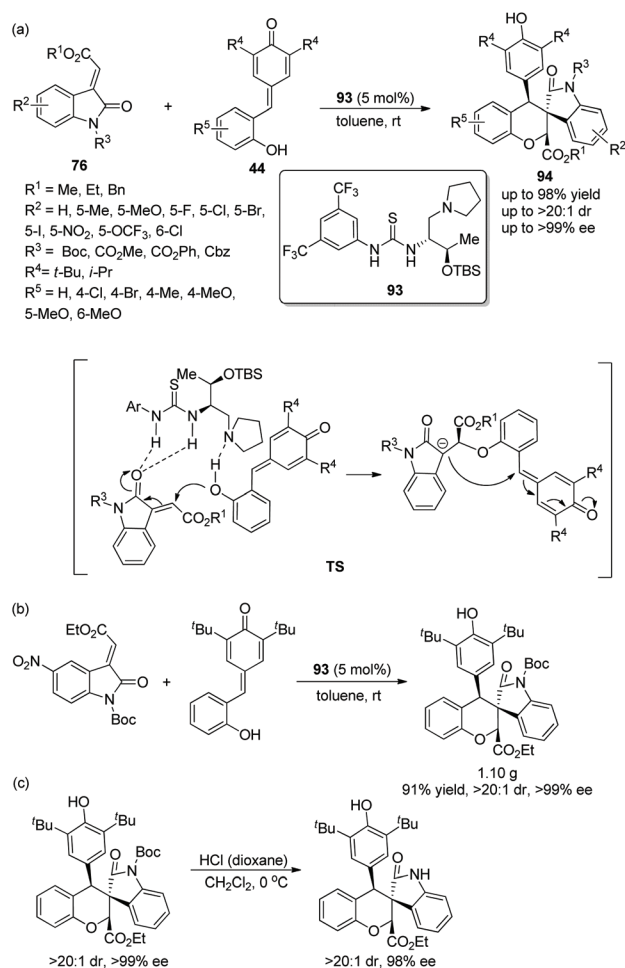
Since there were no precedents for employing *o*-hydroxyphenyl substituted *p*-quinone methides **44** as donor–Michael acceptor synthons in domino addition procedures before 2016, a viable and elegant approach to fill the gap was developed by Enders' group.<sup>43</sup> After the methyleneindolinones **76** and *o*-hydroxyphenyl substituted *p*-quinone methides were catalyzed by chiral bifunctional aminothiurea **93** and underwent an oxa-Michael addition and intramolecular 1,6-addition, the desired chiral functionalized spirochromans **94** were generated with excellent yields and stereoselectivities under mild conditions (Scheme 33a). In the substrate scope of this cascade reaction, electron-donating and electron-withdrawing substituents were both tolerated well, and all of them furnished the corresponding products with excellent enantioselectivities; electron-withdrawing substituents on the aromatic ring of **76** or **44** showed better reactivity than electron-

donating substituents. Nevertheless, when the  $R^4$  substituent was replaced by an isopropyl group, the stereoselectivity of this reaction was not influenced, but the reactivity was decreased, which afforded the product with the lowest yield (43%). The cascade reaction on a gram-scale was accomplished under optimum conditions and maintained the stereoselectivity of the corresponding product (Scheme 33b). In addition, a deprotection transformation to remove the *N*-Boc group was smoothly finished without losing much enantiopurity (Scheme 33c).

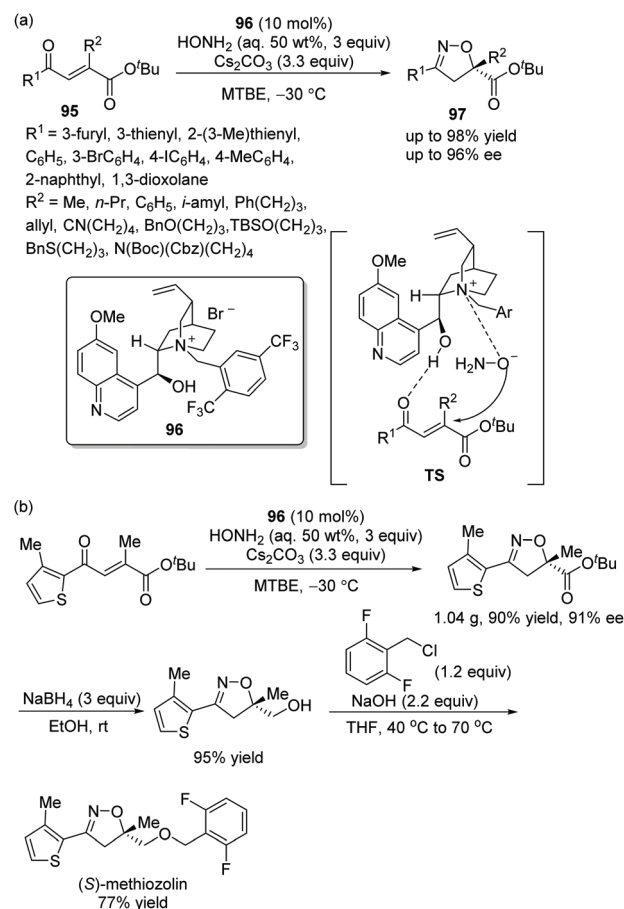
## 2.4 Reactions catalyzed by other chiral organocatalysts

There are some other organocatalytic asymmetric oxa-Michael addition triggered cascade reactions, for example chiral cinchona alkaloid derived catalysts, which cannot be strictly classified into the preceding catalyst groups but still produce a satisfactory result. Moreover, from the perspective of the reaction mechanism, they activate an oxa-Michael cascade reaction by forming hydrogen-bond intermediates.

**2.4.1 Oxa-Michael cascade cyclization reaction of  $\beta$ -carboxy substituted  $\alpha,\beta$ -unsaturated ketones with hydroxylamine catalyzed by chiral quinidine-based catalyst.** Since isoxazoline derivatives play a significant role in bioactive heterocycles, and



**Scheme 33** Oxa-Michael/1,6-addition cascade reaction for the synthesis of spiro chromans-oxindoles and the derivatization of the chiral product.



**Scheme 34** Oxa-Michael cascade reaction for the synthesis of chiral isoxazolines and the derivatization of the chiral product.

the synthesis of some chiral isoxazoline derivatives with potential bioactivity is of great benefit to the pharmaceutical industry and medicine, Cho's group<sup>44</sup> displayed an efficient and useful strategy for the construction of chiral carboxy-substituted isoxazolines *via* an oxa-Michael cascade cyclization reaction in 2018. The  $\beta$ -carboxy substituted  $\alpha,\beta$ -unsaturated ketones **95** and hydroxylamine were phase-transfer-catalyzed by a combination of a chiral quinidine-derived catalyst **96** and cesium carbonate ( $\text{Cs}_2\text{CO}_3$ ), and the corresponding products **97** were generated with high yields and enantioselectivities in methyl *tert*-butyl ether (MTBE) at  $-30^\circ\text{C}$  (Scheme 34a). In the substrate scope of this reaction, all of the substituents provided the corresponding products with satisfactory yields and enantioselectivities, and electron-donating groups were proven to show a better result. Moreover, a gram-scale reaction was carried out and afforded the corresponding product without losing much yield or enantioselectivity. A further transformation for the synthesis of (*S*)-methiozolin in two steps was performed, which demonstrated well the applicability of this cascade strategy (Scheme 34b).

### 3 Conclusions

In summary, we have introduced recent advances in the field of organocatalytic asymmetric oxa-Michael addition triggered cascade reactions and have illustrated that it is a comparatively valuable and powerful tool for the rapid formation of functionalized oxygen heterocyclic molecules containing stereocenters. A great variety of novel and potentially bioactive compounds with excellent stereoselectivities have been synthesized *via* various oxa-Michael addition triggered cascade reactions catalyzed by a diverse number of organocatalysts under different conditions, which has indeed expanded the scope of the methodology of the Michael addition cascade reaction. Many further transformations have been performed to display the applicability of the corresponding oxa-Michael addition triggered cascade reactions. However, there is still a lot of work needed to promote and optimize the process, such as more novel and recyclable organocatalysts, more diversified adducts, more convenient reaction processes and so on. It is expected that many more interesting and valuable developments will be established to increase the diversity of oxa-Michael triggered reactions in the near future.

### Conflicts of interest

There are no conflicts to declare.

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