Iridium-catalyzed asymmetric hydrogenation of racemic \(\alpha\)-substituted lactones to chiral diols\(^{†}\)

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We report a protocol for the highly efficient iridium-catalyzed asymmetric hydrogenation of racemic \(\alpha\)-substituted lactones via dynamic kinetic resolution. Using Ir-SpiroPAP (R)-1d as a catalyst, a wide range of chiral diols were prepared in a high yield (80–95%) with a high enantioselectivity (up to 95% ee) under mild reaction conditions. This protocol was used for enantioselective syntheses of \((\pm)\)-preclamol and a chiral 2,5-disubstituted tetrahydropyran.

**Introduction**

Transition-metal-catalyzed asymmetric hydrogenation of ketones is an efficient and reliable method for the synthesis of optically active chiral secondary alcohols.\(^1\) In contrast, the enantioselective synthesis of chiral primary alcohols by catalytic asymmetric hydrogenation of their corresponding aldehydes or esters is difficult, and work on the development of practical methods is underway. In 2007, we reported the first example of the catalytic asymmetric hydrogenation of racemic \(\alpha\)-branched aldehydes, via dynamic kinetic resolution (DKR), for the synthesis of chiral primary alcohols.\(^2\) Subsequently, List\(^3\) and Lin\(^4\) et al. also reported the synthesis of chiral primary alcohols, by means of ruthenium-catalyzed asymmetric hydrogenation of racemic \(\alpha\)-substituted aldehydes. Although a wide range of catalysts have been developed for the hydrogenation of esters,\(^5\) efficient chiral catalysts for the asymmetric hydrogenation of racemic \(\alpha\)-substituted esters via DKR are rare. The biggest challenge for the direct asymmetric hydrogenation of racemic esters to form optically active primary alcohols is to find a catalyst that can discriminate between the enantiomers of chiral esters and then hydrogenate them to alcohols selectively. In 2011, Ikariya\(^6\) et al. described the enantioselective hydrogenation of a racemic mixture of a \(\alpha\)-substituted \(\gamma\)-lactone to a chiral 1,4-diol via DKR using chiral ruthenium catalysts bearing chiral 1,2-diamine ligands (Scheme 1). However, the enantioselectivity of the reaction was low (up to 32% ee). Recently, as part of our work on the asymmetric hydrogenation of ketones, we found that chiral Ru-SDPs/diamine catalysts and chiral Ir-SpiroPAP catalysts can also mediate the hydrogenation of ester groups.\(^7\)

**Results and discussion**

In this communication, we report a protocol for the Ir-SpiroPAP-catalyzed asymmetric hydrogenation of racemic \(\alpha\)-substituted lactones to afford chiral diols in a high yield (80–95%) with a high enantioselectivity (up to 95% ee, Scheme 1).

**Scheme 1** Asymmetric hydrogenation of racemic \(\alpha\)-substituted lactones via DKR.
(R)-1d (entry 1). However, when the concentration of BuOK was increased to 0.06 M, the hydrogenation reaction occurred and provided (R)-3a in 15% yield with 91% ee (entry 2); further increasing the concentration of BuOK increased the reaction rate and the yield of 3a substantially. For example, when 0.25 M BuOK (2a/BuOK/(R)-1d = 500 : 500 : 1) was used, the reaction was complete within 10 h, providing (R)-3a in 91% yield with 92% ee (entry 3). The absolute configuration of (R)-3a was determined by comparing the sign of its optical rotation with the literature data. Evaluation of various chiral Ir-SpiroPAP catalysts (R)-1 revealed that the substrates on the pyridine and phenyl groups of the catalysts had little effect on the yield or enantioselectivity (entries 4–8), with (R)-1d giving the best results. Experiments with various solvents showed that PrOH was suitable (entries 9–11); the reaction was complete within 10 h, affording (R)-3a in 92% yield with 93% ee. In addition to BuOK, BuONa also gave a high yield with a high enantioselectivity, but the use of KOH, NaOH, or K2CO3 resulted in low yields (entries 12–15).

To evaluate the substrate scope of the reaction, we investigated a wide range of racemic α-substituted ß-valerolactones under the established reaction conditions (Table 2). For racemic α-aryl-substituted ß-valerolactones 2a–i (entries 1–9), neither electron-donating nor electron-withdrawing groups on the phenyl ring of the substrates had much effect on the enantioselectivity of the reaction, but substrates with an electron-withdrawing group (entries 2, 5, and 8) showed a higher reaction rate than those with an electron-donating group.

In addition, owing to steric effects, substrates with an ortho-substituent on the phenyl ring gave lower reaction rates, yields, and enantioselectivities (entries 10–12). The hydrogenation of α-alkyl-substituted ß-valerolactones 2m–p also worked well, affording the corresponding 1,5-diols 3m–p in high yields and high enantioselectivities (entries 13–16). Catalyst (R)-1d also catalyzed the hydrogenation of racemic α-substituted γ-butyrolactones 2q and 2r, providing the 1,4-diols 3q (80% ee) and 3r (69% ee), respectively (entries 17 and 18).

We investigated the pathway of the hydrogenation of racemic α-substituted lactones 2 by 1H NMR. As shown in Fig. 1, after reaction for 0.5 h under the optimal reaction conditions, rac-2a was converted to the hydroxyl ester 4, propyl 5-hydroxy-2-phe-nylpentanoate, in 62% yield with no ee and the diol 3a in 38% yield with 93% ee. Over the following 10 h, the amount of the hydroxyl ester 4 gradually decreased, and the amount of the diol 3a increased. Only a trace amount of lactone 2a was detected from 2 min after the reaction started.

Direct hydrogenation of the hydroxyl ester 4 with catalyst (R)-1d provided the diol 3a in 93% yield with 93% ee, which is the same as the result obtained from the hydrogenation of lactone 2a (Scheme 2). We also conducted the hydrogenation of...
the ester 5, which has a $\delta$-OCH$_2$OMe group instead of a $\delta$-OH group as in the hydroxyl ester 4, and observed no reaction. These results indicated that the hydroxyl group of ester 4 was crucial for the hydrogenation. Thus, although the lactone 2a was readily alcoholized to the hydroxyl ester 4 under the reaction conditions, the hydrogenation of 2a occurred inevitably via its lactone form (Scheme 2).$^{7a,r}

Scheme 2  Asymmetric hydrogenation of esters 4 and 5 with (R)-1d.

Chiral 3-aryl/alkyl substituted piperidines are part of an important class of bioactive heterocyclic compounds, but they are difficult to synthesize in their optically active forms.$^{9}$ By using the catalyst (S)-1d, we synthesized (−)-preclamol,$^{11}$ which is a candidate drug for the treatment of neurological disorders such as Parkinson’s disease.$^{12}$ The hydrogenation of rac-2g (1.65 g) catalyzed by (S)-1d afforded the diol (S)-3g (89% yield and 93% ee), which was subsequently transformed to (−)-preclamol by activation with methanesulfonyl chloride, substitution/cyclization with n-propylamine, and demethylation with hydrobromic acid (84% yield over three steps, Scheme 3).

Diol (S)-3p is a useful building block for the synthesis of chiral 2,5-disubstituted tetrahydropyrans, which occur in many biologically active natural products such as the terpenoids rhopaloic acid A$^{13}$ and barangcadoic acid A$^{14}$ (Scheme 4), isolated from marine sponges. Iodoetherification of (S)-3p with iodine$^{15}$ produced the tetrahydropyran 7 in 94% yield as a 2:1 trans/cis mixture. Nucleophilic substitution of tetrahydropyran 7 with NaCN afforded the nitrile 8 (trans/cis = 2:1). Hydrolysis of the nitrile 8 and its subsequent esterification with MeOH afforded the tetrahydropyran 9 in a 72% yield with a higher trans/cis ratio (5:1). Thus, this protocol represents a potential method for the construction of the chiral core structures of rhopaloic acid A and barangcadoic acid A.$^{16}$

Conclusions

In conclusion, we have developed a protocol for the highly efficient iridium-catalyzed asymmetric hydrogenation of racemic α-substituted lactones via DKR. Using an Ir-SpiroPAP catalyst, a series of racemic α-substituted lactones were hydrogenated to chiral diols in high yield with high enantioselectivity under mild reaction conditions. The protocol was used for the enantioselective syntheses of (−)-preclamol and a chiral 2,5-disubstituted tetrahydropyran.

Fig. 1  The plots of the hydrogenation of rac-2a with (R)-1d.

Scheme 4  Enantioselective synthesis of a chiral 2,5-disubstituted tetrahydropyran.
Experimental

General procedure for the asymmetric hydrogenation of 2

Into a 20 mL hydrogenation vessel in an autoclave, racemic α-substituted δ-valerolactone 2 (1.0 mmol), a solution of iridium catalyst (R)-1d in 9PrOH (dried with MS 4 Å for 12 h, 0.002 mmol mL⁻¹, 1.0 mL, 0.002 mmol), a solution of 'BuOK in 9PrOH (0.5 mmol mL⁻¹, 2.0 mL, 1.0 mmol) and 9PrOH (1.0 mL). The autoclave was purged with hydrogen by pressurizing to 5 atm and releasing the pressure. This procedure was repeated three times and then the autoclave was pressurized to 10 atm of H₂. The reaction mixture was stirred at room temperature (25–30 °C) until no obvious hydrogen pressure drop was observed. The reaction mixture was then quenched with saturated NH₄Cl (5 mL) and extracted with EtOAc (5 mL × 3). The combined extracts were washed with brine, dried over anhydrous MgSO₄ and concentrated in vacuo. The residue was purified by flash column chromatography on a silica gel using petroleum ether/ethyl acetate as the eluent to afford the chiral diols 3. The ee values of the chiral diols 3 were determined by HPLC using a chiral column.

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Notes and references


