## Unusual Effect of a Mixed Solvent on the Asymmetric Reduction of Chiral $\alpha$ -Keto-amides with Sodium Borohydride

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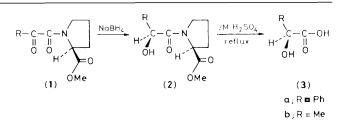
The mixed solvent tetrahydrofuran + methanol (99:1) was much more effective than the individual solvents in the asymmetric reduction of chiral  $\alpha$ -keto-amides with sodium borohydride; this solvent effect in asymmetric induction is unprecedented.

Despite the importance of the role of solvents in asymmetric synthesis, mixed solvents have been used very rarely.<sup>1</sup> We report the first example of a solvent effect on asymmetric induction in which a mixed solvent caused much higher asymmetric induction than either of the individual solvents.

In general, diastereoselectivities are low in the asymmetric reduction of chiral  $\alpha$ -keto-esters or -amides with complex metal hydrides,<sup>2</sup> especially with sodium borohydride.

 $\alpha$ -Keto-amides (1a, b),<sup>3</sup> derived from (S)-proline methyl ester and  $\alpha$ -keto-acids using dicyclohexylcarbodi-imide, were reduced with readily available NaBH<sub>4</sub> at 0 °C to afford (2a, b). Acidic hydrolysis of (2a, b) afforded the optically active  $\alpha$ -hydroxy-acids (3a, b). When (1a) was reduced in tetrahydrofuran (THF) or methanol alone, the resulting enantiomeric excess (e.e.) of the mandelic acid (3a) produced was 36% (THF) and 4% (MeOH).<sup>†</sup>

A striking solvent effect was observed, however, when a THF + methanol (99:1 by volume) co-solvent was used.



The optical purity of the resulting (S)-(+)-mandelic acid { $[\alpha]_{25}^{25}$  +101° (c 1.9, H<sub>2</sub>O)} increased to 64% e.e.† Using the same mixed solvent, (S)-(+)-lactic acid {(3b),  $[\alpha]_{25}^{25}$  -7.4° (c 2.9, 1.5 M NaOH)} was obtained in 55% e.e.‡ from (1b). Water, instead of MeOH, was also found to be effective.

In either mixed solvents, a catalytic amount of the protic solvent (MeOH or  $H_2O$ ) in the aprotic THF was found to be essential for good asymmetric induction. The results are summarized in Table 1.

<sup>†</sup> Enantiomeric excesses were determined by <sup>1</sup>H-n.m.r. analysis of (2a) and/or by reported specific optical rotation of (3a). (S)-(+)-Mandelic acid,  $[\alpha]_D$  +158° (H<sub>2</sub>O), S. Mitsui and A. Kanai, *Nippon Kagaku Zasshi*, 1965, **86**, 627.

<sup>&</sup>lt;sup>‡</sup> Based on the value of (S)-(+)-lactic acid  $[\alpha]_{20}^{20}$  --13.5° (c 2.5, 1.5 M NaOH), 'Aldrich Catalog Handbook of Fine Chemicals,' Aldrich Chemical Co. Inc., Wisconsin, 1980.

Ratio (v/v) THF: MeOH	( <b>3a</b> ) (% e.e.)
100: 0 99: 1 0:100	36 64 4
THF:H₂O 99: 1 85: 15	50 0

This work was partially supported by a Grant-in-Aid for Scientific Research from the Ministry of Education, Science and Culture.

Received, 9th August 1982; Com. 948

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