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## Corrigendum

Corrigendum to "Effect of substrate concentration on the enantioselectivity of cyclohexanone monooxygenase from *Acinetobacter calcoaceticus* and its rationalization" [Tetrahedron: *Asymmetry* 11 (2000) 3653]<sup>†</sup>

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In the legend to Fig. 1, (1R,5S)-2 should be replaced by (1R,5S)-3, (1S,5R)-2 by (1S,5R)-3, (1R,5S)-3 by (1R,5S)-2 and (1S,5R)-3 by (1S,5R)-2. The corrected legend is shown below.

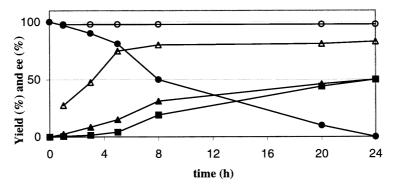


Figure 1. Time course of CYMO-catalyzed oxidation of 1 to lactones 2 and 3. Compound 1 (92 mM) ( $\geq$ 98% pure, provided by Fluka) was dissolved in 10 mL of 0.05 M Tris–HCl buffer, pH 8.6, containing 0.5 mM NADP, 1 M 2-propanol, 50 units of CYMO and 200 units of alcohol dehydrogenase from *Thermoanaerobium brokii* for coenzyme regeneration. The degree of conversion and the enantiomeric excess of the products were determined on ethyl acetate extracts by chiral GC with a CP-cyclodextrin column (50 m, 0.25 mm ID, Chrompack) at 130°C with H<sub>2</sub> as a carrier gas. Retention times were: (1*R*,5*S*)-1, 4.758 min; (1*S*,5*R*)-1, 4.799 min; (1*R*,5*S*)-3, 17.40 min; (1*S*,5*R*)-3, 17.60 min; (1*R*,5*S*)-2, 16.63 min; (1*S*,5*R*)-2, 17.28 min. ( $\bullet$ ) Percentage of remaining substrate 1; ( $\blacksquare$ ) percentage of formed lactone 2; ( $\triangle$ ) percentage of formed lactone 3; ( $\bigcirc$ ) (%) ee of lactone 3

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