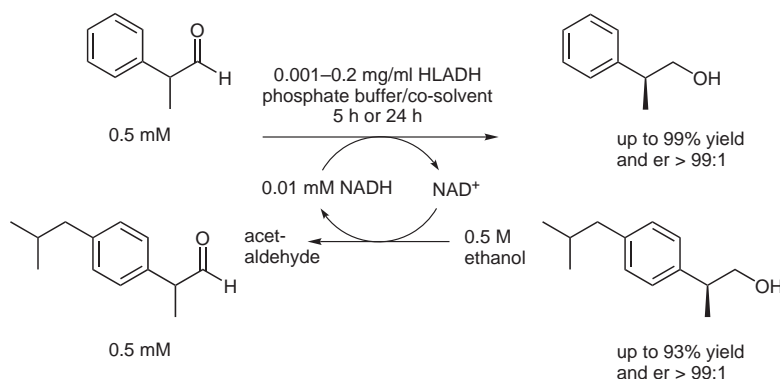


D. GIACOMINI,* P. GALLETTI, A. QUINTAVALLA, G. GUCCIARDO, F. PARADISI
(UNIVERSITY OF BOLOGNA, ITALY AND UNIVERSITY COLLEGE DUBLIN, IRELAND)
Highly Efficient Asymmetric Reduction of Arylpropionic Aldehydes by Horse Liver Alcohol Dehydrogenase through
Dynamic Kinetic Resolution
Chem. Commun. **2007**, 4038-4040.

Arylpropionic Alcohols via Enzyme-Mediated Dynamic Kinetic Resolution



Significance: The authors report their preliminary studies on the reduction of two 2-arylpropionic aldehydes to enantioenriched alcohols via a dynamic kinetic resolution. The enzyme catalyst is commercially available horse liver alcohol dehydrogenase (HLADH), with NADH as a cofactor. Since the enzyme regenerates its cofactor in the presence of ethanol, substoichiometric amounts of NADH are sufficient. It was found that organic co-solvents tetrahydrofuran and acetonitrile, which increase the solubility of the starting material, are tolerated in up to 10 vol% with respect to the buffer solution. With a change in the protocol, almost pure hexane (up to 99 vol%) can also be tolerated as solvent.

Comment: The dynamic kinetic resolution is a powerful tool to convert a starting material into an enantioenriched or even enantiopure product, theoretically in 100% yield, which is a distinct advantage over kinetic resolution methods. The catalyst in this reaction proves to be 'multitasking': Not only does it perform the reduction of the aldehyde, but there is also no need for adding a special racemization catalyst, and it recycles NADH by oxidizing ethanol and reducing NAD⁺. This greatly lowers the amount of expensive NADH employed in the reaction.