## Accepted Manuscript

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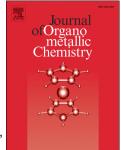
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## ACCEPTED MANUSCRIPT

A New Class of Well-Defined Ruthenium Catalysts for Enantioselective Transfer Hydrogenation of Various Ketones

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## 13 ABSTRACT

14 A pair of novel optically pure phosphinite ligands were synthesized by ring opening reaction of 15 chiral amines with (R)-styrene oxide or (S)-glycidyl phenyl ether oxide using a straightforward 16 method in high yields and their ruthenium complexes were described in detail. The ruthenium 17 complexes proved to be highly efficient catalysts for the enenatioselective hydrogenation of 18 ketones, affording products up to 99% ee. The results showed that the corresponding chiral 19 alcohols could be obtained with high activity and excellent enantioselectivities at the desired 20 (2S)-1-{benzyl[(1S)-1-(naphthalen-1-yl)ethyl]amino}-3-phenoxypropan-2-yl temperature. diphenylphosphinito[dichloro( $\eta^6$ -benzene)ruthenium (II)] acts an excellent catalyst in the 21 22 reduction of ketones, giving the corresponding alcohol up to 99% ee.

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Keywords: Asymmetric Transfer Hydrogenation; Chiral Ruthenium Complexes; Phosphinites;
Epoxide Ring opening; Homogeneous Catalysis.

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