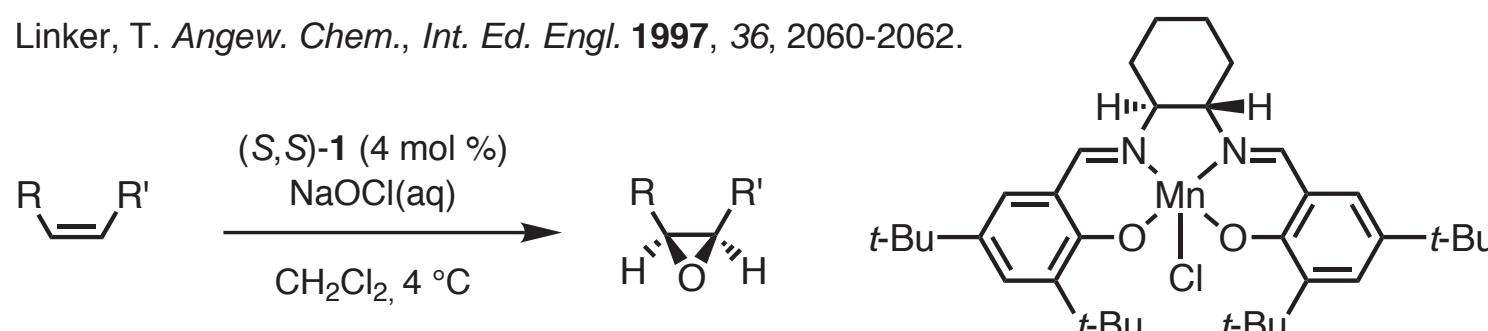


Reviews:

Jacobsen, E. N. In *Catalytic Asymmetric Synthesis*, Ojima, I., Ed.; VCH: New York, 1993; pp. 159-202.

Linker, T. *Angew. Chem., Int. Ed. Engl.* 1997, 36, 2060-2062.



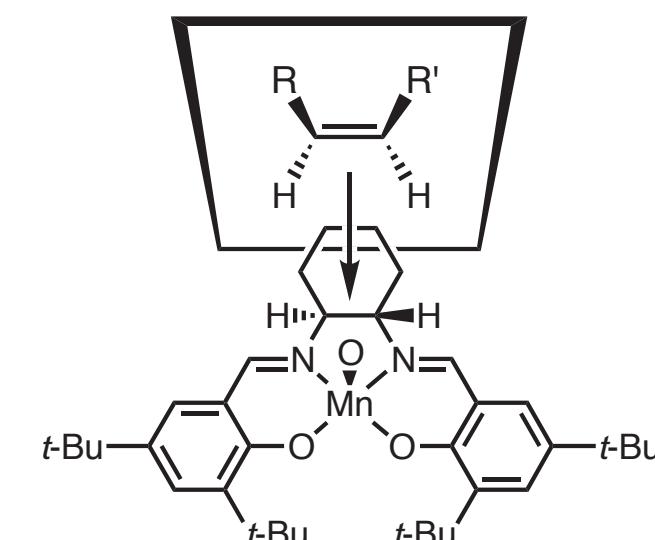
- Selectivity is determined through nonbonded interactions.
- In general, R is aryl, alkenyl or alkynyl and R' is a bulky group.
- *cis*-Disubstituted conjugated olefins are epoxidized with high levels of enantioselectivity.
- *trans*-Disubstituted olefins react more slowly and with diminished selectivity.

olefin	epoxide	yield, %	ee, %	equiv (S,S)-1
		96	97	0.03
		63	94	0.15
		79	84	0.006 ^a

^aReaction carried out in the presence of 4-phenylpyridine *N*-oxide.

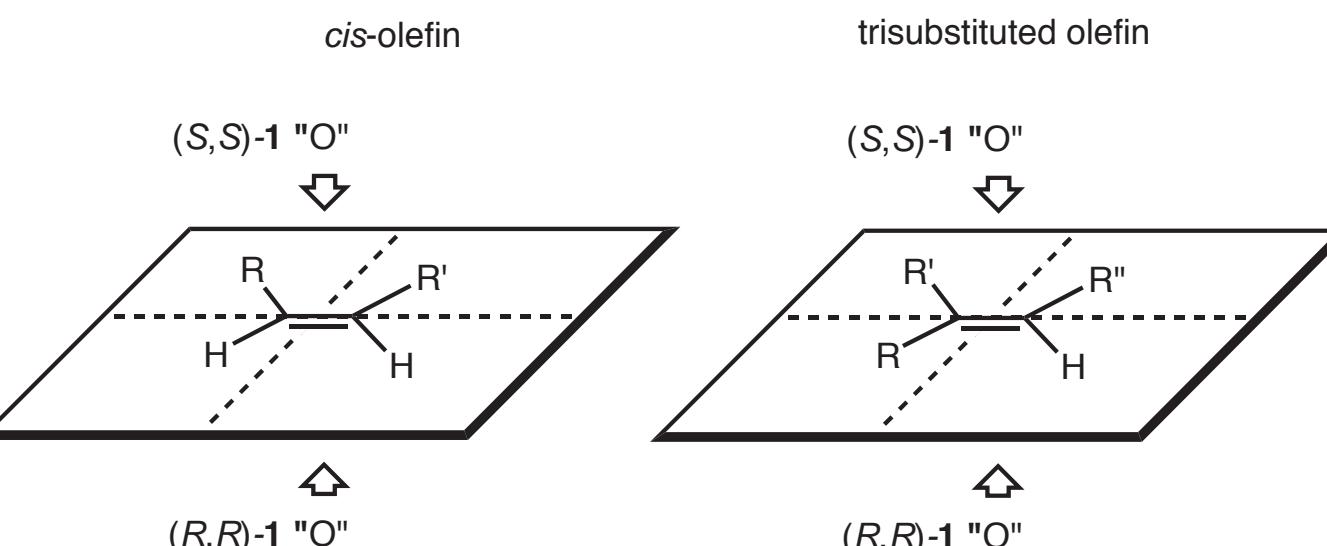
From: Jacobsen, E. N.; Zhang, W.; Muci, A. R.; Ecker, J. R.; Deng, L. *J. Am. Chem. Soc.* 1991, 113, 7063-7064.

- The observed selectivities are explained by a side-on approach of olefin:



- Terminal olefins are poor substrates.
- Addition of stoichiometric amounts of 4-phenylpyridine *N*-oxide improves both catalyst selectivity and turnover numbers.

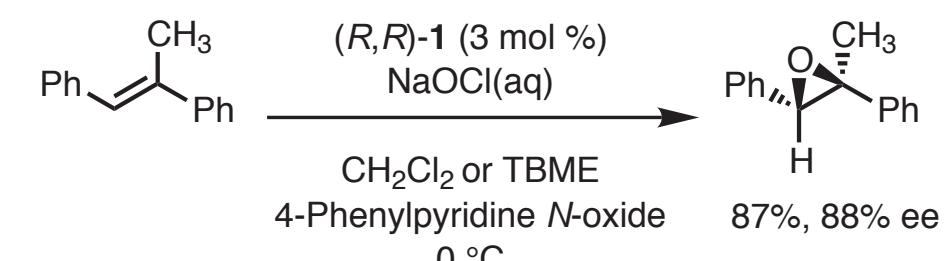
Mnemonics for the observed selectivities:



- *cis*-Olefins: Place Aryl, alkenyl or alkynyl substituent in upper-left quadrant (R) and the corresponding trans hydrogen atom in the lower-right quadrant.
- *trans*-Olefins are poor substrates.
- Trisubstituted olefins: Place the hydrogen atom in the lower-right quadrant.

Trisubstituted olefins:

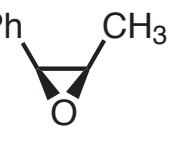
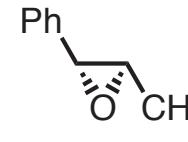
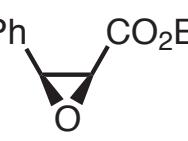
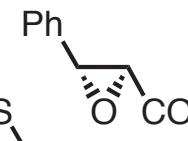
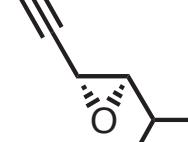
- Trisubstituted alkenes are excellent substrates for the Jacobsen asymmetric epoxidation.



olefin	epoxide ^a	yield, %	ee, %
		69	93
		91	95
		97	92

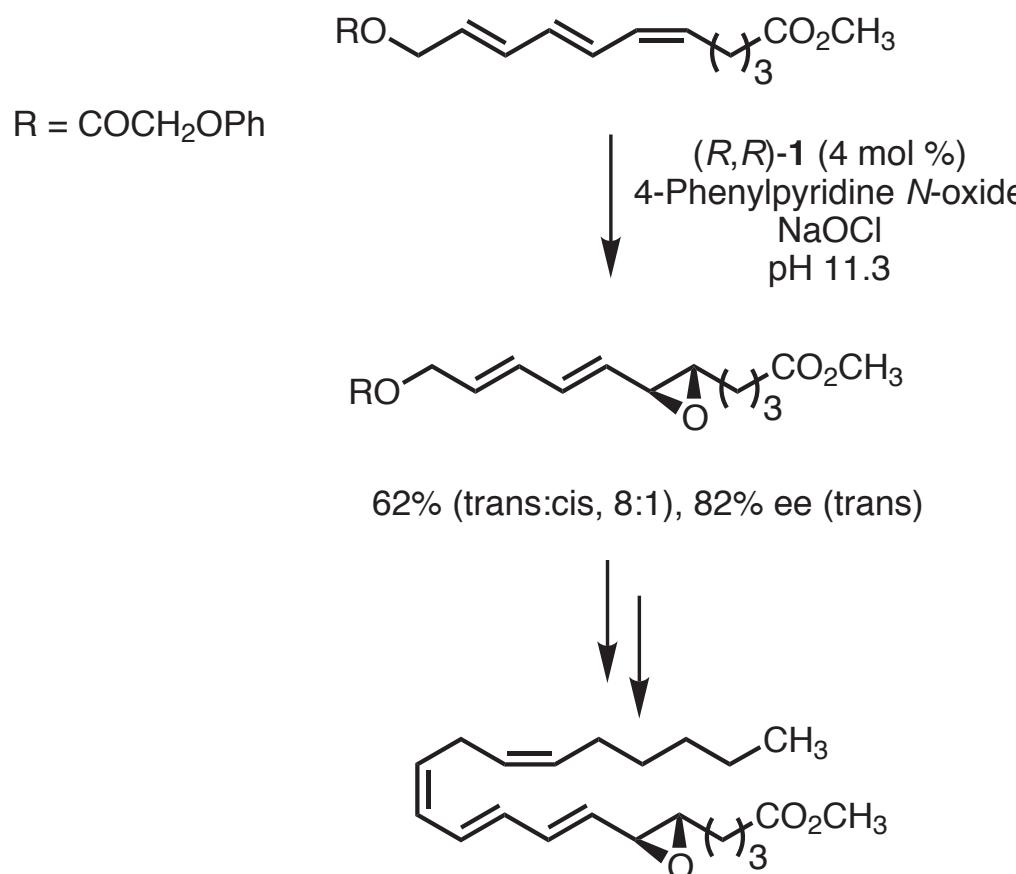
From: Brandes, B. D.; Jacobsen, E. N. *J. Org. Chem.* 1994, 59, 4378-4380.

- cis*- β -Substituted styrene derivatives afford *cis*-epoxides as major products while *cis*-enynes and *cis*-dienes produce *trans*-epoxides.

olefin	equiv (S,S)-1	<i>cis</i> -epoxide	yield (%), ee (%)	<i>trans</i> -epoxide	yield (%), ee (%)
Ph-CH=CH ₂	0.04		77, 92		6.7, 83
Ph-CH=CHCO ₂ Et	0.08		52, 97		15, 78
TMS-CH=CH-Cyclohexane	0.04		10, 64		55, 98

From: Jacobsen, E. N. In *Catalytic Asymmetric Synthesis*, Ojima, I., Ed.; VCH: New York, **1993**; pp. 159-202 and Jacobsen, E. N.; Zhang, W.; Muci, A. R.; Ecker, J. R.; Deng, L. *J. Am. Chem. Soc.* **1991**, *113*, 7063-7064.

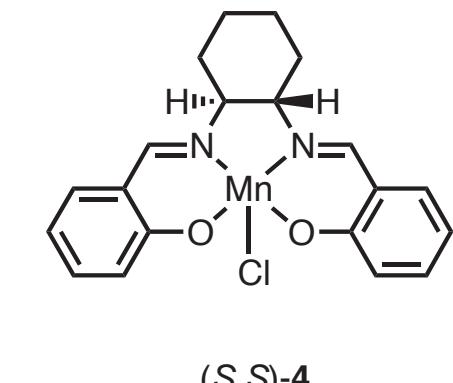
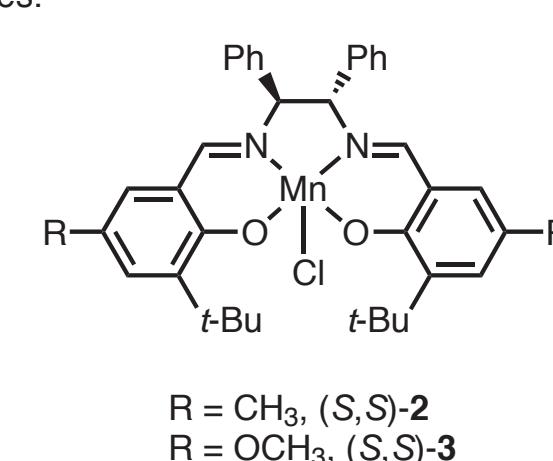
- Rotation of a radical intermediate is proposed to account for the *cis* \rightarrow *trans* isomerization.

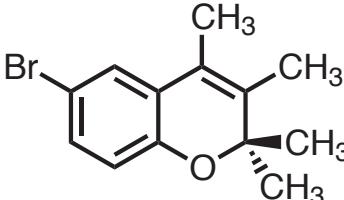
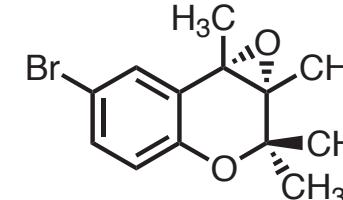
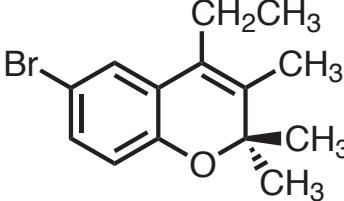
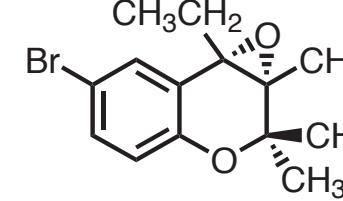
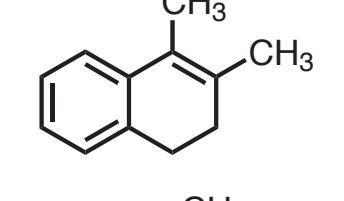
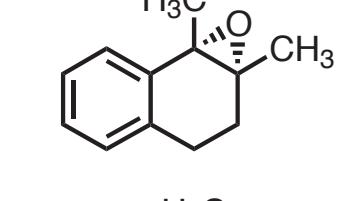
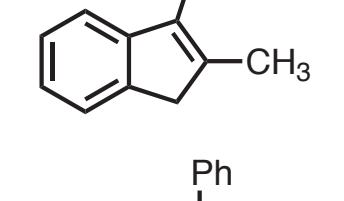
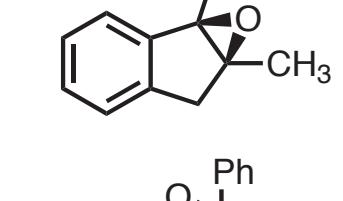
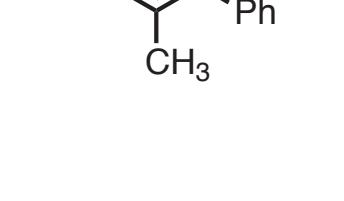
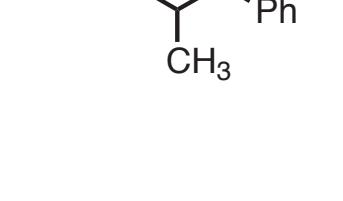


Leukotriene A₄ methyl ester

Tetrasubstituted Olefins:

- High enantioselectivities are not yet general but may be attained in certain cases with catalysts shown.
- Chromene derivatives undergo epoxidation with higher enantioselectivity as compared to indene derivatives.



olefin	catalyst	epoxide	epoxide yield, %	ee, %
	(S,S)-2		84	96
	(S,S)-2		81	97
	(S,S)-3		45	65
	(R,R)-3		37	35
	(S,S)-4		12	46