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Additions and Corrections

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Highly Facialselective Synthesis of Pyranose 1,3-Oxazines and Their Ring Opening with Nucleophiles: A Novel Entry to 2-*C*-Branched Glycosides

Page 2 5089, first column, line 24. The text should read, "generating in situ.[16]" A new reference should be added:
(16) For previous work on the asymmetric synthesis of monocyclic 1,3-dihydrooxazines from chiral acyclic enol ether, see: (a) Gizecki, P.; Dhal, R.; Toupet, L.; Dujardin, G. *Org. Lett.* **2000**, *2*, 585 (from *N*-acylimines). (b) Gizecki, P.; Dhal, R.; Poulard, C.; Gosselin, P.; Dujardin, G. *J. Org. Chem.* **2003**, *68*, 4338 (from *N*-acyl-*O*-alkyl acetals).
Page 5090, first column, line 24. The text should read, "Interestingly, when N-Boc-protected acetal was used as a substrate, tetrahydrooxazinone **8ah** (Figure 2) was produced exclusively.[17]" A new reference should be added:
(17) For previous asymmetric synthesis of monocyclic tetrahydrooxazinones from chiral acyclic enol ether and *N*-acyl-*O*-alkyl acetals, see: Gizecki, P.; Youcef, R. A.; Poulard, C.; Dhal, R.; P.; Dujardin, G. *Tetrahedron Lett.* **2004**, *45*, 9589.
We thank Dr. Gilles Dujardin for bringing the omission for our attention.

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