

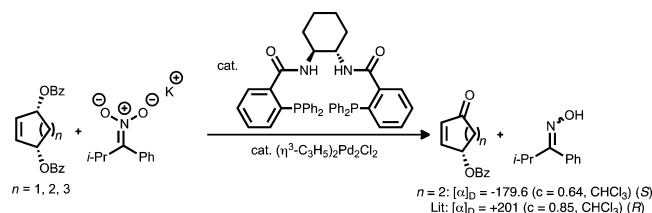
Correction to “Palladium-Catalyzed Chemo- and Enantioselective Oxidation of Allylic Esters and Carbonates”

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S Supporting Information

In the table of contents graphic and Supporting Information, the absolute stereochemistry of the products for Table 2, entries 1–3 (pages S17–S19) should be revised, based on evaluation of the optical rotation data for the product in entry 2. The obtained rotation for this compound, $[\alpha]_D = -179.6$ ($c = 0.64$, CHCl_3), is of comparable magnitude but opposite sign to the literature value for (*R*)-4-benzoyloxycyclohex-2-enone, $[\alpha]_D^{20} = +201$ ($c = 0.85$, CHCl_3).¹ Thus, the corrected stereochemical assignment is (*S*)-4-benzoyloxycyclohex-2-enone. This analysis was performed incorrectly in the Supporting Information, and the incorrect structure was drawn on page S17. By analogy, the structures for entries 1 and 3 in the Supporting Information should also be changed to the (*S*) products. However, the correct structures are drawn in the Communication itself (page 2541, Table 2, entries 1–3). Notably, this corrected sense of stereinduction is in line with empirical models for π -allylpalladium reactions.²



■ ASSOCIATED CONTENT

S Supporting Information

Experimental procedures and characterization data for all new compounds (corrected). This material is available free of charge via the Internet at <http://pubs.acs.org>.

■ REFERENCES

- (1) Yu, L.; Zhang, R.; Wang, Z. *J. Chem. Soc., Perkin Trans. 1* **2001**, 2958.
- (2) Trost, B. M.; Van Vranken, D. L. *Chem. Rev.* **1996**, 96, 395.