

Correction to “3-Fluoro-4-hydroxyprolines: Synthesis, Conformational Analysis, and Stereoselective Recognition by the VHL E3 Ubiquitin Ligase for Targeted Protein Degradation”

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S Supporting Information

Page 9302 (bottom of left column) and Supporting Information (page S10). The concentration of the silyl enol ether **2** was incorrectly quoted as 0.67 M. The correct figure is 0.067 M as follows. The complete corrected Supporting Information is available.

“Under optimized conditions, the silyl enol ether **2** and Selectfluor were dissolved in anhydrous acetonitrile (0.067 and 0.1 M, respectively) and pumped at a rate of 0.77 mL/min in a 10 mL flow reactor”

Page 9310, Figure 7D. It has come to our attention that two values of compound concentrations in the horizontal axis of Figure 7D were not correct. The data analysis and conclusions are not affected by this transcription error. The figure with the corrected values of concentrations in panel D is shown below.

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/jacs.9b03833.

Supplementary results (Tables S1–S6 and Figures S1–S5); materials and methods section; supplementary references; crystal structures have been deposited in the PDB with accession codes 6GFX, 6GFY, and 6GFZ for VBC in complex with compounds **13a**, **14a**, and **14b**, respectively; data collection and refinement statistics; representative ITC titrations; NMR spectra of key intermediates and final compounds (PDF)

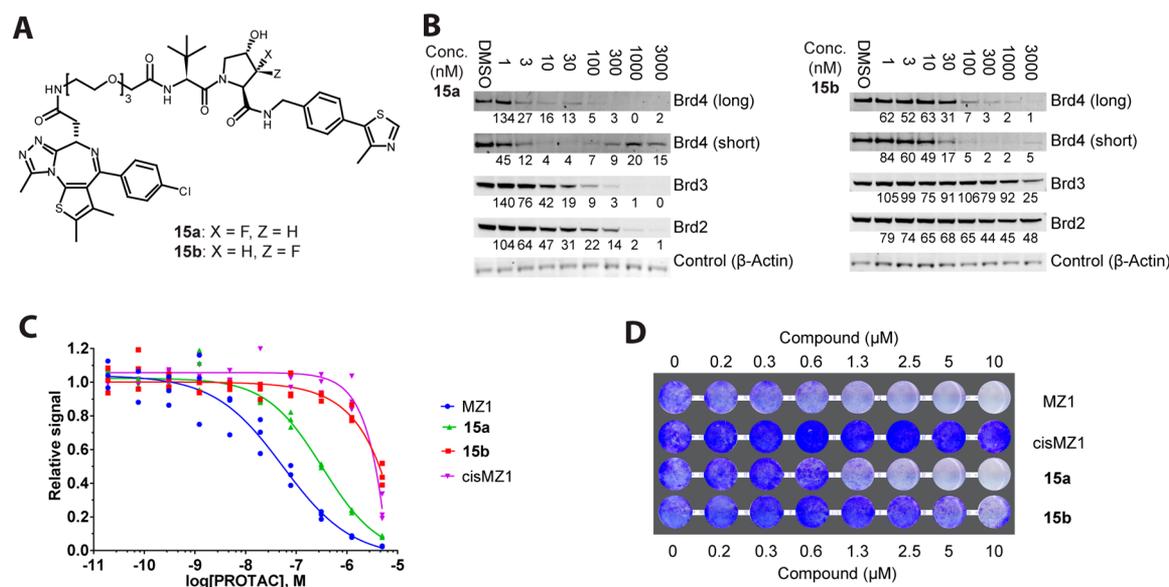


Figure 7. BET protein degradation induced by F-Hyp-containing PROTACs. (A) Chemical structure of PROTACs **15a,b**. (B) HeLa cells were treated with **15a** or **15b** and vehicle control (0.01% DMSO) for 24 h. Abundance of individual BET protein was analyzed by Western blotting using corresponding specific antibodies after SDS-PAGE. (C) Antiproliferative activity of compounds **15a,b**, MZ1, and inactive epimer cisMZ1. MV4;11 cells were treated with compounds for 72 h prior to quantitation of cell proliferation using the CellTiter-Glo luminescent cell viability assay. The relative signal reflects luminescence values normalized to DMSO control. (D) AS49 cells stained with crystal violet dye, following a 7-day incubation with compounds **15a,b**, MZ1, and inactive epimer cisMZ1. Dark background was added to improve legibility.

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