A comparison of enzymatic degradation of *myo-* and *scyllo*inositol hexaskisphosphate

Tony Hollenback 1* Mingjing Sun 1,2 Charles Brearley 3 and deb Jaisi^1

¹Department of Plant and Soil Science, University of Delaware, Newark, Delaware (*correspondence: tonyh@udel.edu)

²Department of Physical Sciences, Emporia State University, Emporia, Kansas

³School of Biological Sciences, University of East Anglia, Norwich, UK

Abstract

Inositol phosphates are the most dominant form of organic phosphorus compounds in soils. Four stereoisomers of inositol hexakisphosphate (IP₆) (myo-, scyllo-, D-chiro-, and neo-) are known to exist in the environment, with myo-IP6 being the most common. Inositiol hexakisphosphates are less labile than other organic P compounds and hence accumulate in the environment, with scyllo-IP6 presumed to be more recalcitrant than myo-IP6. Understanding the relative rates of degradation of various stereoisomers allows quantification of their relative impacts on water quality. We employed ion chromatography and NMR methods to quantify the products of scyllo-inositol hexakisphosphate degradation in the presence of phytase enzymes (from Apergilus niger and wheat) and to compare the degradation pathways to those of myo-IP61. Based on the decay characteristics and isomer composition, our data supports the working hypothesis that scyllo-IP6 decays more slowly than myo-IP6 under the same conditions. Furthermore, the products of scyllo-IP₆ degradation are unique to those of myo-IP6. Additional degradation studies on other enzymes including enzyme specificity of different stereoisomers are underway. These findings fill in knowledge gaps regarding the environmental fate and residence time of these common organic phosphorus compounds.

[1] Sun et al. (2017) Soil Sci. Soc. Am. J. 81, 61-75