ABSTRACT

TITLE:	Synthesis of Haploscleridamine and Biologically Significant Analogs
AUTHOR(S):	Harrison, Jacob A.; Absher, Nathan; Metze, Bryan; Rodriguez, Lydia; Chadeayne, Devon; Wegener, Aaron
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ABSTRACT: Haploscleridamine, a molecule isolated from a sponge of the order Haplosclerida, is a member of a group of natural products that selectively inhibits the E1 ubiquitin pathway and thus shows promise as an anti-cancer drug. We report the development of a reliable synthetic route to Haploscleridamine.