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Douglas Armstrong
Olivet Nazarene University, darmstrg@olivet.edu

Logan Smith

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Progress Toward the Development of a “Catch/Release” Strategy for Isolating Salvinorin A From Plant Materials, Including Microwave Promotion  

Dr. Douglas Armstrong and Logan Smith  
Olivet Nazarene University  
Bourbonnais, IL 60914  
darmstrg@olivet.edu/815-939-5393  
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Proposal and Background

We are investigating using the Diels-Alder (DA) and the retro-Diels-Alder (RDA) reactions as a potential “catch/release” strategy for isolating Salvinorin A from plant samples (Salvia divinorum), hopefully to improve the current method of isolating it, which is very long and laborious. The background for our work is related to the work of Tom Prisinzano (1). Erin Carlson developed a “catch/release” strategy for isolating natural products containing –OH groups, in which alcohols, or phenols, or carboxylic acids are bound to a polymer, which is then rinsed with certain solvents, and then the original compound is released (2). Yli-Kauhaluoma published a review of many DA reactions carried out on solid supports (3).

The synthesis of Salvinorin A is also very long (20 or more steps). (4)

Discussion

Salvinorin A has analgesic activity by activating the kappa opioid receptor, not the mu receptor (1), and thus it has potential for being developed into a non-addicting analgesic.

Strategy and Progress Report

Our strategy focuses on the DA reaction involving the furan ring of Salvinorin A (Fig. 1), as the “diene”, with polymer-bound DEAD (Fig. 2) (diethyl azodicarboxylate), as the dienophile, which is extremely expensive. See below.

Hopefully, our expected product (polymer-bound Salvinorin A, Fig. 3), will allow extraneous compounds and other substances to be rinsed away with suitable solvents, and then hopefully RDA can produce quite pure Salvinorin A. Progress so far includes the IR of the product which lends support to that product, but which we have not had time to fully prove yet.

Prior to doing the above work, we did a “pilot” study, in which we ran DA reactions on Salvinorin A with various dienophiles. One example is the closely related 4-phenyl-1,2,4-triazoline-3,5-dione, PTAD, below, with the expected product shown below, for which we have some evidence (e.g., red color of PTAD disappearing, even at room temperature, and the disappearance of the 1745 cm⁻¹ IR peak of PTAD), but not enough time for complete proof of the product.

Future Research

We now need to finish the proof of the polymer-bound Salvinorin A, and then do the RDA reaction on it. If that fails, we will try the same process using polymer-bound dimethyl acetylenedicarboxylate.

Another future endeavor would be to treat polymer-bound DEAD with an extract of the plant (Salvia divinorum) which was unavailable to us.

References


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