## **Product** Data Sheet

## Debromohymenialdisine

Cat. No.:HY-113632CAS No.:75593-17-8Molecular Formula: $C_{11}H_{11}N_sO_2$ Molecular Weight:245.24Target:Raf; MEK

Pathway: MAPK/ERK Pathway

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

## **BIOLOGICAL ACTIVITY**

Description	Debromohymenialdisine (10Z-Debromohymenialdisine) is a pyrrole alkaloid. Debromohymenialdisine has moderate inhibitory activity with an IC <sub>50</sub> value of 881 nM in the initial Raf/MEK-1/MAPK signaling cascade assay. Debromohymenialdisine can be used for the research of proliferation and differentiation <sup>[1]</sup> .
IC <sub>50</sub> & Target	IC50: 881 nM (in Raf/MEK-1/MAPK signaling cascade assay) <sup>[1]</sup> .
In Vitro	Debromohymenial disine has moderate inhibitory activity with an IC $_{50}$ value of 881 nM in the initial Raf/MEK-1/MAPK signaling cascade assay <sup>[1]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **REFERENCES**

[1]. Deniz Tasdemir, et al. Aldisine alkaloids from the Philippine sponge Stylissa massa are potent inhibitors of mitogen-activated protein kinase kinase-1 (MEK-1). J Med Chem. 2002 Jan 17;45(2):529-32.

Caution: Product has not been fully validated for medical applications. For research use only.

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