PRODUCT INFORMATION



ADB-BUTINACA-do

Item No. 38341

| Formal Name: | N-[(1S)-1-(aminocarbonyl)-2,2-dimethylpropyl]- 1-butyl-d ₉ -1H-indazole-3-carboxamide | |
|------------------|---|---|
| MF: | $C_{18}H_{17}D_{9}N_{4}O_{2}$ | |
| FW: | 339.5 | |
| Chemical Purity: | ≥98% (ADB-BUTINACA) | $\sim \sim $ |
| Deuterium | | a´à a´à N |
| Incorporation: | ≥99% deuterated forms (d ₁ -d ₉); ≤1% d ₀ | H ^N , NH ₂ |
| UV/Vis.: | λ _{max} : 210, 303 nm | |
| Supplied as: | A crystalline solid | |
| Storage: | -20°C | |
| Stability: | ≥2 years | 1 |

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Description

ADB-BUTINACA-d_o (Item No. 38341) is intended for use as an internal standard for the quantification of ADB-BUTINACA (Item No. 29350) by GC- or LC-MS. The accuracy of the sample weight in this vial is between 5% over and 2% under the amount shown on the vial. If better precision is required, the deuterated standard should be quantitated against a more precisely weighed unlabeled standard by constructing a standard curve of peak intensity ratios (deuterated versus unlabeled).

ADB-BUTINACA is categorized as a synthetic cannabinoid.¹ ADB-BUTINACA-d_o is regulated as a Schedule I compound in the United States. This product is intended for research and forensic applications.

Reference

1. Sparkes, E., Cairns, E.A., Kevin, R.C., et al. Structure-activity relationships of valine, tert-leucine, and phenylalanine amino acid-derived synthetic cannabinoid receptor agonists related to ADB-BUTINACA, APP-BUTINACA, and ADB-P7AICA. RSC Med. Chem. 13(2), 156-174 (2021).

WARNING THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

SAFETY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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