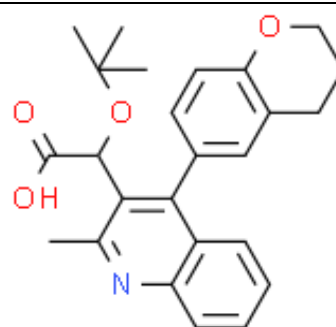


Product Data Sheet

| | | | |
|--|--|--------------|----------|
| Cas No.: | 1416258-16-6 | Cat. No: | PL04811 |
| Product Name: | (±)-BI-D | | |
| Product synonym: | (±)-BI-D | | |
| Chemical name: | (±)-BI-D | | |
| MF: | C25H27NO4 | FW: | 405.4862 |
| Purity: | ≥98% | Batch No.: | - |
| Storage: | | | |
| Structural formula: |  | | |
| λmax: | - | Formulation: | - |
| Solubility : | | | |
| SMILES : | O(C(C([H])([H])([H])(C([H])([H])([H])(C([H])([H])([H])(C([H])(C(=O)O[H])C1C(C([H])([H])([H])=NC2=C([H])C([H])=C([H])C([H])=C2C=1C1C([H])=C([H])C2=C(C=1[H])C([H])([H])C([H])([H])C([H])([H])O2 | | |
| InChI Code: | - | | |
| InChI Key: | | | |
| WARNING This product is not for human or veterinary use. | | | |

Product Description

(±)-BI-D 是HIV整合酶（integrase）变构抑制剂，作用于整合酶与LEDGF/p75结合部位。

| | |
|-------|---|
| 生物活性 | (±)-BI-D is a potent ALLINI(An allosteric IN inhibitor) that binds integrase at the LEDGF/p75 binding site. IC ₅₀ value: 2.4–2.9 μM(HIV-Luc infection of WT and Hdgfrp2 KO cells) [1] Target: integrase inhibitor in vitro: Approximately 2.4–2.9 μM of BI-D was required to inhibit 50% of HIV-Luc infection of WT and Hdgfrp2 KO cells, while the IC ₅₀ decreased dramatically, to 160–200 nM, in Psp1 and double-KO cells [1]. |
| 包装储存 | Powder -20°C 3 years; 4°C 2 years |
| 溶解度数据 | In Vitro: DMSO : ≥ 100 mg/mL (246.62 mM)配制储备液 |