



## Product Manual

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## Ritonavir-d6(利托那韦 d6)

## 产品描述

Ritonavir-d6 is intended for use as an internal standard for the quantification of ritonavir by GC- or LC-MS. Ritonavir is an HIV protease inhibitor. It inhibits recombinant HIV-1 protease by 79% when used at a concentration of 0.5 nM. It inhibits HIV-13B-induced cell death in MT-4 human T cell leukemia cells (EC50 = 25 nM) as well as cell death induced by HIV-1LAI, HIV-2ROD, and HIV-2EHO in human MT-2 cells (IC50s = 0.045, 0.13, and 0.24  $\mu$ M, respectively). Ritonavir also inhibits the cytochrome P450 (CYP) isoform CYP3A (IC50 = 0.14  $\mu$ M). It inhibits CYP-mediated oxidative metabolism of the HIV protease inhibitors saquinavir , indinavir , nelfinavir , and amprenavir in rat and human liver microsomes in a concentration-dependent manner. Ritonavir (10 mg/kg) also prevents decreases in plasma levels of these four compounds in rats. Formulations containing ritonavir have been used in the treatment of HIV-1 infection.

## 化学数据

Catalog No.: MR720201 Cas No.: 1217720-20-1 分子式: C<sub>37</sub>H<sub>42</sub>D<sub>6</sub>N<sub>6</sub>O<sub>5</sub>S<sub>2</sub>

分子量: **727** Purity: >99.00%

溶解度: Chloroform: Slightly Soluble, DMSO: Slightly Soluble, Methanol: Slightly Soluble

储存条件: Store at -20°C

**General tips:** For obtaining a higher solubility , please warm the tube at 37  $^{\circ}{\mathbb{C}}$  and shake it in the

ultrasonic bath for a while.

Shipping Condition: Evaluation sample solution: ship with blue ice

All other available size: ship with RT , or blue ice upon request  $% \left( 1\right) =\left( 1\right) \left( 1\right) \left$ 

Chemical Structure:



