

Produktinformation



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Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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PRODUCT INFORMATION



Maraviroc-d₆ Item No. 28767

Formal Name: 4,4-difluoro-N-[(1S)-3-[(3-exo)-3-[3-methyl-

> 5-(1-(methyl-d₃)ethyl-2,2,2-d₃)-4H-1,2,4triazol-4-yl]-8-azabicyclo[3.2.1]oct-8-yl]-1phenylpropyl]-cyclohexanecarboxamide

C₂₉H₃₅D₆F₂N₅O 519.7 MF:

FW:

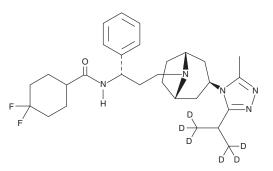
Chemical Purity: ≥98% (Maraviroc)

Deuterium

Incorporation: \geq 99% deuterated forms (d₁-d₆); \leq 1% d₀

Supplied as: A solid -20°C Storage: Stability: ≥2 years

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



Laboratory Procedures

Maraviroc-d₆ is intended for use as an internal standard for the quantification of maraviroc (Item No. 14641) 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).

Maraviroc- d_6 is supplied as a solid. A stock solution may be made by dissolving the maraviroc- d_6 in the solvent of choice, which should be purged with an inert gas. Maraviroc-d_k is slightly soluble in chloroform.

Description

Maraviroc is an antagonist of the chemokine (C-C motif) receptor 5 (CCR5; K; = 0.24 nM for the rhesus monkey recombinant receptor).1 It inhibits binding of chemokine (C-C motif) ligand 3 (CCL3), CCL4, and CCL5 to HEK293 cell membranes expressing CCR5 (IC $_{50}$ s = 3.3, 7.2, and 5.2 nM, respectively) and inhibits HIV-1 binding to CCR5 via glycoprotein 120 (gp120) and gp160 (IC50S = 11 and 0.22 nM, respectively).² Maraviroc is selective for CCR5 over CCR1, -2, -3, -4, -7, and -8, and chemokine (C-X-C motif) receptor 1 (CXCR1) and CXCR2 in a panel of immunological assays for ligand-induced cell chemotaxis and ligand-receptor binding (IC $_{50}$ s = >10 μ M for all). It has antiviral activity against laboratory and clinical isolates of CCR5-tropic (IC $_{50}$ s = 0.1-1.1 nM), but not CXCR4-tropic or dual tropic (IC₅₀S = >10 μ M for all), HIV-1 in isolated human peripheral blood mononuclear cells (PBMCs). Maraviroc prevents infection upon exposure to HIV-1 in humanized RAG-hum mice when administered at a dose of 62 mg/kg.3

References

- 1. Yuan, Y., Arnatt, C.K., El-Hage, N., et al. Medchemcomm. 4(5), 847-851 (2013).
- 2. Dorr, P., Westby, M., Dobbs, S., et al. Antimicrob. Agents Chemother. 49(11), 4721-4732 (2005).
- 3. Neff, C.P., Ndolo, T., Tandon, A., et al. PLoS One 5(12), e15257 (2010).

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

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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