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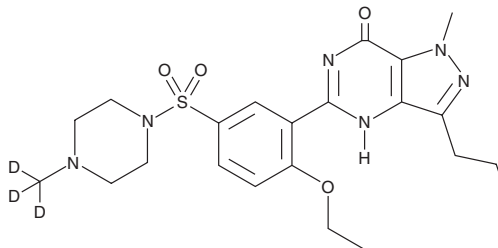


PRODUCT INFORMATION



Sildenafil-d₃ Item No. 18261

CAS Registry No.: 1126745-90-1
Formal Name: 5-[2-ethoxy-5-[[4-(methyl-d₃)-1-piperazinyl]sulfonyl]phenyl]-1,6-dihydro-1-methyl-3-propyl-7H-pyrazolo[4,3-d]pyrimidin-7-one
MF: C₂₂H₂₇D₃N₆O₄S
FW: 477.6
Chemical Purity: ≥98% (Sildenafil)
Deuterium Incorporation: ≥99% deuterated forms (d₁-d₃); ≤1% d₀
Supplied as: A solid
Storage: -20°C
Stability: ≥2 years



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

Laboratory Procedures

Sildenafil-d₃ is intended for use as an internal standard for the quantification of sildenafil (Item Nos. 10008671 | 14008) 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).

Sildenafil-d₃ is supplied as a solid. A stock solution may be made by dissolving the sildenafil-d₃ in the solvent of choice. Sildenafil-d₃ is soluble in organic solvents such as DMSO and dimethyl formamide, which should be purged with an inert gas. The solubility of sildenafil-d₃ in these solvents is approximately 10 and 5 mg/ml, respectively.

Description

Sildenafil-d₃ is intended for use as an internal standard for the quantification of sildenafil (Item Nos. 10008671 | 14008) by GC- or LC-MS. Sildenafil is a potent inhibitor of phosphodiesterase 5 (PDE5) with IC₅₀ values of 3.6 and 3 nM for PDE5 activity in isolated rabbit platelets and human corpus cavernosum, respectively.¹ It is selective for PDE5 over PDE1 and PDE3 (IC₅₀s = 0.26 and 65 μM, respectively). Sildenafil reverses glucose-induced decreases in angiotensin 1 (Ang1) expression and reduction of capillary-like tube formation by mouse dermal endothelial cells *in vitro* and increases the number of functional blood vessels and regional blood flow in the sciatic nerve in a *db/db* mouse model of diabetic peripheral neuropathy.² It increases the ratio of maximum intracavernosal pressure to mean arterial blood pressure (ICP/MAP), a measure of erectile function, in castrated rats when administered at a dose of 20 mg/kg per day.³ Sildenafil (0.5 mg/kg) also reduces cardiac arrest and resuscitation-induced increases in angiotensin II (Item No. 17150), angiotensin converting enzyme (ACE), ACE2, and various angiotensin receptors and increases survival in a porcine model of ischemia/reperfusion injury.⁴ Formulations containing sildenafil have been used in the treatment of erectile dysfunction, pulmonary arterial hypertension, and high-altitude pulmonary edema associated with altitude sickness.

References

1. Terrett, N.K., Bell, A.S., Brown, D., *et al.* *Bioorg. Med. Chem. Lett.* **6**(15), 1819-1824 (1996).
2. Wang, L., Chopp, M., Szalad, A., *et al.* *PLoS One* **10**(2), e0118134 (2015).
3. Mulhall, J.P., Verma, N., Deveci, S., *et al.* *BJU Int.* **113**(4), 656-661 (2014).
4. Wang, G., Zhang, Q., Yuan, W., *et al.* *Int. J. Mol. Sci.* **16**(11), 27015-27031 (2015).

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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