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**Product Data Sheet**

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Product Name: Sildenafil mesylate

Cat. No.: GC15037

**Chemical Properties**

Cas. No. 1308285-21-3

Chemical Name 5-(2-ethoxy-5-((4-methylpiperazin-1-yl)sulfonyl)phenyl)-1-methyl-3-propyl-1H-pyrazolo[4,3-d]pyrimidin-7(6H)-one methanesulfonate

SMILES CCCC(C(N=C(N1)C2=C(OCC)C=CC(S(N3CCN(CC3)C)(=O)=O)=C2)=C4C1=O)=NN4C.CS(O)(=O)=OFormula  $C_{23}H_{34}N_6O_7S_2$  M.Wt 570.68Solubility  $\geq 57.1\text{mg/mL}$  in DMSO Storage Store at  $-20^\circ\text{C}$ General tips For obtaining a higher solubility, please warm the tube at  $37^\circ\text{C}$  and shake it in the ultrasonic bath for a while. Stock solution can be stored below  $-20^\circ\text{C}$  for several months.

Shipping Condition Evaluation sample solution: ship with blue ice. All other available size: ship with RT, or blue ice upon request.

Structure **Background**

IC50: 3.5 nM

Sildenafil mesylate is an inhibitor of cyclic guanosine monophosphate (cGMP)-specific phosphodiesterase type 5 (PDE5).

PDE5 is the predominant phosphodiesterase in the corpus cavernosum. The catalytic site of PDE5 degrades cGMP, and PDE5 inhibitors can potentiate endogenous increases in cGMP.

In vitro: Sildenafil mesylate had been identified as a potent PDE5 reversible and selective inhibitor. Sildenafil enhanced sodium nitroprusside- or transmural electrical stimulation-induced relaxation of precontracted corpus cavernosum muscle strips in organ baths. Sildenafil also increased intracellular cGMP concentrations in cultured smooth muscle cells treated with sodium nitroprusside [1].

**Caution: Product has not been fully validated for medical applications. For research use only.**

Tel: (909) 407-4943 Fax: (626) 353-8530 E-mail: tech@glpbio.com

Address: 10292 Central Ave. #205, Montclair, CA, USA

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In vivo: In anesthetized dogs, sildenafil could enhance the erectile function following pelvic nerve stimulation as measured by increased intracavernosal pressure [1].

Clinical trial: Ten previous randomised controlled trials had been conducted. Dose optimisation resulted in >60% of attempts at sexual intercourse, which was successful in 49% of men. Sildenafil treatment related adverse events occurred in 30% of men compared with 11% on placebo. Dose optimisation of sildenafil showed efficacy equivalent to the highest fixed doses, and adverse events equivalent to the lowest fixed doses [2].

### References:

[1] Nehra A, Colreavy F, Khandheria BK, Chandrasekaran K. Sildenafil citrate, a selective phosphodiesterase type 5 inhibitor: urologic and cardiovascular implications. *World J Urol.* 2001 Feb;19(1):40-5.

[2] Moore RA, Edwards JE, McQuay HJ. Sildenafil (Viagra) for male erectile dysfunction: a meta-analysis of clinical trial reports. *BMC Urol.* 2002 May 22;2:6.

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