Proteins

TPN729

Cat. No.: HY-116808 936951-20-1 CAS No.: Molecular Formula: $C_{25}H_{36}N_{6}O_{4}S$ Molecular Weight: 516.66

Phosphodiesterase (PDE) Target: Pathway: Metabolic Enzyme/Protease

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

Product Data Sheet

BIOLOGICAL ACTIVITY

Description TPN729 is a selective phosphodiesterase type 5 (PDE5) inhibitor with an IC $_{50}$ value of 2.28 nM. TPN729 affects erectile function and it can be used for the research of erectile dysfunction^[1].

IC₅₀ & Target PDE5 PDE6 PDE1 PDE4

2.28 nM (IC₅₀) 45.2 nM (IC₅₀) 566 nM (IC₅₀) 834 nM (IC₅₀)

PDE11 6090 nM (IC₅₀)

TPN729 (1.25, 2.5 and 5.0 mg/kg; intraduodenal injection, once) increases intracavernous pressure/blood pressure (ICP/BP) In Vivo

with the dose of 5.0 mg/kg at all time points and with the dose of 2.5 mg/kg at 75, 90, 105, and 120 minutes time points^[1]. TPN729 (5.0 μg/kg; i.v., once) significantly increases ICP and ICP/BP of male Beagle dogs^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Male Sprague-Dawley rats with corpora cavernosa complete $\operatorname{exposure}^{[1]}$
Dosage:	1.25, 2.5 and 5.0 mg/kg
Administration:	Intraduodenal (i.d.) injection; 1.25, 2.5 and 5.0 mg/kg, once
Result:	Dose-dependently increased the maximum intracavernous pressure (ICP) and ICP/blood pressure (BP).

REFERENCES

[1]. Wang Z, et al. The selectivity and potency of the new PDE5 inhibitor TPN729MA. J Sex Med. 2013 Nov;10(11):2790-7.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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