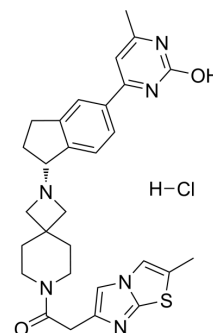


PF-6870961 hydrochloride

Cat. No.:	HY-153095A
CAS No.:	2857112-07-1
Molecular Formula:	C ₂₉ H ₃₃ ClN ₆ O ₂ S
Molecular Weight:	565.13
Target:	GHSR
Pathway:	GPCR/G Protein
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	PF-6870961 hydrochloride is an inverse agonist of GHSR1a with K _i values of 73.6 nM (human GHSR), 239 nM (rat GHSR), and 217 nM (dog GHSR), respectively. PF-6870961 hydrochloride inhibits the constitutive GHSR1a-induced IP accumulation with an IC ₅₀ value of 300 nM. PF-6870961 hydrochloride also inhibits constitutive GHSR1a β-arrestin mobilization with an IC ₅₀ value of 1.10 nM ^[1] .								
In Vivo	<p>PF-6870961 hydrochloride (40 mg/kg; i.p.) suppresses food intake in both satiated and fasted rats^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Satiated and fasted rats mode^[1]</td> </tr> <tr> <td>Dosage:</td> <td>2.5 mg/kg, 10 mg/kg, and 40 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Intraperitoneal injection</td> </tr> <tr> <td>Result:</td> <td>Significantly reduced food intake at 40 mg/kg.</td> </tr> </table>	Animal Model:	Satiated and fasted rats mode ^[1]	Dosage:	2.5 mg/kg, 10 mg/kg, and 40 mg/kg	Administration:	Intraperitoneal injection	Result:	Significantly reduced food intake at 40 mg/kg.
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REFERENCES

[1]. Deschaine SL, et al. Initial pharmacological characterization of a major hydroxy metabolite of PF-5190457: inverse agonist activity of PF-6870961 at the ghrelin receptor. J Pharmacol Exp Ther. 2023 Jan 11:JPET-AR-2022-001393.

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

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