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Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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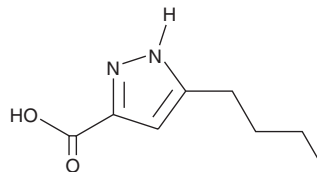
PRODUCT INFORMATION



LUF6283

Item No. 15423

CAS Registry No.: 92933-48-7
Formal Name: 5-butyl-1H-pyrazole-3-carboxylic acid
MF: $C_8H_{12}N_2O_2$
FW: 168.2
Purity: $\geq 98\%$
UV/Vis.: λ_{max} : 218 nm
Supplied as: A crystalline 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

LUF6283 is supplied as a crystalline solid. A stock solution may be made by dissolving the LUF6283 in the solvent of choice. LUF6283 is soluble in organic solvents such as ethanol and DMSO, which should be purged with an inert gas. The solubility of LUF6283 in these solvents is approximately 16 mg/ml.

Description

LUF6283 is a partial agonist of GPR109A/HCA₂ G protein-coupled receptors ($K_i = 0.55 \mu\text{M}$ in a radioligand binding assay).¹ It stimulates [³⁵S]-GTPγS binding and increases phosphorylation of ERK in HEK293T cells expressing GPR109A/HCA₂ receptors (EC_{50} s = 3.1 and 0.32 μM , respectively). *In vivo*, LUF6283 (400 mg/kg per day) reduces plasma levels of VLDL and hepatic expression of apolipoprotein B in mice.

Reference

1. Li, Z., Blad, C.C., van der Sluis, R.J., *et al.* Effects of pyrazole partial agonists on HCA₂-mediated flushing and VLDL-triglyceride levels in mice. *Br. J. Pharmacol.* **167**(4), 818-825 (2012).

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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