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Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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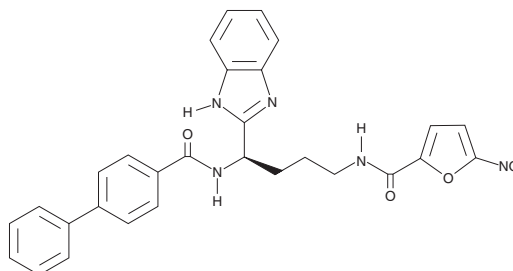
PRODUCT INFORMATION



LB244

Item No. 39832

Formal Name: (R)-N-(4-([1,1'-biphenyl]-4-carboxamido)-4-(1H-benzo[d]imidazol-2-yl)butyl)-5-nitrofuran-2-carboxamide
MF: C₂₉H₂₅N₅O₅
FW: 523.5
Purity: ≥95%
Supplied as: A solid
Storage: -20°C
Stability: ≥3 years



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

Laboratory Procedures

LB244 is supplied as a solid. A stock solution may be made by dissolving the LB244 in the solvent of choice, which should be purged with an inert gas. LB244 is soluble in chloroform.

Description

LB244 is an irreversible STING antagonist.¹ It binds to STING in a cell-free assay and inhibits STING activation induced by the STING agonist diABZI in cells (EC₅₀ = 0.8 nM). LB244 (1 μM) inhibits diABZI-induced STING oligomerization and phosphorylation of IFN regulatory factor 3 (IRF3) and TANK-binding kinase 1 (TBK1) in primary mouse bone marrow-derived macrophages (BMDMs). *In vivo*, LB244 (5 mg/kg, i.p.) reduces diABZI-induced increases in serum IL-6 and IFN-β levels in mice.

Reference

1. Barasa, L., Chaudhuri, S., Zhou, J.Y., *et al.* Development of LB244, an irreversible STING antagonist. *J. Am. Chem. Soc.* **145**(37), 20273-20288 (2023).

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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