

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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PRODUCT INFORMATION



Lipid Catechol

Item No. 38665

Formal Name: ditetradecyl (((2-(diethylamino)

ethyl)amino)(3,4-dihydroxyphenyl)

methyl)phosphonate

MF: $C_{41}H_{79}N_2O_5P$

FW: 711.1 **Purity:** ≥95% Supplied as: A solid -20°C Storage: Stability: ≥2 years

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

Laboratory Procedures

Lipid catechol is supplied as a solid. A stock solution may be made by dissolving the lipid catechol in the solvent of choice, which should be purged with an inert gas. Lipid catechol is soluble in organic solvents such as ethanol and DMSO. The solubility of lipid catechol in these solvents is approximately 1 mg/ml.

Description

Lipid catechol is a lipid that contains an α-aminophosphonate group, two 14-carbon acyl chains, and a catechol ring that forms a covalent bond with boronic acid-containing compounds to form lipid prodrug nanoassemblies (LPNA).1 LPNAs composed of lipid catechol conjugated to phenylboronic acid-modified ciprofloxacin (CIP-PBA) inhibit the formation of, and disrupt preformed, S. aureus biofilms and eradicate staphylococci in a mouse model of peritoneal S. aureus infection. LPNAs composed of lipid catechol conjugated to bortezomib (BTZ) reduce tumor growth and increase survival in a 4T1 murine mammary carcinoma model.

Reference

1. Ding, Y., Hu, X., Piao, Y., et al. Lipid prodrug nanoassemblies via dynamic covalent boronates. ACS Nano 17(7), 6601-6614 (2023).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

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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