mPEG-DSPE, MW 2000

| Cat. No.: CAS No.: Molecular Formula: Target: Pathway: | HY-139385 147867-65-0 (C ₂ H ₄ O) _n C ₄₃ H ₈₄ NO ₁₀ P Liposome Metabolic Enzyme/Protease | |
|--|--|--|
| Storage: | -20°C, protect from light, stored under nitrogen * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light, stored under nitrogen) | |

| SOLVENT & SOLUE | |
|-----------------|---|
| | |
| In Vitro | DMSO : 50 mg/mL (ultrasonic and warming and heat to 60°C) |
| In Vivo | 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (Infinity mM); Clear solution |
| | 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (Infinity mM); Clear solution |

| BIOLOGICAL ACT | |
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| DIOLOGICALACI | |

Description mPEG-DSPE, MW 2000 is a phospholipid PEG conjugate, has both hydrophilicity and hydrophobility. mPEG-DSPE, MW 2000 is used in the synthesis of liposomes, increases the hydrophilicity on the surface of liposomes, consequently increasing the longevity of liposomes in the blood circulation for the study of anticancer and antimalarial agents^{[1][2]}.

REFERENCES

Page 1 of 1

[1]. Rathore SS, et al. Effect of surface charge and density of distearylphosphatidylethanolamine-mPEG-2000 (DSPE-mPEG-2000) on the cytotoxicity of liposome-entrapped ricin: effect of lysosomotropic agents. Int J Pharm. 2008;350(1-2):79-94.;Hasan GM, et al. Inhibition of the Growth of Plasmodium falciparum in Culture by Stearylamine-Phosphatidylcholine Liposomes. J Parasitol Res. 2011;2011:120462.

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

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Product Data Sheet

