



# SZABO SCANDIC

Part of Europa Biosite

## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten!  
See the following pages for more information!



### Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

### SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

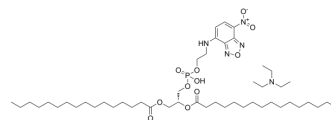
[mail@szabo-scandic.com](mailto:mail@szabo-scandic.com)

[www.szabo-scandic.com](http://www.szabo-scandic.com)

[linkedin.com/company/szaboscandic](https://www.linkedin.com/company/szaboscandic) 

## NBD-PE

<b>Cat. No.:</b>	HY-D1683
<b>CAS No.:</b>	178119-00-1
<b>Molecular Formula:</b>	C <sub>49</sub> H <sub>90</sub> N <sub>5</sub> O <sub>11</sub> P
<b>Molecular Weight:</b>	956.24
<b>Target:</b>	Fluorescent Dye
<b>Pathway:</b>	Others
<b>Storage:</b>	-20°C, protect from light, stored under nitrogen * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light, stored under nitrogen)



## BIOLOGICAL ACTIVITY

<b>Description</b>	NBD-PE is an effective lipid fluorescent probe (Excitation/Emission: 465/535 nm; Color: Green). NBD-PE offers a wide array of applications in membrane and cell biology <sup>[1][2][3]</sup> .
<b>In Vitro</b>	<p>Guidelines (Following is our recommended protocol. This protocol only provides a guideline, and should be modified according to your specific needs).</p> <p>Assay of phospholipid transfer activity<sup>[3]</sup>:</p> <ol style="list-style-type: none"> <li>1. The reaction mixture contains 2 mg mitochondrial protein, 0.02 μmol of NBD-PE, 0.5 mmol of Tris-HCl, pH 7.4, and 0-200 μg of pH 5.1-supernatant protein.</li> <li>2. Samples are incubated at 37°C for 1 hour with gentle agitation.</li> <li>3. The reaction is terminated by placing samples in an ice bath for 10 min, followed by centrifugation at 15,000 g for 5 min.</li> <li>4. The supernatants are decanted, allowed to come to room temperature, and the absorbance (460 nm) and the relative fluorescence at 535 nm (excitation at 465 nm) are determined.</li> <li>5. Blanks containing only NBD-PE and exchange protein are run concomitantly to correct for possible interaction between NBD-PE and exchange protein.</li> <li>6. The percent exchange of NBD-PE is calculated from the differences between the absorbances or relative fluorescence in the supernatant fractions in the presence and absence of exchange protein.</li> </ol> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>

## REFERENCES

- [1]. M Esfahani, et al. 19F-NMR and fluorescence polarization of yeast plasma membrane and isolated lipids. *Biochem Biophys Res Commun*. 1981 Jul 16;101(1):306-11.
- [2]. P S Uster, et al. Resonance energy transfer microscopy: visual colocalization of fluorescent lipid probes in liposomes. *Methods Enzymol*. 1989;171:850-7.
- [3]. J A Monti, et al. Synthesis and properties of a highly fluorescent derivative of phosphatidylethanolamine. *J Lipid Res*. 1978 Feb;19(2):222-8.

---

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: [tech@MedChemExpress.com](mailto:tech@MedChemExpress.com)

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA