

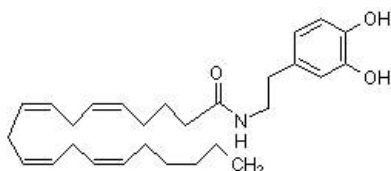
## Product datasheet

# NADA (N-Arachidonyldopamine), endogenous CB1 / TRPV1 agonist ab120099

[1 References](#) [2 图像](#)

### 概述

产品名称	NADA (N-Arachidonyldopamine), endogenous CB1 / TRPV1激动剂
描述	Endogenous CB <sub>1</sub> / TRPV1激动剂
CAS编号	199875-69-9
化学结构	



### 性能

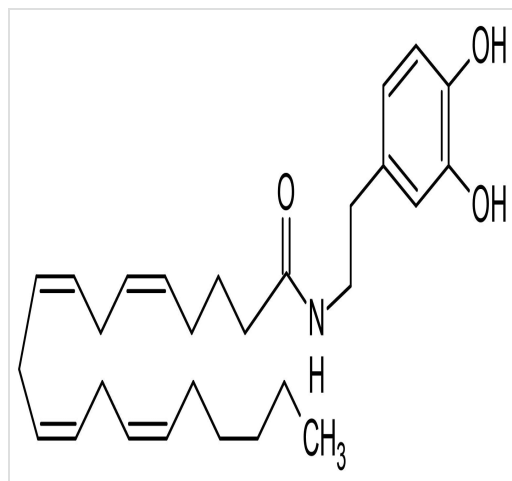
化学名称	(5Z,8Z,11Z,14Z)-N-(3,5-dihydroxyphenethyl)icosa-5,8,11,14-tetraenamide
分子量	439.63
分子式	C <sub>28</sub> H <sub>41</sub> NO <sub>3</sub>
PubChem识别号	5282105
存放说明	Store at -20°C (desiccating conditions).
溶解度概述	Supplied in ethanol (5 mg/ml)
处理	<p>Providing storage is as stated on the product vial and the vial is kept tightly sealed, the product can be stored for up to 6 months. Before use, and prior to opening the vial we recommend that you allow your product to equilibrate to room temperature for at least 1 hour.</p> <p>Refer to SDS for further information.</p> <p>Need more advice on solubility, usage and handling? Please visit our <a href="#">frequently asked questions (FAQ) page</a> for more details.</p>
SMILES	<chem>Oc1ccc(CCNC(=O)CCCC/C=CC/C=CC/C=CC/C=CCCCC)cc1O</chem>
来源	Synthetic

### 应用

“应用说明”部分 下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

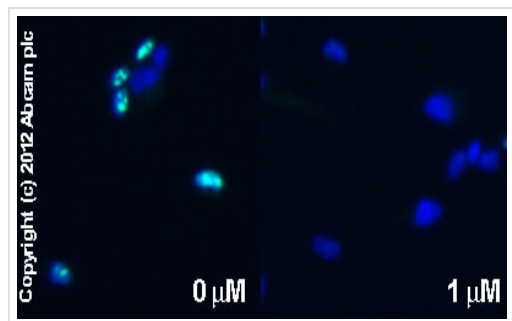
应用	Ab评论	说明
Functional Studies		Use at an assay dependent concentration.

## 图片



Chemical Structure - NADA (N-Arachidonyldopamine), endogenous CB<sub>1</sub> / TRPV1 agonist (ab120099)

2D chemical structure image of ab120099, NADA (N-Arachidonyldopamine), endogenous CB<sub>1</sub> / TRPV1 agonist



Immunocytochemistry/ Immunofluorescence - NADA (N-Arachidonyldopamine), endogenous CB<sub>1</sub> / TRPV1 agonist (ab120099)

**ab15580** staining Ki67 in SK-N-SH cells treated with NADA (N-Arachidonyldopamine) (ab120099), by ICC/IF. Decrease in Ki67 expression correlates with increased concentration of NADA (N-Arachidonyldopamine), as described in literature.

The cells were incubated at 37°C for 10 minutes in media containing different concentrations of ab120099 (NADA (N-Arachidonyldopamine)) in ethanol, fixed with 4% formaldehyde for 10 minutes at room temperature and blocked with PBS containing 10% goat serum, 0.3 M glycine, 1% BSA and 0.1% tween for 2h at room temperature. Staining of the treated cells with **ab15580** (1 μg/ml) was performed overnight at 4°C in PBS containing 1% BSA and 0.1% tween. A DyLight 488 goat anti-rabbit polyclonal antibody (**ab96899**) at 1/250 dilution was used as the secondary antibody. Nuclei were counterstained with DAPI and are shown in blue.

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