

Catalog # 10-1297 Palmitoylethanolamide

CAS# 544-31-0 N-(2-Hydroxyethyl)palmitamide PEA Lot # X101408

Endogenous cannabinoid. Weak ligand for CB₁ ($K_i = 23.8 \mu M$) and CB₂ ($K_i = 13.9 \mu M$) receptors. Inhibits fatty acid amide hydrolase (FAAH) ((IC₅₀ = 5.1 μM). Modulates mast cell activation. Displays immunosuppressant, anti-inflammatory and anti-nociceptive activity.

- 1) De Filippis et al. (2011), Palmitoylethanolamide reduces granuloma-induced hyperalgesia by modulation of mast cell activation in rats; Mol. Pain, **7** 3
- 2) Re et al. (2007), Palmitoylethanolamide, endocannabinoids and related cannabimimetic compounds in protection against tissue inflammation and pain: potential use in companion animals; Vet J., **173** 21
- 3) Lambert et al. (2002), *The palmitoylethanolamide family: a new class of anti-inflammatory agents?;* Curr. Med. Chem., **9** 663

PHYSICAL DATA

Molecular Weight: 299.50
Molecular Formula: C₁₈H₃₇NO₂
Purity: 98% by TLC:

NMR: Conforms
DMSO (up to 25 mg/ml), Ethanol (up to 25 mg/ml)

Physical Description: White solid

Solubility:

Storage and Stability: Store as supplied at -20°C for up to 1 year from the date of purchase.

Solutions in DMSO or ethanol may be stored at -20°C for up to 3 months.

Materials provided by Focus Biomolecules are for laboratory research use only and are not intended for human or veterinary applications.