

Catalog # 10-1435 Amlexanox

CAS# 68302-57-8 2-Amino-7-(1-methylethyl)-5-oxo-5H-[1]benzopyrano[2,3-b]pyridine-3-carboxylic acid Lot # X105810



Improves obesity-related metabolic dysfunction in mouse models.¹ Increases thermogenesis and weight loss, decreases steatosis and improves insulin sensitivity. Inhibits TANK-binding kinase 1 (TBK1) and IKK ϵ (IC₅₀=1.2 μ M).² Clinically useful in the treatment of aphthus ulcers.³ Inhibits prostate tumor metastasis by targeting IKK ϵ /TBK1/NF κ B signaling.⁴

- 1) Koch et al. (2013), Obesity: Teaching an old drug new tricks amlexanox targets inflammation to improve metabolic dysfunction; Nat. Rev. Endocrinol., **9** 185
- Reilly et al. (2013), An inhibitor of the protein kinases TBK1 and IKK-ε improves obesity-related metabolic dysfunctions in mice; Nat. Med., 19 313
- Nasry et al. (2016), Different modalities for treatment of recurrent aphthous stromatitis. A Randomized clinical trial; J. Clin. Exp. Dent. 8 e517
- 4) Cheng *et al.* (2018), *Aphthous ulcer drug inhibits prostate tumor metastasis by targeting IKKε/TBK1/NFκB signaling*; Theranostics **8** 4633

PHYSICAL DATA

Molecular Weight:	298.29
Molecular Formula:	C ₁₆ H ₁₄ N ₂ O ₄
Purity:	>98%
	NMR: (Conforms)
Solubility:	Soluble in DMSO (60 mg/ml). Compound is not soluble in ethanol.
Physical Description:	White solid
Storage and Stability:	Store as supplied at room temperature for up to 2 years from the date of purchase. Store solutions in DMSO at -20°C for up to 3 months.

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