

Catalog # 10-4498 PF-06840003

CAS# 198474**-05-4**

3-(5-Fluoro-1H-indol-3-yl)pyrrolidine-2,5-dione; EOS200271 Lot # FBS1121

PF-06840003 is potent ($IC_{50} = 150$ and 410 nM via two methods) and selective inhibitor of Indoleamine-2,3-dioxygenase (IDO1).¹ In mice with syngeneic tumor grafts, it reduced intratumoral kynurenine levels by over 80%, inhibited tumor growth as monotherapy, and increased the efficacy of anti-PD-L1 therapy.² PF-06840003 treatment combined with GM-CSF-secreting allogenic pancreatic tumor cell vaccine (GVAX) resulted in increased antitumor efficacy in a murine model of pancreatic ductal adenocarcinoma.³ PF-06840003 rescued hippocampal synaptic plasticity and memory function in preclinical models of amyloid and tau pathology via restoration of neuronal metabolic support.⁴

- 1) Crosignani et al. (2017), Discovery of a Novel and Selective Indoleamine-2,3-Dioxygenase (IDO-1) Inhibitor 3-(5-Fluoro-1H-indol-3-yl)pyrrolidine-2,5-dione (EOS200271/PF-06840003) and Its Characterization as a Potential Clinical Candidate; J.Med.Chem. **60** 9617
- Gomes et al. (2018), Characterization of the Selective Indoleamine-2,3-Dioxygenase-1 (IDO1) Catalytic Inhibitor EOS200271/PF-06840003 Supports IDO1 as a Critical Resistance Mechanism to PD-(L)1 Blockade Therapy; Mol.CancerTher. 9 2530
- 3) Blair et al. (2019), IDO1 inhibition potentiates vaccine-induced immunity against pancreatic adenocarcinoma; J.Clin.Invest. 124077
- 4) Minhas et al. (2024), Restoring hippocampal glucose metabolism rescues cognition across Alzheimer's disease pathologies; Science **385** eabm6131

PHYSICAL DATA

Molecular Weight: 232.21

Molecular Formula: $C_{12}H_9FN_2O_2$ Purity: >98% by HPLC

NMR: (Conforms)

Solubility: DMSO (>25 mg/ml)

Physical Description: White solid

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

DMSO may be stored at -20°C for up to 1 month.

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