

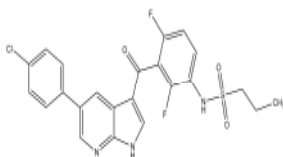
PRODUCT: PLX-4032

ALTERNATE NAME: Vemurafenib; RG7204, RO5185426

CATALOG #: 2235-5, 25

AMOUNT: 5 mg, 25 mg

STRUCTURE:

MOLECULAR FORMULA: C₂₃H₁₈ClF₂N₃O₃S

MOLECULAR WEIGHT: 489.92

CAS NUMBER: 1029872-54-5

APPEARANCE: White to off-white solid

SOLUBILITY: DMSO

PURITY: ≥99%

STORAGE: At -20° C. Protect from light and moisture

DESCRIPTION: Cell-permeable. A potent, selective and ATP-competitive, inhibitor of BRAF(V600E) kinase with potential antineoplastic activity. PLX-40432 selectively binds to the ATP-binding site of BRAF(V600E) kinase and inhibits its activity, which may result in an inhibition of an over-activated MAPK signaling pathway downstream in BRAF(V600E) kinase-expressing tumor cells and a reduction in tumor cell proliferation.

REFERENCE: Lee, J.T., *et al.* (2010). *Pigment Cell Melanoma Res.* **23**, 820-827.

HANDLING: Do not take internally. Wear gloves and mask when handling the product! Avoid contact by all modes of exposure.

RELATED PRODUCTS:

Active RAF1 (Cat. No. 7726-5, 100)

BAY 43-9006 (Cat. No. 1594-25, 100)

BAY 43-9006, Free base (Cat. No. 2142-25, 100)

EZSolution™ BAY 43-9006 (Cat. No. 2031-25)

ZM 336372 (Cat. No. 1789-1, 5)

USAGE: **FOR RESEARCH CH USE ONLY! Not to be used in humans**