

Catalog # 10-4810 NU7441

CAS# 503468-95-9 8-Dibenzothiophen-4-yl-2-morpholin-4-ylchromen-4-one; Ku-57788 Lot # FBS2031

NU7441 is a potent ($IC_{50} = 13$ nM) and selective ATP-competitive inhibitor of DNA-dependent protein kinase (DNA-PK).^{1,2} It increased the toxicity of ionizing radiation and etoposide on SW620 human colon cancer lines via impeding DNA double-strand break repair.³ NU7441 also increased the radio/chemosensitivity of various other cancer cell lines.³⁻⁶ Reduces the frequency of NHEJ repair while enhancing the rate of HDR repair following CRISPR/Cas9-mediated gene editing.⁷

- 1) Leahy et al. (2004), Identification of a highly potent and selective DNA-dependent protein kinase (DNA-PK) inhibitor (NU7441) by screening of chromenone libraries; Bioorg. Med. Chem. Lett. **14** 6083
- 2) Hardcastle et al. (2005), Discovery of potent chromen-4-one inhibitors of the DNA-dependent protein kinase (DNA-PK) using a small-molecule library approach: J. Med. Chem. **48** 7829
- 3) Zhao et al. (2006), Preclinical evaluation of a potent novel DNA-dependent protein kinase inhibitor NU7441: Cancer Res. 66 5354
- 4) Ciszewski et al. (2014), DNA-PK inhibition by NU7441 sensitizes breast cancer cells to ionizing radiation and doxorubicin: Breast Cancer Res. Treat. 143 83
- 5) Yang et al. (2016), NU7441 Enhances the Radiosensitivity of Liver Cancer Cells: Cell Physiol. Biochem. 38 1897
- 6) Geng et al. (2019), DNA-PKcs inhibitor increases the sensitivity of gastric cancer cells to radiotherapy: Oncol. Rep. 42 561
- 7) Robert et al. (2015), Pharmacological inhibition of DNA-PK stimulates Cas9-mediated genome editing: Genome Med., 7 93

PHYSICAL DATA

Molecular Weight: 413.49

Molecular Formula: $C_{25}H_{19}NO_3S$ Purity: 98% by TLC

NMR: (Conforms)

Solubility: DMSO (10 mg/mL with warming)

Physical Description: Off-white to beige 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 3 months.

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