

Catalog # 10-3730 Fasentin

CAS# 392721-37-8 N-[-4-Chloro-3-(trifluoromethyl)phenyl]-3-oxobutanamide Lot # X109521



A novel inhibitor of glucose uptake which acts via inhibition of the glucose transporter GluT1 and GluT4 (IC_{50} =68 μ M).¹ Inhibition of glucose uptake is cell type specific² and is a promising approach to new cancer therapeutics³. Rescues cardiac progenitor cell dysfunction and mitochondrial fission induced by high glucose.⁴ Sensitizes cells to FAS-induced cell death.^{5,6}

- 1) Granchi et al. (2016), Anticancer agents interacting with membrane glucose transporters; Med. Chem. Comm., 7 1716
- Kraus et al. (2018), Targeting glucose transport and the NAD pathway in tumor cells with STF-31: a re-evaluation; Cell. Oncol. (Dordr), 41 485
- 3) Adekola et al. (2012), Glucose transporters in cancer metabolism; Curr. Opin. Oncol., 24 650
- 4) Choi et al. (2016), High Glucose Causes Human Cardiac Progenitor Cell Dysfunction by Promoting Mitochondrial Fission: Role of a GLUT1 Blocker, Prog. Biomol. Ther. (Seoul), **24** 363
- 5) Schimmer et al. (2006), Identification of small molecules that sensitize resistant tumor cells to tumor necrosis factor-family death receptors; Cancer Res., **66** 2367
- 6) Wood et al. (2008), A novel inhibitor of glucose uptake sensitizes cells to FAS-induced cell death; Mol. Cancer Ther., 7 3546

PHYSICAL DATA

Molecular Weight:	279.64
Molecular Formula:	C ₁₁ H ₉ CIF ₃ NO ₂
Purity:	98% by TLC
	NMR: (Conforms)
Solubility:	DMSO (up to 28 mg/ml)
Physical Description:	Off-white solid
Storage and Stability:	Store as supplied desiccated at room temperature for up to 2 years from the date of purchase.
	Solutions in DMSO may be stored at -20°C for up to 3 months.

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