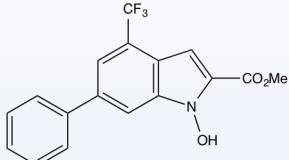
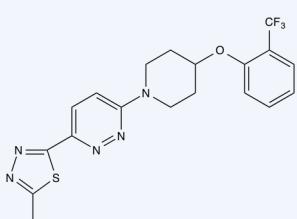


Cancer Cell Metabolism



NHI-2



MF-438

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NHI-2

NHI-2 is a selective inhibitor of Lactate Dehydrogenase A (LDHA) – IC₅₀ *in vitro* = 14.7 µM (NADH), 10.5 µM (Pyruvate)^{1,2}

10-4555 10 mg / \$95.00, 50 mg / \$395.00

CBR-5884

Potent and selective inhibitor of 3-phosphoglycerate dehydrogenase (PHGDH), IC₅₀=33 µM³

10-1588 5 mg / \$70.00, 25 mg / \$280.00

AGI-5198

AGI-5198 is a potent (IC₅₀ = 70 nM) inhibitor of R132H mutant isocitrate dehydrogenase 1(IDH1).⁴

10-4599 5 mg / \$70.00, 25 mg / \$225.00

MF-438

MF-438 is a potent inhibitor of stearoyl-CoA desaturase 1 (SCD1) - IC₅₀ = 2.3 nM.⁵

10-4558 5 mg / \$60.00, 25 mg / \$250.00

CT-2584

Cytotoxic agent that alters phospholipid biosynthesis via inhibition of CTP:choline-phosphate cytidylyltransferase.^{6,7}

10-4560 5 mg / \$125.00, 25 mg / \$475.00

R162

R162 is a selective inhibitor of Glutamate Dehydrogenase 1. Selective over 6-phosphogluconate and fumarate hydratase.⁸

10-4603 10 mg / \$50.00, 50 mg / \$185.00

Lonidamine

Lonidamine is a mitochondrial hexokinase inhibitor.⁹ The disruption of the mitochondrial transmembrane potential has also been linked to its apoptotic effects.¹⁰

10-4563 10 mg / \$45.00, 50 mg / \$175.00

PFK15

PFK15 is a selective inhibitor of 6-Phosphofructo-2-kinase/fructose-2,6-biphosphatase 3(PFKFB3), an enzyme that is constitutively expressed in neoplastic cells and is required for the high glycolytic rate (Warburg effect) of these cells.^{11,12,13}

10-4580 5 mg / \$45.00, 25 mg / \$170.00

CB-839

CB-839 is a potent (IC₅₀ = 24 nM) inhibitor of glutaminase (KGA and GAC).^{14,15}

10-4556 5 mg / \$90.00, 25 mg / \$325.00