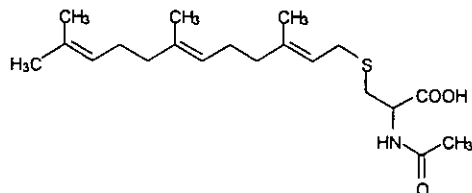


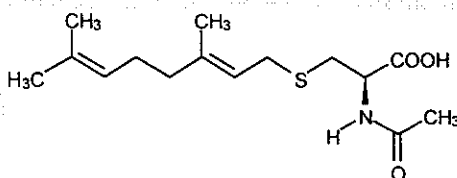
INHIBITORS

N-Acetyl-S-farnesyl-L-cysteine (AFC); Inhibits S-farnesylcysteine methyl transferase ($K_m = 20 \mu\text{M}$) in cell-free extracts and in whole cells. Molecular Formula: $\text{C}_{20}\text{H}_{33}\text{NO}_3\text{S}$; Molecular Weight: 367.5 ; Purity: > 98%



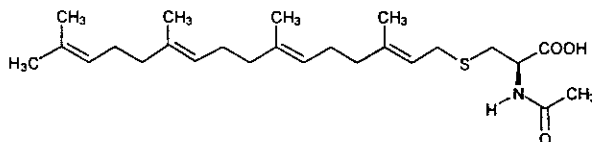
I-0105 5 mg

N-Acetyl-S-geranyl-L-cysteine (AGC); Used as a negative control for either AFC or AGGC; Molecular Formula: $\text{C}_{15}\text{H}_{25}\text{NO}_3\text{S}$; Molecular Weight: 299.4; Purity: > 98%



I-0119 5 mg

N-Acetyl-S-geranylgeranyl-L-cysteine (AGGC); Blocks receptor-mediated signaling in human neutrophils ($\text{IC}_{50} = 4 \mu\text{M}$). Molecular Formula: $\text{C}_{25}\text{H}_{41}\text{NO}_3\text{S}$; Molecular Weight: 435.7 ; Purity: > 98%



I-0123 5 mg

4-(3-Butoxy-4-methoxybenzyl)imidazolidin-2-one (RO-20-1724); Potent inhibitor of cAMP-specific phosphodiesterase ($\text{IC}_{50} = 2 \mu\text{M}$). Inhibits superoxide generation and arachidonic-induced platelet aggregation

I-0270 10 mg

AACOCF3 (Arachidonyltrifluoromethyl ketone); Potent and selective slow binding inhibitor of cytosolic (85 kDa) phospholipase A_2

I-0102 10 mg

Ascomycin; immunosuppressant

I-0150 1 mg

Bisindolylmaleimide I, HCl; Bisindolylmaleimide (GF 109203X) is a protein kinase inhibitor that is structurally similar to staurosporine but is highly selective for the inhibition of Protein Kinase C (PKC). GF 109203X inhibits PKC activity by acting as a competitive inhibitor of the ATP-binding site on PKC. Its reported K_i value is 10 nM.

I-0225 1 mg



1-800-701-3353

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