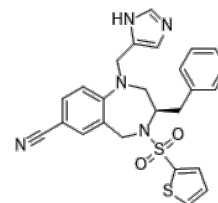


**Product Name** : BMS-214662  
**Cat. No.** : PC-24422  
**CAS No.** : 195987-41-8  
**Molecular Formula** : C<sub>25</sub>H<sub>23</sub>N<sub>5</sub>O<sub>2</sub>S<sub>2</sub>  
**Molecular Weight** : 489.61  
**Target** : Farnesyl transferase (FTase)  
**Solubility** : 10 mM in DMSO



CAS: 195987-41-8

## Biological Activity

BMS-214662 is a potent, selective **farnesyltransferase (FTase)** inhibitor with IC<sub>50</sub> of 1.35 nM, inhibits anchorage-independent growth of H-ras transformed NIH3T3 cells in soft agar EC<sub>50</sub> of 25 nM. BMS-214662 is over 1000-fold selective over the related enzyme GGT1. BMS-214662 has IC<sub>50</sub> values for inhibition of geranylgeranylation of Ras-CVLL and K-Ras of 1.3 and 2.3 μM, respectively. BMS-214662 reversed the H-Ras-transformed phenotype but not that of K-Ras or other oncogenes. BMS-214662 (300 mg/kg) achieved curative efficacy when given orally in the HCT-116 human colon tumor model. BMS-214662 demonstrated broad spectrum activity against human tumors, but murine tumors were not as sensitive.

## References

Rose WC, et al. **Cancer Res.** 2001 Oct 15;61(20):7507-17.  
 Hunt JT, et al. **J Med Chem.** 2000 Oct 5;43(20):3587-95.

**Caution: Product has not been fully validated for medical applications. Lab Use Only!**

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