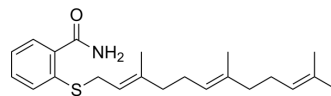


## Farnesyl thiosalicylic acid amide

Cat. No.:	HY-112666
CAS No.:	1092521-74-8
Molecular Formula:	C <sub>22</sub> H <sub>31</sub> NOS
Molecular Weight:	358
Target:	Ras
Pathway:	GPCR/G Protein; MAPK/ERK Pathway
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Farnesyl thiosalicylic acid amide (FTS-A) is an orally active derivative of farnesyl thiosalicylic acid (HY-14754). Farnesyl thiosalicylic acid amide reduces Ras-GTP levels and inhibits cell growth with IC <sub>50</sub> s of 20 and 10 μM for Panc-1 and U87 cells, respectively. Farnesyl thiosalicylic acid amide can be used for the research of cancer <sup>[1]</sup> .												
<b>In Vitro</b>	<p>Farnesyl thiosalicylic acid amide (50 μM, 24 h) decreases the levels of total Ras-GTP and K-Ras-GTP in Panc-1 and U87 cells<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Western Blot Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>U87 cells</td> </tr> <tr> <td>Concentration:</td> <td>50 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Reduced the level of N-Ras-GTP, phospho-ERK and phospho-AKT.</td> </tr> </table>	Cell Line:	U87 cells	Concentration:	50 μM	Incubation Time:	24 h	Result:	Reduced the level of N-Ras-GTP, phospho-ERK and phospho-AKT.				
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Concentration:	50 μM												
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Result:	Reduced the level of N-Ras-GTP, phospho-ERK and phospho-AKT.												
<b>In Vivo</b>	<p>Farnesyl thiosalicylic acid amide (100 mg/kg, i.g., once time) inhibits tumor growth in Panc-1 nude mouse model<sup>[1]</sup>.</p> <p>Farnesyl thiosalicylic acid amide (100 mg/kg, p.o., twice a day, 14 days) inhibits tumor growth in U87 nude mouse model<sup>[1]</sup>.</p> <p>Pharmacokinetic Analysis in Panc-1 Nude Mouse Model<sup>[1]</sup></p> <table border="1"> <thead> <tr> <th>Route</th> <th>Dose (mg/kg)</th> <th>half-life (min)</th> <th>T<sub>max</sub> (h)</th> <th>C<sub>max</sub> (ng/mL)</th> <th>AUC<sub>0-inf</sub> (min*ng/mL)</th> </tr> </thead> <tbody> <tr> <td>i.g.</td> <td>100</td> <td>314</td> <td>110</td> <td>371</td> <td>108783</td> </tr> </tbody> </table> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>	Route	Dose (mg/kg)	half-life (min)	T <sub>max</sub> (h)	C <sub>max</sub> (ng/mL)	AUC <sub>0-inf</sub> (min*ng/mL)	i.g.	100	314	110	371	108783
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i.g.	100	314	110	371	108783								

### REFERENCES

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[1]. Goldberg L, et al. New derivatives of farnesylthiosalicylic acid (salirasib) for cancer treatment: farnesylthiosalicylamide inhibits tumor growth in nude mice models. J Med Chem. 2009 Jan 8;52(1):197-205.

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**Caution: Product has not been fully validated for medical applications. For research use only.**

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