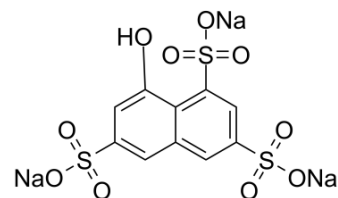


ζ-Stat trisodium

Cat. No.:	HY-123979A
CAS No.:	31894-34-5
Molecular Formula:	C ₁₀ H ₅ Na ₃ O ₁₀ S ₃
Molecular Weight:	450.31
Target:	PKC; Apoptosis
Pathway:	Epigenetics; TGF-beta/Smad; Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	ζ-Stat trisodium (NSC37044 trisodium) is a specific and atypical PKC-ζ inhibitor, with an IC ₅₀ of 5 μM. ζ-Stat trisodium can reduce melanoma cell lines proliferation and induce apoptosis, and has antitumor activity in vitro ^{[1][2]} .																
IC₅₀ & Target	aPKC-ζ 5 μM (IC ₅₀)																
In Vitro	<p>ζ-Stat (0.1-20 μM) shows only 13% inhibition on PKC-ι at 20 μM, but shows a significant inhibition on PKC-ζ as 51% at 5 μM level^[1].</p> <p>ζ-Stat (0.1-10 μM; 3 d) significantly decreases cell proliferation of SK-MEL-2 and MeWo upon increasing the concentrations^[1].</p> <p>ζ-Stat (7 or 10 μM; 24-72 h) and 5-FU in combination is able to decrease the viability of LoVo CRC cells by more than 75%^[2].</p> <p>ζ-Stat (5 μM; 3 d) shows a significant diminution of phosphorylated, total PKC-ζ, Bcl-2 and PARP levels, and increases Caspase-3 and cleaved-PARP levels in SK-MEL-2 and MeWo cells^[1].</p> <p>ζ-Stat (5 μM; 1-10 h) does not show significant cytotoxicity on MEL-F-NEO, SK-MEL-2 and MeWo cells^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Proliferation Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>MEL-F-NEO, SK-MEL-2 and MeWo cells</td> </tr> <tr> <td>Concentration:</td> <td>0.1, 0.5, 1, 2.5, 5, 7.5, 10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>3 days</td> </tr> <tr> <td>Result:</td> <td>Decreased proliferation by 47.7% for 5 μM in SK-MEL-2 cells and by 50.6% for 5 μM in MeWo cells. Showed significant inhibitions on MEL-F-NEO cells 19.3% (P ≤ 0.05) at 10 μM.</td> </tr> </table> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>SK-MEL-2 and MeWo cells</td> </tr> <tr> <td>Concentration:</td> <td>5 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>3 days</td> </tr> <tr> <td>Result:</td> <td>Decreased phosphorylated and total PKC-ζ levels.</td> </tr> </table>	Cell Line:	MEL-F-NEO, SK-MEL-2 and MeWo cells	Concentration:	0.1, 0.5, 1, 2.5, 5, 7.5, 10 μM	Incubation Time:	3 days	Result:	Decreased proliferation by 47.7% for 5 μM in SK-MEL-2 cells and by 50.6% for 5 μM in MeWo cells. Showed significant inhibitions on MEL-F-NEO cells 19.3% (P ≤ 0.05) at 10 μM.	Cell Line:	SK-MEL-2 and MeWo cells	Concentration:	5 μM	Incubation Time:	3 days	Result:	Decreased phosphorylated and total PKC-ζ levels.
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REFERENCES

- [1]. Ratnayake WS, et, al. Oncogenic PKC- α activates Vimentin during epithelial-mesenchymal transition in melanoma; a study based on PKC- α and PKC- ζ specific inhibitors. Cell Adh Migr. 2018; 12(5):447-463.
- [2]. Islam SMA, et, al. Atypical Protein Kinase-C inhibitors exhibit a synergistic effect in facilitating DNA damaging effect of 5-fluorouracil in colorectal cancer cells. Biomed Pharmacother. 2020 Jan; 121:109665.
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Caution: Product has not been fully validated for medical applications. For research use only.

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