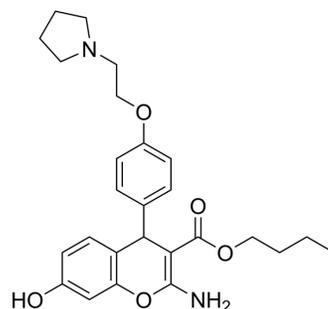


## Estrogen receptor $\alpha$ antagonist 1

Cat. No.:	HY-150692
CAS No.:	2580941-45-1
Molecular Formula:	C <sub>26</sub> H <sub>32</sub> N <sub>2</sub> O <sub>5</sub>
Molecular Weight:	452.54
Target:	Estrogen Receptor/ERR
Pathway:	Others
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Estrogen receptor $\alpha$ antagonist 1 (compound 35) is a highly selective antagonist of estrogen receptor $\alpha$ , with IC <sub>50</sub> s of 0.02, 6.55 and 7.73 $\mu$ M for estrogen receptor $\alpha$ , estrogen receptor $\beta$ and MCF-7 cells, respectively. Estrogen receptor $\alpha$ antagonist 1 can be used for the research of cancer <sup>[1]</sup> .								
<b>IC<sub>50</sub> &amp; Target</b>	IC <sub>50</sub> : 0.02 $\mu$ M (estrogen receptor $\alpha$ ), 6.55 $\mu$ M (estrogen receptor $\beta$ ), 7.73 $\mu$ M (MCF-7 cells) <sup>[1]</sup>								
<b>In Vitro</b>	<p>Estrogen receptor <math>\alpha</math> antagonist 1 (compound 35) (1 nM-100 <math>\mu</math>M; 72 h) selectively inhibits estrogen receptor <math>\alpha</math>, estrogen receptor <math>\beta</math> and MCF-7 cells<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>MCF-7 cells</td> </tr> <tr> <td>Concentration:</td> <td>1 nM-100 <math>\mu</math>M</td> </tr> <tr> <td>Incubation Time:</td> <td>72 hours</td> </tr> <tr> <td>Result:</td> <td>Exhibited IC<sub>50</sub>s of 0.02, 6.55 and 7.73 <math>\mu</math>M for estrogen receptor <math>\alpha</math>, estrogen receptor <math>\beta</math> and MCF-7 cells, respectively at 72 hours.</td> </tr> </table>	Cell Line:	MCF-7 cells	Concentration:	1 nM-100 $\mu$ M	Incubation Time:	72 hours	Result:	Exhibited IC <sub>50</sub> s of 0.02, 6.55 and 7.73 $\mu$ M for estrogen receptor $\alpha$ , estrogen receptor $\beta$ and MCF-7 cells, respectively at 72 hours.
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### REFERENCES

[1]. Carr M, et al. Optimisation of estrogen receptor subtype-selectivity of a 4-Aryl-4H-chromene scaffold previously identified by virtual screening. *Bioorg Med Chem.* 2020 Mar 1;28(5):115261.

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

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