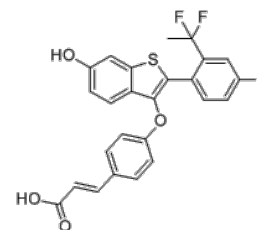


**Product Name** : LSZ102  
**Cat. No.** : PC-63293  
**CAS No.** : 2135600-76-7  
**Molecular Formula** : C<sub>25</sub>H<sub>17</sub>F<sub>3</sub>O<sub>4</sub>S  
**Molecular Weight** : 470.462  
**Target** : Estrogen Receptor/ERR  
**Solubility** : 10 mM in DMSO



## Biological Activity

LSZ102 is a potent, orally bioavailable, selective estrogen receptor degrader (**SERD**) with ER $\alpha$  transcription IC<sub>50</sub> of 6 nM and ER $\alpha$  degradation IC<sub>50</sub> of 0.2 nM.

LSZ102 robustly causes inhibition of cell proliferation in MCF-7 cells with IC<sub>50</sub> of 1.7 nM, inhibits the mRNA level of canonical endogenous ER target gene GREB1 with IC<sub>50</sub> of 8.9 nM.

LSZ102 demonstrates anti-tumor efficacy in the ER<sup>+</sup> human breast cancer MCF-7 xenograft model (10 mg/kg).

## References

Tria GS, et al. *J Med Chem.* 2018 Apr 12;61(7):2837-2864.

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

E-mail: tech@probechem.com