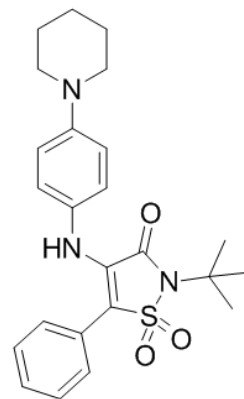


**Product Name** : AZ876  
**Cat. No.** : PC-45444  
**CAS No.** : 898800-26-5  
**Molecular Formula** : C<sub>24</sub>H<sub>29</sub>N<sub>3</sub>O<sub>3</sub>S  
**Molecular Weight** : 439.5704  
**Target** : Liver X Receptor (LXR)  
**Solubility** : DMSO: ≥ 2.6 mg/mL



### Biological Activity

AZ876 (AZ 876) is a potent, highly selective **LXR agonist** with Ki/EC<sub>50</sub> of 7/6 nM and 11/73 nM for hLXR $\alpha$  and hLXR $\beta$  respectively.

AZ876 is a more potent binder and activator of LXR $\alpha$  and LXR $\beta$  than GW3965.

AZ876 is highly selective against FXR, RXR, TR $\alpha$ , TR $\beta$  etc.

AZ876 reduces lesion area, and strongly decreases lesion area, lesion number and severity in vivo.

AZ876 protects against pathological cardiac hypertrophy and fibrosis without lipogenic side effects in mice.

### References

van der Hoorn J, et al. *Br J Pharmacol.* 2011 Apr;162(7):1553-63.

Cannon MV, et al. *Eur J Heart Fail.* 2015 Mar;17(3):273-82.

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

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