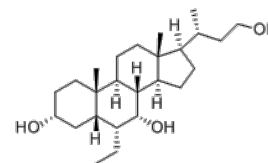


**Product Name** : BAR502  
**Cat. No.** : PC-20485  
**CAS No.** : 1612191-86-2  
**Molecular Formula** : C<sub>25</sub>H<sub>44</sub>O<sub>3</sub>  
**Molecular Weight** : 392.62  
**Target** : Farnesoid X Receptor (FXR)  
**Solubility** : 10 mM in DMSO



CAS: 1612191-86-2

## Biological Activity

BAR502 (NorEcdcoH) is a potent, selective, dual farnesoid X receptor (FXR) and G-protein coupled bile acid receptor 1 (GP-BAR1) ligand (agonist) with EC<sub>50</sub> of 2.0 and 0.4 uM, respectively.

BAR502 potently inhibits binding of LIF to LIFR with an IC<sub>50</sub> of 3.8 uM.

BAR502 protected against liver damage caused by HFD by promoting the browning of adipose tissue.

BAR502 partially protected against liver damage caused by Western diet.

The combination of BAR502 and UDCA reversed the pro-atherogenic lipid profile and completely reversed the histopathology damage, attenuating liver steatosis, ballooning, inflammation and fibrosis.

## References

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Carino A, et al. Sci Rep. 2017 Feb 16;7:42801.

Festa C. et al. J Med Chem 57, 8477–95 (2014).

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

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