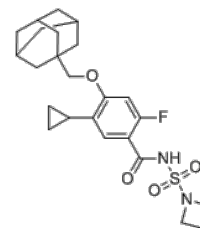


**Product Name** : GDC-0276  
**Cat. No.** : PC-72192  
**CAS No.** : 1494581-70-2  
**Molecular Formula** : C<sub>24</sub>H<sub>31</sub>FN<sub>2</sub>O<sub>4</sub>S  
**Molecular Weight** : 462.58  
**Target** : Sodium Channel  
**Solubility** : 10 mM in DMSO (4.6 mg/mL)



## Biological Activity

GDC-0276 (GDC 0276) is a highly potent, selective **NaV1.7** inhibitor with IC<sub>50</sub> of with binding K<sub>i</sub> of 1.1 nM (hNaV1.7), cellular sodium influx IC<sub>50</sub> of 49 nM.

GDC-0276 displays 18-fold selectivity over NaV1.5 (cellular sodium influx IC<sub>50</sub>=870 nM), as well as 26-1200-fold selectivity over NaV1.1/1.2/1.6 (potency measured by voltage clamp).

GDC-0276 showed a concentration dependent reduction in nociceptive events, with a EC<sub>50</sub> of 1.7 μM in inherited erythromelalgia (IEM) mouse model.

## References

Brian S Safina, et al. *J Med Chem.* 2021 Mar 25;64(6):2953-2966.

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

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