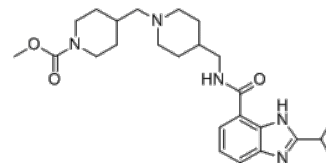


**Product Name** : Felcisetrag  
**Cat. No.** : PC-73307  
**CAS No.** : 916075-84-8  
**Molecular Formula** : C<sub>25</sub>H<sub>37</sub>N<sub>5</sub>O<sub>3</sub>  
**Molecular Weight** : 455.603  
**Target** : 5-HT Receptor  
**Solubility** : 10 mM in DMSO



## Biological Activity

Felcisetrag (TAK-954, TD-8954) is a highly selective and potent **5-HT<sub>4</sub>** receptor agonist with pK<sub>i</sub> of 9.4 (human 5-HT<sub>4</sub>). Felcisetrag (TAK-954, TD-8954) displays high selectivity (>2,000-fold) over all other 5-HT receptors and non-5-HT receptors, ion channels, enzymes and transporters.

Felcisetrag (TAK-954, TD-8954) produced an elevation of cAMP in HEK-293 cells expressing the h5-HT<sub>4</sub>(c) receptor (pEC=9.3), and contracted the guinea pig colonic longitudinal muscle/myenteric plexus preparation (pEC<sub>50</sub>=8.6).

Felcisetrag (TAK-954, TD-8954) (0.03-3 mg/kg) increased the colonic transit of carmine red dye, reducing the time taken for its excretion.

Felcisetrag (TAK-954, TD-8954) demonstrated robust in vivo stimulatory activity in the gastrointestinal (GI) tract of guinea pigs, rats, dogs, and humans.

## References

Shen F, et al. *Neuropharmacology*. 2011 Jul-Aug;61(1-2):69-79.

Beattie DT, et al. *Front Pharmacol*. 2011 May 30;2:25.

Beattie DT, et al. *Vascul Pharmacol*. 2013 Jan;58(1-2):150-6.

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

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