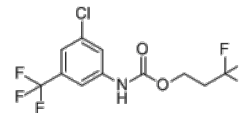


**Product Name** : ETG-5773  
**Cat. No.** : PC-49269  
**CAS No.** : 23794-80-1  
**Molecular Formula** : C<sub>11</sub>H<sub>8</sub>ClF<sub>6</sub>NO<sub>2</sub>  
**Molecular Weight** : 335.630  
**Target** :  
**Solubility** : 10 mM in DMSO



### Biological Activity

ETG-5773 (ETG5773) is a selective, cross-species active, non-competitive, non-substrate-like inhibitor of **NaCT (SLC13A5, INDY)**, reduces citrate uptake in cell lines expressing mouse or human NaCT with IC<sub>50</sub> of 180 and 160 nM, respectively. ETG-5773 displays no activity on the closely related human NaDC3 (SLC13A3) transporter at 20 μM. ETG-5773 (15 mg/kg twice-daily, oral) ameliorates diet-induced obesity in DIO mice. ETG-5773 attenuates hepatic steatosis and regulates lipogenesis-related genes, significantly downregulates Acly mRNA levels. ETG-5773 is able to reduce liver fat and other features of metabolically associated fatty liver disease (MAFLD).

### References

Grit Zahn, et al. *Metabolites*. 2022 Aug 8;12(8):732.

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

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