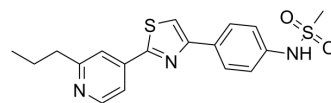


## FGH10019

|                    |  |       |         |
|--------------------|--|-------|---------|
| Cat. No.:          | HY-16207   |       |         |
| CAS No.:           | 1046045-61-7   |       |         |
| Molecular Formula: | C <sub>18</sub> H <sub>19</sub> N <sub>3</sub> O <sub>2</sub> S <sub>2</sub> |       |         |
| Molecular Weight:  | 373.49   |       |         |
| Target:            | Fatty Acid Synthase (FASN)   |       |         |
| Pathway:           | Metabolic Enzyme/Protease  |       |         |
| Storage:           | Powder   | -20°C | 3 years |
|                    |  | 4°C   | 2 years |
|                    | In solvent   | -80°C | 2 years |
|                    |  | -20°C | 1 year  |



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : ≥ 38 mg/mL (101.74 mM)  
 \* "≥" means soluble, but saturation unknown.

| Preparing Stock Solutions | Solvent       | Mass | 1 mg      | 5 mg       | 10 mg      |
|---------------------------|---------------|------|-----------|------------|------------|
|                           | Concentration |      |           |            |            |
|                           | 1 mM          |      | 2.6774 mL | 13.3872 mL | 26.7745 mL |
|                           | 5 mM          |      | 0.5355 mL | 2.6774 mL  | 5.3549 mL  |
|                           | 10 mM         |      | 0.2677 mL | 1.3387 mL  | 2.6774 mL  |

Please refer to the solubility information to select the appropriate solvent.

### BIOLOGICAL ACTIVITY

|                                     |  |
|-------------------------------------|--|
| <b>Description</b>                  | FGH10019 is a novel sterol regulatory element-binding protein (SREBP) inhibitor with IC <sub>50</sub> of 1 μM.   |
| <b>IC<sub>50</sub> &amp; Target</b> | IC <sub>50</sub> : 1 μM (SREBP)  |
| <b>In Vitro</b>                     | Treatment of the CHO-K1 cells with analog FGH10019 decreases the percentage of the mature form of SREBP-2 (68 kDa) at lower concentrations than treatment with fatostatin. Densitometric analysis of the gels indicates that the IC <sub>50</sub> of analog FGH10019 is approximately 1 μM, which is 5-10 times lower than the IC <sub>50</sub> of fatostatin (appr 10 μM) <sup>[1]</sup> .<br>MCE has not independently confirmed the accuracy of these methods. They are for reference only. |
| <b>In Vivo</b>                      | FGH10019-treated chow is fed at a dose rate calculated to provide about 0.7 mg analog FGH10019 per day, at about 23 mg/kg body weight, to 5-wk-old male ob/ob mice weighing an average of appr 30 g. After 8 wk on the analog 24-treated chow, the mice gain 8-9% less weight than control mice <sup>[1]</sup> .<br>MCE has not independently confirmed the accuracy of these methods. They are for reference only.  |

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## PROTOCOL

### Animal Administration <sup>[1]</sup>

Five-week-old homozygous male obese (ob/ob) mice (C57BL/6J) are housed five per cage, and had ad libitum access to normal chow and water for 1 wk after their arrival. On day 1 of the experiment, the animals (10 per group) are fed normal chow (control diet) or chow that contains 200 mg/kg of analogue 24. These doses are estimated to provide approximately 0.7 mg analogue 24 per day (appr 23 mg/kg body weight per day). Daily food intake and body weight are carefully monitored and recorded between 3:00 and 5:00 p.m. Serum constituents, and TG levels in livers are determined.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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## CUSTOMER VALIDATION

- Sci Rep. 2017 May 23;7(1):2303.

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## REFERENCES

[1]. Kamisuki S, et al. Synthesis and evaluation of diarylthiazole derivatives that inhibit activation of sterol regulatory element-binding proteins. J Med Chem. 2011 Jul 14;54(13):4923-7.

**Caution: Product has not been fully validated for medical applications. For research use only.**

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