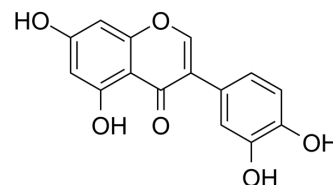


## Orobol

<b>Cat. No.:</b>	HY-N3127
<b>CAS No.:</b>	480-23-9
<b>Molecular Formula:</b>	C <sub>15</sub> H <sub>10</sub> O <sub>6</sub>
<b>Molecular Weight:</b>	286.24
<b>Target:</b>	Casein Kinase; PI3K
<b>Pathway:</b>	Cell Cycle/DNA Damage; Stem Cell/Wnt; PI3K/Akt/mTOR
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Orobol is one of the major soy isoflavones and has various pharmacological activities, including anti-skin-aging and anti-obesity effects. Orobol inhibits CK1ε, VEGFR2, MAP4K5, MNK1, MUSK, TOPK, and TNIK (IC <sub>50</sub> =1.24-4.45 μM). Orobol also inhibits PI3K isoforms (IC <sub>50</sub> =3.46-5.27 μM for PI3K α/β/γ/κ/δ) <sup>[1][2]</sup> .
<b>In Vitro</b>	Orobol binds to CK1ε in an ATP-competitive manner and exerts anti-obesity effects by targeting casein kinase 1 epsilon <sup>[2]</sup> . Orobol (5-20 μM) effectively suppresses MDI (isobutylmethylxanthine, dexamethasone and insulin (MDI))-induced phosphorylation of 4E-BP1 <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
<b>In Vivo</b>	Orobol attenuates high fat diet-induced weight gain and lipid accumulation without affecting food intake in C57BL/6J mice <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
<b>Animal Model:</b>	HFD-induced obesity in C57BL/6J mice <sup>[2]</sup>
<b>Dosage:</b>	10 mg/kg
<b>Administration:</b>	Intragastrically; daily for 23 weeks
<b>Result:</b>	Significantly reduced body weight by 17.3% compared to the HFD group.

### REFERENCES

[1]. Kim MH, et al. Lipid Nanoparticles for Enhancing the Physicochemical Stability and Topical Skin Delivery of Orobol. *Pharmaceutics*. 2020;12(9):845. Published 2020 Sep 3.

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

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