

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

linkedin.com/company/szaboscandic in



PRODUCT INFORMATION



Sobetirome

Item No. 30814

CAS Registry No.: 211110-63-3

Formal Name: 2-[4-[[4-hydroxy-3-(1-methylethyl]

phenyl]methyl]-3,5-dimethylphenoxy]-

acetic acid

Synonyms: GC-1, QRX 431

MF: $C_{20}H_{24}O_4$ FW: 328.4 **Purity:** ≥98% Supplied as: A solid Storage: -20°C Stability: ≥2 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

Sobetirome is supplied as a solid. A stock solution may be made by dissolving the sobetirome in the solvent of choice, which should be purged with an inert gas. Sobetirome is soluble in ethanol and DMSO. The solubility of sobetirome in these solvents is approximately 45 and 30 mg/ml, respectively.

Description

Sobetirome is an agonist of thyroid hormone receptor β (TR β ; EC₅₀ = 7 nM in a reporter assay). It selectively binds to TR β over TR α with K_d values of 0.1 and 1.8 nM, respectively. Sobetirome (97 nmol/kg, i.p.) increases hepatic levels of the HDL receptor SR-B1 and serum levels of C4, a marker of increased bile acid synthesis, in mice fed normal chow or high-cholesterol diets.² It reduces serum cholesterol and triglyceride levels in these same models. Sobetirome (154 nmol/kg, i.p.) also decreases plasma cholesterol levels, without increasing heart rate, in a mouse model of hypothyroidism induced by an iodine-deficient diet containing the thyroid hormone biosynthesis inhibitor 5-propyl-2-thio-uracil.3

References

- 1. Nguyen, N.-H., Apriletti, J.W., Cunha Lima, S.T., et al. Rational design and synthesis of a novel thyroid hormone antagonist that blocks coactivator recruitment. J. Med. Chem. 45(15), 3310-3320 (2002).
- 2. Johansson, L., Rudling, M., Scanlan, T.S., et al. Selective thyroid receptor modulation by GC-1 reduces serum lipids and stimulates steps of reverse cholesterol transport in euthyroid mice. Proc. Natl. Acad. Sci. USA 102(29), 10297-10302 (2005).
- 3. Trost, S.U., Swanson, E., Gloss, B., et al. The thyroid hormone receptor-β-selective agonist GC-1 differentially affects plasma lipids and cardiac activity. Endocrinology 141(9), 3057-3064 (2000).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

WARRANTY AND LIMITATION OF REMEDY

subject to Cayman's Terms and Conditions. Complete Terms and Conditions including Warranty and Limitation of Liability information Buyer agrees to purchase the material can be found on our website.

Copyright Cayman Chemical Company, 09/09/2020

CAYMAN CHEMICAL

1180 EAST ELLSWORTH RD ANN ARBOR, MI 48108 · USA PHONE: [800] 364-9897

[734] 971-3335

FAX: [734] 971-3640 CUSTSERV@CAYMANCHEM.COM WWW.**CAYMANCHEM**.COM