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- Trockeneiszuschlag
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- Expressversand

SZABO-SCANDIC Handels GmbH

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](https://www.linkedin.com/company/szaboscandic) 

PRODUCT INFORMATION



Fructose-1,6-bisphosphatase-1 Inhibitor

Item No. 18860

CAS Registry No.: 883973-99-7

Formal Name: 2,5-dichloro-N-(5-chloro-2-benzoxazolyl)-benzenesulfonamide

Synonyms: F1,6BPase-1 Inhibitor, FBP1 Inhibitor, FBPase-1 Inhibitor

MF: $C_{13}H_7Cl_3N_2O_3S$

FW: 377.6

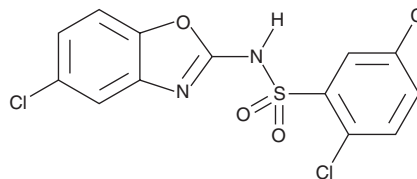
Purity: $\geq 98\%$

UV/Vis.: λ_{max} : 204, 287 nm

Supplied as: A crystalline solid

Storage: -20°C

Stability: As supplied, 2 years from the QC date provided on the Certificate of Analysis, when stored properly



Laboratory Procedures

Fructose-1,6-bisphosphatase-1 (FBPase-1) inhibitor is supplied as a crystalline solid. A stock solution may be made by dissolving the FBPase-1 inhibitor in the solvent of choice. FBPase-1 inhibitor is soluble in organic solvents such as DMSO and dimethyl formamide (DMF), which should be purged with an inert gas. The solubility of FBPase-1 inhibitor in these solvents is approximately 14 and 20 mg/ml, respectively.

FBPase-1 inhibitor is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, FBPase-1 inhibitor should first be dissolved in DMF and then diluted with the aqueous buffer of choice. FBPase-1 inhibitor has a solubility of approximately 0.14 mg/ml in a 1:6 solution of DMF:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Description

FBPase-1 is an enzyme that catalyzes the conversion of fructose-1,6-bisphosphate to fructose-6-phosphate, which is one of the rate-limiting steps in gluconeogenesis. Excess hepatic FBPase-1 activity contributes to hyperglycemia in patients. Thus, the development of specific FBPase-1 inhibitors is of great clinical interest for the treatment of patients with type 2 diabetes. FBPase-1 inhibitor is a cell-permeable benzoxazolo-sulfonamide compound that blocks human FBPase-1 enzymatic activity with an IC_{50} value of $3.4 \mu\text{M}$ ($K_i = 1.1 \mu\text{M}$) by competing at the AMP allosteric binding site.^{1,2} It has been shown to block glucose production in rat hepatoma cells that are starved of nutrients with an IC_{50} value of $6.6 \mu\text{M}$.^{1,2}

References

1. Lai, C., Gum, R.J., Daly, M., *et al.* Benzoxazole benzenesulfonamides as allosteric inhibitors of fructose-1,6-bisphosphatase. *Bioorg. Med. Chem. Lett.* **16**(7), 1807-1810 (2006).
2. von Geldern, T.W., Lai, C., Gum, R.J., *et al.* Benzoxazole benzenesulfonamides are novel allosteric inhibitors of fructose-1,6-bisphosphatase with a distinct binding mode. *Bioorg. Med. Chem. Lett.* **16**(7), 1811-1815 (2006).

WARNING

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

SAFETY DATA

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.

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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