

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



SU 5402

Purity:

Item No. 13182

CAS Registry No.: 215543-92-3

Formal Name: 2-[(1,2-dihydro-2-oxo-3H-indol-3-ylidene)

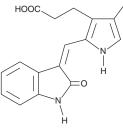
methyl]-4-methyl-1H-pyrrole-3-propanoic acid

MF: $C_{17}H_{16}N_2O_3$ FW: 296.3

≥95% λ_{max} : 210, 275, 430 nm A crystalline solid UV/Vis.: Supplied as:

Storage: -20°C Stability: ≥2 years

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



Laboratory Procedures

SU 5402 is supplied as a crystalline solid. A stock solution may be made by dissolving the SU 5402 in the solvent of choice, which should be purged with an inert gas. SU 5402 is soluble in organic solvents such as DMSO and dimethyl formamide (DMF). The solubility of SU 5402 in these solvents is approximately 30 and 5 mg/ml, respectively.

SU 5402 is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, SU 5402 should first be dissolved in DMF and then diluted with the aqueous buffer of choice. SU 5402 has a solubility of approximately 0.5 mg/ml in a 1:1 solution of DMF:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Description

SU 5402 is an inhibitor of the tyrosine kinase domains of VEGFR2, FGFR1, and PDGFRB $(IC_{50}s = 0.02, 0.03, and 0.51 \mu M, respectively)$. It is much less effective against other receptor tyrosine kinases.¹ SU 5402 is commonly used to evaluate the role of FGFR1 in cellular functions.²⁻⁵

References

- 1. Sun, L., Tran, N., Liang, C., et al. Design, synthesis, and evaluations of substituted 3-[(3- or 4-carboxyethylpyrrol-2-yl)methylidenyl]indolin-2-ones as inhibitors of VEGF, FGF, and PDGF receptor tyrosine kinases. J. Med. Chem. 42(25), 5120-5130 (1999).
- 2. Heryanto, B., Lipson, K. E., and Rogers, P. A. W. Effect of angiogenesis inhibitors on oestrogen-mediated endometrial endothelial cell proliferation in the ovariectomized mouse. Reproduction 125, 337-346 (2003).
- 3. Ying, Q. L., Wray, J., Nichols, J., et al. The ground state of embryonic stem cell self-renewal. Nature 453, 519-524 (2008).
- 4. Hasse, C., Holz, O., Lange, E., et al. FGFR-ERK signaling is an essential component of tissue separation. Dev. Biol. 395(1), 154-166 (2014).
- 5. Kachel, P., Trojanowicz, B., Sekulla, C., et al. Phosphorylation of pyruvate kinase M2 and lactate dehydrogenase A by fibroblast growth factor receptor 1 in benign and malignant thyroid tissue. BMC Cancer 15:140, (2015).

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, 09/20/2019

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