# PRODUCT INFORMATION



# **Abrocitinib**

Item No. 34869

CAS Registry No.: 1622902-68-4

Formal Name: N-[cis-3-(methyl-7H-pyrrolo[2,3-d]

pyrimidin-4-ylamino)cyclobutyl]-1-

propanesulfonamide

Synonym: PF-04965842 MF:  $C_{14}H_{21}N_5O_2S$ FW:

323.4 Purity: ≥98%  $\lambda_{max}$ : 288 nm UV/Vis.: Supplied as: A solid -20°C Storage: Stability: ≥4 years

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

# **Laboratory Procedures**

Abrocitinib is supplied as a solid. A stock solution may be made by dissolving the abrocitinib in the solvent of choice, which should be purged with an inert gas. Abrocitinib is soluble in DMSO.

## Description

Abrocitinib is a JAK1 inhibitor (IC $_{50}$  = 0.029  $\mu$ M).<sup>1</sup> It is selective for JAK1 over JAK2, JAK3, and TYK2 (IC<sub>50</sub>s = 0.803, >10, and 1.253  $\mu$ M, respectively) and a panel of 40 kinases at 1  $\mu$ M. Abrocitinib inhibits  $\widetilde{IFN}$ - $\alpha$ -, IL-21-, or IL-27-induced phosphorylation of STAT3 (IC<sub>50</sub>s = 0.189, 0.511, and 0.271  $\mu$ M, respectively) and IFN- $\gamma$ -induced phosphorylation of STAT1 (IC<sub>50</sub> = 0.163  $\mu$ M) in isolated human whole blood. It reduces paw edema in a rat model of adjuvant-induced arthritis when administered at doses of 1, 5, and 15 mg/kg. Formulations containing abrocitinib have been used in the treatment of atopic dermatitis.

## Reference

1. Vazquez, M.L., Kaila, N., Strohbach, J.W., et al. Identification of N-{cis-3-[methyl(7H-pyrrolo[2,3-d] pyrimidin-4-yl)amino]cyclobutyl}propane-1-sulfonamide (PF-04965842): A selective JAK1 clinical candidate for the treatment of autoimmune diseases. J. Med. Chem. 61(3), 1130-1152 (2018).

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.

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