PRODUCT INFORMATION



MRS2578

Item No. 19704

CAS Registry No.: 711019-86-2

N,N"-1,4-butanediylbis[N'-(3-Formal Name:

isothiocyanatophenyl)-thiourea

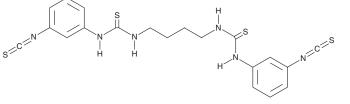
MF: $C_{20}H_{20}N_6S_4$

FW: 472.7 **Purity:**

 λ_{max} : 217, 273 nm A crystalline solid UV/Vis.: Supplied as:

-20°C Storage: Stability: ≥4 years

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



Laboratory Procedures

MRS2578 is supplied as a crystalline solid. A stock solution may be made by dissolving the MRS2578 in the solvent of choice, which should be purged with an inert gas. MRS2578 is soluble in organic solvents such as DMSO and dimethyl formamide (DMF). The solubility of MRS2578 in these solvents is approximately 10 and 25 mg/ml, respectively.

MRS2578 is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, MRS2578 should first be dissolved in DMF and then diluted with the aqueous buffer of choice. MRS2578 has a solubility of approximately 0.2 mg/ml in a 1:4 solution of DMF:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Description

MRS2578 is an antagonist of the purinergic P2Y₆ receptor (IC_{50} s = 37 and 98 nM for human and rat receptors, respectively). It is without effect at other purinergic receptors. MRS2578 is used to study the cellular and physiological roles of P2Y₆.²⁻⁵

References

- 1. Mamedova, L.K., Joshi, B.V., Gao, Z.-G., et al. Diisothiocyanate derivatives as potent, insurmountable antagonists of P2Y₆ nucleotide receptors. Biochem. Pharmacol. 67(9), 1763-1770 (2004).
- Koizumi, S., Shigemoto-Mogami, Y., Nasu-Tada, K., et al. UDP acting at P2Y, receptors is a mediator of microglial phagocytosis. Nature 446(78139), 1091-1095 (2007).
- Kukulski, F., Ben Yebdri, F., Lefebvre, J., et al. Extracellular nucleotides mediate LPS-induced neutrophil migration in vitro and in vivo. J. Leukoc. Biol. 81(5), 1269-1275 (2007).
- Ma, X., Pan, X., Wei, Y., et al. Chemotherapy-induced uridine diphosphate release promotes breast cancer metastasis through P2Y₆ activation. Oncotarget 7(20), 29036-29050 (2016).
- Sil, P., Hayes, C.P., Reaves, B.J., et al. P2Y₆ receptor antagonist MRS2578 inhibits neutrophil activation and aggregated neutrophil extracellular trap formation induced by gout-associated monosodium urate crystals. J. Immunol. 198(1), 428-442 (2017).

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

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