

## Produktinformation



Forschungsprodukte & Biochemikalien
Zellkultur & Verbrauchsmaterial
Diagnostik & molekulare Diagnostik
Laborgeräte & Service

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# **PRODUCT** INFORMATION



**MRS2578** 

Item No. 19704

Formal Name:N,N"-1,4-butanediylbis[N'-(3-isothiocyanatophenyl)-thioureaMF: $C_{20}H_{20}N_6S_4$ FW:472.7Purity: $\geq 98\%$ UV/Vis.: $\lambda_{max}$ : 217, 273 nm
MF: $C_{20}H_{20}N_6S_4$ FW:       472.7         Purity: $\geq 98\%$ UV/Vis.: $\lambda_{max}$ : 217, 273 nm
FW: $472.7$ $s = c = N$ N       N       N $r = c = N$ Purity: $\geq 98\%$ $V = V = 10^{-10}$ $V = 10^{-10}$ $V = 10^{-10}$ $V = 10^{-10}$ $V = 10^{-10}$ UV/Vis.: $\lambda_{max}$ : 217, 273 nm $V = 10^{-10}$ $V = 10^{-10}$ $V = 10^{-10}$ $V = 10^{-10}$
FW:       472.7 $s = c^{-r}$ $H$
<b>UV/Vis.:</b> $\lambda_{max}$ : 217, 273 nm
UV/Vis.: $\lambda_{\text{max}}$ : 217, 273 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

MRS2578 is supplied as a crystalline solid. A stock solution may be made by dissolving the MRS2578 in the solvent of choice. MRS2578 is soluble in organic solvents such as DMSO and dimethyl formamide (DMF), which should be purged with an inert gas. 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  $P2Y_6$  receptor (IC<sub>50</sub>s = 37 and 98 nM for human and rat receptors, respectively).<sup>1</sup> It is without effect at other purinergic receptors. MRS2578 is used to study the cellular and physiological roles of P2Y<sub>6</sub>.<sup>2-5</sup>

#### References

- 1. Mamedova, L.K., Joshi, B.V., Gao, Z.-G., et al. Diisothiocyanate derivatives as potent, insurmountable antagonists of P2Y<sub>6</sub> nucleotide receptors. Biochem. Pharmacol. 67(9), 1763-1770 (2004).
- 2. Koizumi, S., Shigemoto-Mogami, Y., Nasu-Tada, K., et al. UDP acting at P2Y<sub>6</sub> receptors is a mediator of microglial phagocytosis. Nature 446(78139), 1091-1095 (2007).
- 3. 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 4. metastasis through P2Y<sub>6</sub> activation. Oncotarget 7(20), 29036-29050 (2016).
- 5. Sil, P., Hayes, C.P., Reaves, B.J., et al. P2Y<sub>6</sub> 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.

#### 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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