# PRODUCT INFORMATION



## **AA43279**

Item No. 40428

CAS Registry No.: 354812-16-1

Formal Name: 3-amino-5-(4-methoxyphenyl)-2-

thiophenecarboxamide

MF:  $C_{12}H_{12}N_2O_2S$ 

FW: 248.3 **Purity:** ≥98% Supplied as: A solid Storage: -20°C Stability: ≥4 years

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

#### **Laboratory Procedures**

AA43279 is supplied as a solid. A stock solution may be made by dissolving the AA43279 in the solvent of choice, which should be purged with an inert gas. AA43279 is soluble in DMSO and acetonitrile.

### Description

AA43279 is an activator of voltage-gated sodium channel 1.1 (Na $_{v}$ 1.1; EC $_{50}$  = 9.5  $\mu$ M in a patch-clamp assay using HEK293 cells expressing the human channel). It is selective for Na, 1.1 over Na, 1.4 and Na, 1.7 at 30  $\mu$ M but also activates Na<sub>v</sub>1.2, Na<sub>v</sub>1.5, and Na<sub>v</sub>1.6 (EC<sub>50</sub>s = 22.8, 14.4, and 11.6  $\mu$ M, respectively). AA43279 also inhibits IkB kinase  $\beta$  (IKK $\beta$ ; IC<sub>50</sub> = 2.9  $\mu$ M). It increases the seizure threshold in the maximal electroshock seizure threshold (MEST) test in mice when administered at a dose of 300 mg/kg.<sup>1</sup>

#### References

- 1. Frederiksen, K., Lu, D., Yang, J., et al. A small molecule activator of Na., 1.1 channels increases fast-spiking interneuron excitability and GABAergic transmission in vitro and has anti-convulsive effects in vivo. Eur. J. Neurosci. 46(3), 1887-1896 (2017).
- 2. Morwick, T., Berry, A., Brickwood, J., et al. Evolution of the thienopyridine class of inhibitors of IKB kinase-β: part I: Hit-to-lead strategies. J. Med. Chem. 49(10), 2898-2908 (2006).

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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