# **PRODUCT** INFORMATION



NSC 405020

Item No. 14859

CAS Registry No.:	7497-07-6	
Formal Name:	3,4-dichloro-N-(1-methylbutyl)-benzamide	0
MF:	C <sub>12</sub> H <sub>15</sub> Cl <sub>2</sub> NO	
FW:	260.2	
Purity:	≥98%	
UV/Vis.:	λ <sub>max</sub> : 238 nm	CI
Supplied as:	A crystalline solid	
Storage:	-20°C	ĊI
Stability:	≥4 years	

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

## Laboratory Procedures

NSC 405020 is supplied as a crystalline solid. A stock solution may be made by dissolving the NSC 405020 in the solvent of choice. NSC 405020 is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF), which should be purged with an inert gas. The solubility of NSC 405020 in ethanol and DMF is approximately 30 mg/ml and 20 mg/ml in DMSO.

NSC 405020 is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, NSC 405020 should first be dissolved in ethanol and then diluted with the aqueous buffer of choice. NSC 405020 has a solubility of approximately 0.25 mg/ml in a 1:3 solution of ethanol:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

## Description

NSC 405020 binds to the hemopexin (PEX) domain of the membrane type-1 matrix metalloproteinase (MT<sub>1</sub>-MMP/MMP-14).<sup>1</sup> It disrupts MT<sub>1</sub>-MMP homodimerization without affecting its catalytic activity in an assay using the MT<sub>1</sub>-MMP CAT domain, which is involved in its tumor promoting function (IC<sub>50</sub> > 100  $\mu$ M). In a mouse tumor xenograft model, it reduces tumor growth through a cell death-independent mechanism when injected intratumorally at a dose of 0.5 mg/kg three times per week for two weeks. NSC 405020  $(1 \mu M)$  applied to HSC5 cells prevents arsenic-mediated invasion while also decreasing the expression of MT1-MMP and pERK.<sup>2</sup>

## References

- 1. Remacle, A.G., Golubkov, V.S., Shiryaev, S.A., et al. Novel MT<sub>1</sub>-MMP small-molecule inhibitors based on insights into hemopexin domain function in tumor growth. Cancer Res. 72(9), 2339-2349 (2012).
- Thang, N.D., Yajima, I., Kumasaka, M.Y., et al. Bidirectional functions of arsenic as a carcinogen and an 2. anti-cancer agent in human squamous cell carcinoma. PLoS One 9(5), e96945 (2014).

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.

## WARRANTY AND LIMITATION OF REMEDY

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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