# PRODUCT INFORMATION



# **SAICAR** (hydrate)

Item No. 14087

Formal Name: N-[[5-amino-1-(5-O-phosphono-β-

D-ribofuranosyl)-1H-imidazol-4-yl]

carbonyl]-L-aspartic acid, hydrate

Synonyms: Succino-AICAR, Succinyl-5-

aminoimidazole-4-carboxamide-1-

ribose-5'-phosphate

MF:  $C_{13}H_{19}N_4O_{12}P \bullet XH_2O$ 

FW: 454.3 **Purity:** ≥98% Supplied as: A solid -20°C Storage: Stability: ≥4 years

• XH<sub>2</sub>O NH<sub>2</sub>

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

## **Laboratory Procedures**

SAICAR (hydrate) is supplied as a solid. A stock solution may be made by dissolving the SAICAR (hydrate) in the solvent of choice, which should be purged with an inert gas. SAICAR (hydrate) is slightly soluble in

SAICAR (hydrate) is sparingly soluble in aqueous solutions. To enhance aqueous solubility, dilute the organic solvent solution into aqueous buffers or isotonic saline. If performing biological experiments, ensure the residual amount of organic solvent is insignificant, since organic solvents may have physiological effects at low concentrations. We do not recommend storing the aqueous solution for more than one day.

### Description

SAICAR is an intermediate in the de novo purine nucleotide synthesis pathway. It is reported to stimulate pyruvate kinase isoform M2 and to promote cancer cell survival in conditions of limited glucose availability. 1

#### Reference

1. Keller, K.E., Tan, I.S., and Lee, Y.-S. SAICAR stimulates pyruvate kinase isoform M2 and promotes cancer cell survival in glucose-limited conditions. Science 338(6110), 1069-1072 (2012).

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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