

ProductInformation

NICOTINIC ACID ADENINE DINUCLEOTIDE PHOSPHATE SODIUM SALT Sigma Prod. No. N5655

CAS NUMBER: 5502-96-5

SYNONYMS: Nicotine-TPN; ß-NAADP

PHYSICAL DESCRIPTION:

Appearance: white powder

Molecular formula of free acid: C₂₁H₂₇N₆O₁₈P₃

Molecular weight of free acid: 744.4

 $E^{mM}(260nm) = 18 (0.1 M phosphate, pH 7)^{1}$

 $E^{mM}(260nm) = 17.5(pH 12)^2$

Extensive analytical data have been published.^{2,3}

STORAGE / STABILITY AS SUPPLIED:

ß-NAADP showed less than 2% loss per year when stored at -20°C with desiccant.¹

SOLUBILITY / SOLUTION STABILITY:

β-NAADP dissolves readily in water at 50 mg/mL, giving a clear colorless solution. Solutions of 10 mg/mL stored at 2-8°C at neutral pH are expected to lose 2-5% per week. Solutions stored at -20°C lost between 2-5% after six months.¹

METHOD OF PREPARATION:

This was enzymatically prepared by substituting nicotinic acid for nicotinamide (of NADP). It was purified using ion exchange chromatography, followed by precipitation and desiccation. The compound can be chemically prepared (by alkaline treatment of NADP) or enzymatically prepared. ²⁻⁵

GENERAL REMARKS:

 $\&Bar{B}$ -NAADP, an analog of $\&Bar{B}$ -NADP, is a specific activator of intracellular calcium ion release from sea urchin egg homogenates, active in nanomolar concentrations ($ED_{50} = 16 \text{ nM}^4$; 160 nM for total activation⁵). (HPLC-purified NADP does not trigger such release.) The mechanism appears distinct from the release systems triggered by inositol 1,4,5-triphosphate (IP_3) or by cyclic ADP-ribose (cADPR), since the NAADP-induced Ca^{2^+} release is not blocked by heparin (antagonist of IP_3) or by procaine or ruthenium red (antagonists of cADPR). The mechanism is selectively blocked by thionicotinamide-NADP which does not inhibit IP_3 or cADPR.

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REFERENCES:

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- 2. Imai, T., J. Biochem., 118, 196-203 (1995).
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- 4. Chini, E.N., Beers, K.W. and Dousa, T.P., *J. Biol. Chem.*, 270, 3216-3223 (1995).
- 5. Lee, H.C. and Aarhus, R., J. Biol. Chem., 270, 2152-2157 (1995).

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