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Product Information

β-Nicotinamide adenine dinucleotide sodium salt N0632

Product Description

CAS Registry Number: 20111-18-6

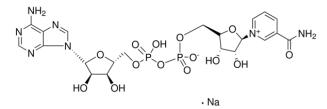
Molecular Formula: C₂₁H₂₆N₇O₁₄P₂Na

Molecular Weight: 685.4

Synonyms: β-NAD, β-DPN

Extinction coefficient (free acid): $E^{mM} = 18.0$ (260 nm, pH 7)¹

Structure:



β-NAD is a pyridine nucleotide and biologically active form of nicotinic acid. β-NAD is a coenzyme necessary for the catalytic reaction of certain enzymes. β-NAD is a carrier for hydride ion, forming β-NADH. Hydride ion is enzymatically removed from a substrate molecule by the action of dehydrogenases, such as malic dehydrogenase and lactic dehydrogenase. Such enzymes catalyze the reversible transfer of a hydride ion from malate or lactate to β-NAD to form the reduced product, β-NADH. Unlike β-NAD, which has no absorbance at 340 nm, β-NADH absorbs at 340 nm (E^{mM} = 6.22). The increase in absorbance at 340 nm with the formation of β-NADH is the basis for measurement of activity of many enzymes.^{2,3}

Many metabolites and enzymes of biological interest are present in tissues at low concentrations. With the use of β -NAD as a catalyst intermediate and several enzymes in a multistep system, known as enzyme cycling, much greater sensitivity for detection of these components is achieved. The reduced form, β -NADH, is fluorescent, whereas β -NAD is not. This difference in fluorescence provides a sensitive fluorescent measurement of the oxidized or reduced pyridine nucleotides at concentrations down to 10^{-7} M.^{3,4} Several theses⁵⁻⁸ and dissertations⁹⁻¹⁷ have cited use of N0632 in their research protocols.

Precautions and Disclaimer

For R&D use only. Not for drug, household, or other uses. Please consult the Safety Data Sheet for information regarding hazards and safe handling practices.

Storage/Stability

 β -NAD is very hygroscopic and should be stored desiccated,¹⁸ at -20 °C.

Solubility

 β -NAD, sodium salt, is tested for solubility in water at 50 mg/mL.

Aqueous solutions between pH 2-6, when stored as single-use aliquots at -70 °C, are stable for at least 6 months. Neutral or slightly acidic solutions are stable at 0 °C for at least 2 weeks. Solutions are rapidly degraded upon heating and are very labile in alkaline solutions, especially in the presence of phosphate, maleate, or carbonate. The rates of degradation of solutions at different pH and temperature conditions have been reported. Solutions are also sensitive to light.^{19,20}

References

- 1. Specifications and Criteria for Biochemical Compounds, 3rd edition. National Academy of Sciences (Washington, DC), p. 87 (1972).
- 2. Bergmeyer, Hans-Ulrich, *Methods of Enzymatic Analysis*, Volume 4. Academic Press, Inc. (New York, NY), pp. 2066-2072 (1974).
- Passonneau, J.V., and Lowry, O.H., *Enzymatic Analysis, A Practical Guide*. The Humana Press, Inc. (Totowa, NJ), pp. 3-4 (1993).
- 4. Passonneau and Lowry, pp. 85-110.



- Damian, Alexis, "An Electrochemical Study of NAD⁺ Interaction with a Polycrystalline Gold Surface". McGill University, M.Eng. thesis, p. 10 (2006).
- Gudiksen, Anders, "AICAR-mediated PDH and GS regulation in skeletal muscle in wild type and PGC-1a MKO mice ". University of Tasmania, M.Sc. thesis, p. 100 (2013).
- Schlattl, Wolfgang, "Protein Folding Mechanism: Analysing the Folding Mechanism of myPGK Utilizing Simulations and Experiments". Universität Wien, Mag. pharm. thesis, p. 19 (2014).
- Škovroňová, Renata, "Renální *in vitro* modely v toxikologii" ("Renal *in vitro* models in toxicology"). Masarykova Univerzita, M.Sc. thesis, p. 54 (2018).
- Liaw, Ean-Tun, "The Relative Activity of the Cellulose Enzyme System of *Trichoderma reesei* with Native and Modified Cellulosic Substrates". Oregon State University, Ph.D. dissertation, p. 61 (1994).
- Rajendran, Ramkumar, "The transcriptional cofactor PCAF as mediator of the interplay between p53 and HIF-1 alpha and its role in the regulation of cellular energy metabolism ". University of Manchester, Ph.D. dissertation, p. 67 (2011).
- Zhou, Gaofeng, "Enhancing Aluminium Resistance in Barley through Over-expression of *MATE* Genes". University of Tasmania, Ph.D. dissertation, p. 27 (2012).
- 12. Guest, Patrick, "The identification and characterisation of novel inhibitors of the 17 β -HSD10 enzyme for the treatment of Alzheimer's disease ". University of St. Andrews, Ph.D. dissertation, pp. 75, 80 (2016).
- Stiers, Kyle M., "Molecular Mechanisms of Enzyme Dysfunction in Human Phosphoglucomutase-1 Deficiency". University of Missouri – Columbia, Ph.D. dissertation, p. 95 (2019).
- 14. Kaschubowski, Klaus Eric, "Use of Fluorescent Sensors to Visualize P2X7-Mediated Changes in Local ATP Concentrations in the Cytosol and at the Cell Surface". Universität Hamburg, Dr. rer. hum. biol. dissertation, p. 23 (2020).

- 15. Abbas, Hafsa, "Glyoxalase 1 overexpressionassociated multi-drug resistance in cancer chemotherapy". University of Warwick, Ph.D. dissertation, p. 147 (2020).
- Khachatoorian, Careen, "Identification, Quantification, and Evaluation of the Cytotoxicity of Electronic Cigarette Exhaled Aerosol Residue (ECEAR)". University of California Riverside, Ph.D. dissertation, p. 94 (2020).
- Geller, Sarah G., "Establishment of Metabolomics Methods Based on Microseparations Coupled to Mass Spectrometry". Northeastern University, Ph.D. dissertation, p. 94 (2021).
- 18. *The Merck Index*, 13th ed., Entry# 6370.
- Dawson, R.M.C. *et al.* (eds.), *Data for Biochemical Research*, 3rd edition. Oxford University Press (New York, NY), pp. 130-131 (1986).
- 20. Lowry, O. H. *et al.*, *J. Biol. Chem.*, **236(10)**, 2756-2759 (1961).

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