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REGULATORY STATUS STATEMENT

β-Nicotinamide Mononucleotide 99.0%

This statement summarises the regulatory position of β-Nicotinamide Mononucleotide 99.0% in the principal markets as of the document date and supports the customer's own compliance assessment of its finished product. It is provided for general guidance and does not constitute legal advice.

1. Regulatory status in major markets

Market / jurisdiction	Regulatory framework	Status / classification	Basis / documentation	Remarks
United States (FDA)	FD&C Act; DSHEA 1994	Not a lawful dietary ingredient: the FDA has stated that NMN is excluded from the definition of a dietary supplement. No New Dietary Ingredient notification or GRAS notice is held by us.	FD&C Act 201(ff)(3)(B); FDA correspondence 2022	Supply is for research and industrial use only.
European Union (EFSA / European Commission)	Regulation (EU) 2015/2283 (Novel Food)	Not authorised as a novel food; no novel-food authorisation is held by us and no application has been lodged.	Reg. (EU) 2015/2283	Not for food use in the EU.
United Kingdom (FSA)	UK Novel Food framework	Not authorised; the substance does not appear on the UK novel-food authorisation register.	UK Novel Food regime	Not for food use in the UK.
Canada (Health Canada / NNHPD)	Natural Health Products Regulations	No Natural Product Number or product licence is held by us.	NHP Regulations	Customer to confirm acceptability before any use.
Australia / New Zealand (TGA / FSANZ)	Therapeutic Goods Act; Food Standards Code	Not listed on the Australian Register of Therapeutic Goods and not permitted as a food ingredient.	TGA; FSANZ	Not for therapeutic or food use.
China (SAMR / NMPA)	Food-safety and health-food raw-material requirements	Supplied as a research raw material for export; not registered for domestic food use.	Food-safety law; GB standards	Export-oriented.

2. Other

• Not listed on the WADA Prohibited List; not a controlled substance in major sports jurisdictions.

• The substance is listed in the U.S. TSCA inventory and is subject to EU REACH where applicable.

Classification applies to the ingredient as supplied; finished-product and supplement status remain the customer's responsibility.

Prepared by: Mia Yu Approved by: Charlie Wang Date: Nov. 06, 2026