Stevia-Erythritol Blend 1%

Steviol glycosides 1% on erythritol · CAS — · Spray-dried or dry-blended steviol glycosides on erythritol

INDICATIVE SPECIFICATION — CONFIRMED IN WRITING AT QUOTATION

IDENTITY / FORM	FORMULA	CAS	PRIMARY ASSAY
Spray-dried or dry-blended steviol glycosides on erythritol	Steviol glycosides 1% on erythritol	—	0.9–1.1% (HPLC-UV)
METHOD	**MINIMUM ORDER**	**LEAD TIME**	**STANDARD PACK**
HPLC-UV for steviol glycosides and HPLC-RI for erythritol	1 kg	3–10 days	1 kg foil pouch

SPECIFICATION

PARAMETER	SPECIFICATION	METHOD / NOTE
Steviol glycosides	0.9–1.1% (HPLC-UV)	Analytical
Rebaudioside A	0.5–0.8% (HPLC-UV)	Analytical
Erythritol	≥98.0% (HPLC-RI)	Analytical
Appearance	White to off-white powder	Specification
Moisture	≤0.5%	Specification
Solubility	Soluble in water	Specification
Particle size	≥95% through 60 mesh	Sieve
Homogeneity	Specified by the blend protocol	Specification
Heavy metals	Pb ≤0.5 · As ≤0.5 · Cd ≤0.5 · Hg ≤0.1 mg/kg	Specification
Microbiology	APC ≤1,000 cfu/g · yeast and mould ≤100 cfu/g · E. coli and Salmonella absent	Specification
Marker / profile	Steviol glycosides 0.9–1.1%, erythritol ≥98%	Identity / ratio
Appearance	White to off-white powder	Visual
Solubility	Soluble in water; the erythritol dissolves endothermically.	Matrix dependent
Standard packaging	1 kg foil pouch · 25 kg fiber drum	Food-grade liner

APPLICATIONS

- Tabletop
- Beverages

AVAILABLE FORMS

- Powder, 60 mesh
- Granular

- Dairy alternatives
- Confectionery
- Supplements
- Custom ratio

WHAT TO CONFIRM BEFORE ORDERING

- Confirm the exact identity, carrier and particle size for Stevia-Erythritol Blend 1% before the first order.
- Taste at the target sucrose equivalence in the real matrix, not in water alone.

Indicative document. This sheet describes the grade we can quote; it is not a batch COA and does not confirm a result for a specific lot. Final specification, contaminant limits and documentation are agreed in writing before the order. The identity, assay basis, profile and method are release parameters, not marketing descriptions.

DOCUMENTATION RELEASED WITH EACH LOT

DOCUMENT	CONTENT	SCOPE
Certificate of analysis	Assay, identity, physical parameters, contaminant panel and the release decision for the lot.	Batch-specific
Technical data sheet	Identity, form, CAS, specification ranges, solubility, bulk density, storage and packing.	Product-level
Safety data sheet	Sixteen sections covering hazards, handling, storage, transport and emergency response.	Grade-specific
Allergen and GMO statements	Major allergens, gluten, animal-derived inputs and GMO status.	Batch-specific
Origin and traceability record	Manufacturing site and the link between the lot and the retained sample.	Batch-specific
Residual-solvent statement	The crystallisation system and the solvent class, where applicable.	Batch-specific

HOW TO REQUEST THIS GRADE

01 · BRIEF

Send the target specification, quantity, application and destination market.

02 · COUNTER-SPEC

We confirm what can be matched and the documents available for the grade.

03 · SAMPLE & QUOTE

A sample with its own COA, followed by packing, lead time and terms in writing.

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