Urolithin A

$C_{13}H_8O_4$ · CAS 1143-70-0 · Urolithin A, powder

INDICATIVE SPECIFICATION — CONFIRMED IN WRITING AT QUOTATION

CHEMICAL FORM	FORMULA	CAS	PRIMARY ASSAY
Urolithin A, powder	$C_{13}H_8O_4$	1143-70-0	98.0–102.0%
METHOD	**MINIMUM ORDER**	**LEAD TIME**	**STANDARD PACK**
HPLC	1 kg	1–7 days	1 kg foil pouch

SPECIFICATION

PARAMETER	SPECIFICATION	METHOD / NOTE
Assay (HPLC)	98.0–102.0%	Specification
Appearance	Off-white to yellow powder	Specification
Identification	HPLC and NMR conform	Analytical
Loss on drying	≤1.0%	Specification
Residual solvents	Methanol ≤3,000 ppm	Specification
Heavy metals	Pb ≤1 · As ≤1 · Cd ≤1 · Hg ≤0.1 mg/kg	Specification
Microbiology	APC ≤1,000 cfu/g · yeast and mould ≤100 cfu/g · E. coli and Salmonella absent	Specification
Particle size	≥95% through 40 mesh	Sieve
Marker set	Urolithin A 98.0–102.0% (HPLC)	Identity / ratio
Appearance	Off-white to yellow, free-flowing powder	Visual
Solubility	Sparingly soluble in water; soluble in dimethyl sulfoxide.	Matrix dependent
Standard packaging	1 kg foil pouch · 25 kg fiber drum	Food-grade liner

APPLICATIONS

- Capsules and tablets
- Powder sticks and drink mixes

AVAILABLE FORMS

- Powder, 40 mesh
- Granulated on request
- Dilution on request

WHAT TO CONFIRM BEFORE ORDERING

- Confirm the destination market and the specific authorization before quoting; urolithin A is a novel food in some markets and not in others.
- Ask for the assay and the residual-solvent profile; the purity of the lot is the starting point, not the regulatory answer.
- Urolithin A is light-sensitive; the pack and the process exposure belong in the specification.

- Do not use the page as a label or a claim: the finished product's status is a separate legal question.

> **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 form, hydrate state, ligand ratio 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	Form, hydrate state, 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, dosage form 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.

No claim is made here. This sheet does not claim a therapeutic effect, a product certification or a regulatory approval. Health claims belong to the brand that makes them and to the market that authorises them.

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