Chondroitin Sulfate

Chondroitin sulfate sodium (glycosaminoglycan mixture) · CAS 9007-28-7 · Chondroitin sulfate sodium, bovine cartilage

INDICATIVE SPECIFICATION — CONFIRMED IN WRITING AT QUOTATION

CHEMICAL FORM	FORMULA	CAS	PRIMARY ASSAY
Chondroitin sulfate sodium, bovine cartilage	Chondroitin sulfate sodium (glycosaminoglycan mixture)	9007-28-7	90.0–105.0% (CPC titration)
METHOD	**MINIMUM ORDER**	**LEAD TIME**	**STANDARD PACK**
CPC titration	1 kg	1–7 days	1 kg foil pouch

SPECIFICATION

PARAMETER	SPECIFICATION	METHOD / NOTE
Assay (dried basis)	90.0–105.0% (CPC titration)	Specification
Appearance	White to off-white powder	Specification
Identification	IR and electrophoretic profile conform	Specification
Loss on drying	≤10.0%	Specification
pH (5% solution)	5.5–7.5	Specification
Protein	≤6.0%	Specification
Heavy metals	Pb ≤3 · As ≤2 · 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 80 mesh	Sieve
Marker set	Chondroitin sulfate sodium 90.0–105.0% (dried basis)	Identity / ratio
Appearance	White to off-white, free-flowing powder	Visual
Solubility	Soluble in water; the solution is viscous.	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, 80 mesh
- Granulated on request
- Milled to 100 mesh on request

WHAT TO CONFIRM BEFORE ORDERING

- Name the source and the salt: bovine, porcine and shark cartilage are different label ingredients and different BSE/TSE files.
- Ask for the assay method and the electrophoretic profile; a total glycosaminoglycan figure is not the same as chondroitin sulfate.
- Confirm the protein and moisture limits; both change the dose and the stability.
- Chondroitin sulfate is hygroscopic once dried; the pack and the dose-room humidity belong in the specification.

> **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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