Chymotrypsin

— · CAS 9004-07-3 · Chymotrypsin, powder

INDICATIVE SPECIFICATION — CONFIRMED IN WRITING AT QUOTATION

CHEMICAL FORM	FORMULA	CAS	PRIMARY ASSAY
Chymotrypsin, powder	—	9004-07-3	≥1,000 USP U/mg
METHOD	**MINIMUM ORDER**	**LEAD TIME**	**STANDARD PACK**
USP activity assay (ATEE)	1 kg	1–7 days	1 kg foil pouch

SPECIFICATION

PARAMETER	SPECIFICATION	METHOD / NOTE
Activity (USP units)	≥1,000 USP U/mg	Specification
Appearance	White to off-white powder	Specification
Identification	Activity assay and IR conform	Specification
Loss on drying	≤5.0%	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 40 mesh	Sieve
Marker set	Chymotrypsin ≥1,000 USP U/mg	Identity / ratio
Appearance	White to off-white, free-flowing powder	Visual
Solubility	Freely soluble in water; the solution is clear.	Matrix dependent
Standard packaging	1 kg foil pouch · 25 kg fiber drum	Food-grade liner

APPLICATIONS

- Capsules and tablets
- Powder blends

AVAILABLE FORMS

- Powder
- Granulated on request

WHAT TO CONFIRM BEFORE ORDERING

- Name the assay: chymotrypsin is reported in USP units by the ATEE method; a trypsin unit is a different number.
- Ask for the trypsin limit and the autolysis profile; a protease blend is not the same as purified chymotrypsin.
- The enzyme loses activity with heat and moisture; the pack and the storage condition 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.

Shaanxi Rainwood Biotech Co., Ltd. · Rm 218, Tangxing Digital Bldg, #6 Tangxing Rd, Xi'an, Shaanxi, China

info@rainwoodbio.com · nutrition.rainwoodbio.com