Phosphatidylserine

$C_{42}H_{82}NO_{10}P$ · CAS 51446-62-9 · Soy-derived phosphatidylserine 50% powder

INDICATIVE SPECIFICATION — CONFIRMED IN WRITING AT QUOTATION

CHEMICAL FORM	FORMULA	CAS	PRIMARY ASSAY
Soy-derived phosphatidylserine 50% powder	$C_{42}H_{82}NO_{10}P$	51446-62-9	50.0–55.0% (HPLC-ELSD)
METHOD	**MINIMUM ORDER**	**LEAD TIME**	**STANDARD PACK**
HPLC-ELSD	1 kg	1–7 days	1 kg foil pouch

SPECIFICATION

PARAMETER	SPECIFICATION	METHOD / NOTE
Phosphatidylserine	50.0–55.0% (HPLC-ELSD)	Analytical
Appearance	Light yellow to yellow powder	Specification
Identification	HPLC-ELSD profile conforms	Analytical
Loss on drying	≤5.0%	Specification
Peroxide value	≤10 meq/kg	Specification
Acid value	Reported	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
Source	Soybean lecithin	Specification
Marker set	Phosphatidylserine (50.0–55.0%, HPLC-ELSD)	Identity / ratio
Appearance	Light yellow to yellow, free-flowing powder	Visual
Solubility	Dispersible in water; not a clear solution. Soluble in chloroform and other non-polar solvents.	Matrix dependent
Standard packaging	1 kg foil pouch · 25 kg fiber drum	Food-grade liner

APPLICATIONS

- Capsules and tablets
- Powder sticks and drink mixes
- Gummies and chews

AVAILABLE FORMS

- 50% powder, 80 mesh
- 20% or 70% on request
- Granulated on request

WHAT TO CONFIRM BEFORE ORDERING

- Name the source and the PS content: soy and sunflower PS are different label ingredients and different allergen declarations.
- Ask for peroxide and acid values; an assay can pass while the lipid has oxidised.
- Confirm the HPLC-ELSD method and the phospholipid profile, not only the PS percentage.
- PS powder is hygroscopic and prone to oxidation; the pack and the storage temperature 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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