Reishi Mushroom Extract (Ganoderma lucidum) – Triterpenes 2–20%

Ganoderma lucidum · Ganodermataceae

DOCUMENT	PRODUCT CODE	DOCUMENT NO.	REVISION	PRODUCT FORM
Technical Data Sheet	AP0513	TDS-AP0513	1.0 · 24 October 2026	Mushroom Extract

1 Product description

Reishi extract is produced from the fruiting body / spores of Ganoderma lucidum (Ganodermataceae family) and supplied as a standardized mushroom extract, standardized to Triterpenes. Available grades: 2%, 4%, 10%, 20%. Typical sourcing: China, Japan, Korea. Intended for B2B formulation in nutraceuticals & supplements.

2% 4% 10% 20%

2 Identification

Botanical source	*Ganoderma lucidum* (Ganodermataceae)
Common name(s)	Reishi, Lingzhi
Plant part used	Fruiting Body (mushroom)
Typical origin	China, Japan, Korea
Marker compound	Triterpenes (Triterpenoids & triterpene saponins)
Manufacturing process	Hot-water and/or hydro-alcoholic extraction of fruiting body or mycelium, concentrated and spray-dried
Standardisation	2–20% (HPLC, UV-Vis spectrophotometry)
Assay method	HPLC, UV-Vis spectrophotometry
Pharmacopoeia & standards	USP, Chinese Pharmacopoeia, American Herbal Pharmacopoeia

3 Specification

PARAMETER	SPECIFICATION	METHOD
Appearance	Light brown to brown fine powder	Visual
Odour	Characteristic	Organoleptic
Identification	Conforms (HPTLC / HPLC fingerprint vs. reference)	HPTLC / HPLC
Assay – Triterpenes	Not less than the labelled grade (2%, 4%, 10%, 20%)	HPLC, UV-Vis spectrophotometry
Loss on drying	≤ 5.0%	USP ‹731›
Total ash	≤ 5.0%	USP ‹561›
Particle size	≥ 95% through 80 mesh (180 µm)	USP ‹786›
Bulk density	Report (typically 0.40–0.70 g/mL)	USP ‹616›
Residual solvents	Complies with ICH Q3C / USP ‹467›	GC-HS
Lead (Pb)	≤ 3.0 mg/kg	ICP-MS
Arsenic (As)	≤ 2.0 mg/kg	ICP-MS
Cadmium (Cd)	≤ 1.0 mg/kg	ICP-MS
Mercury (Hg)	≤ 0.1 mg/kg	ICP-MS
Pesticide residues	Complies with USP ‹561› and Regulation (EC) No 396/2005	GC-MS/MS, LC-MS/MS
Aflatoxins	B1 ≤ 2 µg/kg; sum of B1, B2, G1, G2 ≤ 4 µg/kg	HPLC-FLD
Total aerobic microbial count	≤ 10,000 cfu/g	USP ‹2021›

Yeasts & moulds	≤ 1,000 cfu/g	USP ‹2021›
Escherichia coli	Absent in 10 g	USP ‹2022›
Salmonella spp.	Absent in 25 g	USP ‹2022›
Staphylococcus aureus	Absent in 10 g	USP ‹2022›

Limits above are the standard release specification for this product class. Each lot is released against its Certificate of Analysis, which is the governing document and is available on request.

4 Grades & pack sizes

GRADE	SKU PREFIX	PACK SIZES
2%	AP0513-2	Sample (50 g) · 250 g · 1 Kg · 5 Kg · 25 Kg
4%	AP0513-4	Sample (50 g) · 250 g · 1 Kg · 5 Kg · 25 Kg
10%	AP0513-10	Sample (50 g) · 250 g · 1 Kg · 5 Kg · 25 Kg
20%	AP0513-20	Sample (50 g) · 250 g · 1 Kg · 5 Kg · 25 Kg

5 Packaging

- Samples: food-grade HDPE jar or aluminium-laminate pouch.
- 250 g – 1 Kg: food-grade LDPE bag sealed in an aluminium-laminate pouch.
- 5 Kg – 25 Kg: double food-grade LDPE bags in a fibre drum or HDPE pail.
- Packaging may vary by lot and is stated on the packing list.

6 Storage & shelf life

- Shelf life: 24 months from date of manufacture in the original, unopened packaging.
- Store in a cool (below 25 °C), dry place, protected from light.
- Hygroscopic: reseal immediately after use and minimise exposure to humid air.

7 Applications

Industries	Nutraceuticals & Supplements
Formulation focus	Stress & Sleep Support, Immune Support
Traditional systems	Traditional Chinese Medicine, Medicinal mushrooms

Use levels depend on the finished product and the target market. The customer is responsible for establishing safe and permitted use levels and for any claims made on finished products.

8 Regulatory & compliance

- Pharmacopoeia / standard references: USP, Chinese Pharmacopoeia, American Herbal Pharmacopoeia.
- Allergen information: not derived from a source listed in Annex II of Regulation (EU) No 1169/2011.
- Regulatory status for food, supplement, cosmetic or pharmaceutical use (e.g. EU novel food, US NDI/GRAS, FSSAI) must be confirmed by the customer for the intended market and use.

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