Senna Extract (Senna alexandrina) – Sennoside B 1–30%

Senna alexandrina · Fabaceae

DOCUMENT	PRODUCT CODE	DOCUMENT NO.	REVISION	PRODUCT FORM
Technical Data Sheet	AP1114	TDS-AP1114	1.0 · 24 October 2026	Standardized Extract

1 Product description

Senna extract is produced from the leaf / pod of Senna alexandrina (Fabaceae family) and supplied as a standardized extract powder, standardized to Sennoside B. Available grades: 1%, 3%, 20%, 30%. Typical sourcing: India, Egypt, Sudan. Intended for B2B formulation in nutraceuticals & supplements.

1% 3% 20% 30%

2 Identification

Botanical source	*Senna alexandrina* (Fabaceae)
Common name(s)	Senna
Plant part used	Leaf, Fruit
Typical origin	India, Egypt, Sudan
Marker compound	Sennoside B (Anthraquinones & anthrones)
Manufacturing process	Water and/or hydro-alcoholic (ethanol) extraction, concentration and spray-drying; the solvent system for each grade is stated on the CoA
Standardisation	1–30% (HPLC)
Assay method	HPLC
Pharmacopoeia & standards	USP, Ph. Eur. (EP), British Pharmacopoeia, Indian Pharmacopoeia, Ayurvedic Pharmacopoeia of India, Japanese Pharmacopoeia, EMA herbal monograph, ESCOP monograph, WHO monograph

3 Specification

PARAMETER	SPECIFICATION	METHOD
Appearance	Yellow-orange to reddish-brown powder	Visual
Odour	Characteristic	Organoleptic
Identification	Conforms (HPTLC / HPLC fingerprint vs. reference)	HPTLC / HPLC
Assay – Sennoside B	Not less than the labelled grade (1%, 3%, 20%, 30%)	HPLC
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
1%	AP1114-1	Sample (50 g) · 250 g · 1 Kg · 5 Kg · 25 Kg
3%	AP1114-3	Sample (50 g) · 250 g · 1 Kg · 5 Kg · 25 Kg
20%	AP1114-20	Sample (50 g) · 250 g · 1 Kg · 5 Kg · 25 Kg
30%	AP1114-30	Sample (50 g) · 250 g · 1 Kg · 5 Kg · 25 Kg

Made-to-order grades: 3%.

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
Traditional systems	Ayurveda, European herbal tradition

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, Ph. Eur. (EP), British Pharmacopoeia, Indian Pharmacopoeia, Ayurvedic Pharmacopoeia of India, Japanese Pharmacopoeia, EMA herbal monograph, ESCOP monograph, WHO monograph.
- **EU hydroxyanthracene restriction:** Hydroxyanthracene derivatives are restricted in foods in the EU by Regulation (EU) 2021/468: preparations from Rheum, Cassia and Rhamnus species are under Union scrutiny (Annex III, Part C).
- 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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