

The Extract Brief

A monthly briefing on the trade in botanical extracts — regulation, quality, supply & freight — compiled for the professional buyer.

NUMBER 1 · VOLUME I · JULY 2026



PLATE I — *Silybum marianum* L., THE MILK THISTLE · A. ACHENE · B. FLORET
THE SPECIMEN OF THIS NUMBER — SEE P. 4

IN THIS NUMBER

- | | | | |
|--|------|--|------|
| I. Regulatory Watch — <i>ashwagandha</i> in Brussels | p. 2 | V. The Specimen — <i>milk thistle</i> | p. 4 |
| II. Quality Desk — <i>lessons of the moringa outbreak</i> | p. 3 | VI. Logistics & Trade — <i>tariffs settle, freight softens</i> | p. 4 |
| III. Supply & Price — <i>turmeric firms</i> | p. 3 | VII. Dates for the Diary & a letter from the desk | p. 4 |
| IV. Knowledge Series — <i>why two honest CoAs disagree</i> | p. 3 | | |

I. Europe moved a step closer to a formal safety review of ashwagandha — the largest Indian botanical export line.	II. A new EU pesticide regulation names concentrates and extracts explicitly; compliance falls due 29 January 2027.	III. US tariffs on Indian goods settled at 18 per cent , down from a peak of 50 — and ocean freight kept softening.
---	--	--

§ 1 Regulatory Watch

BRUSSELS · CANBERRA · THE BORDER

1.1 Ashwagandha edges toward an EU Article 8 review

On the fourteenth of July, a meeting at the European Parliament — hosted by MEP Sean Kelly and organised by Kerry Group — launched *Safeguard Ashwagandha*, an industry initiative to put the botanical's peer-reviewed safety record in front of regulators before any formal assessment begins. At stake is an **Article 8 review** under Regulation (EC) 1925/2006: the process that can end in restrictions, mandatory label warnings, or a ban. No EFSA assessment is open yet; the initiative exists precisely because the sector expects one. Australia's TGA, meanwhile, issued its own ashwagandha safety update on 20 July.

👉 **To do now:** if ashwagandha sits in a 2027 pipeline, ask the supplier for species and plant-part authentication (root versus leaf), the withanolide assay method behind the label claim, and batch documentation fit for a safety dossier. The suppliers who can produce that file quickly are the ones whose material keeps moving.

1.2 The MRL rule that names extracts

Regulation (EU) 2026/1546, adopted **9 July** and applicable **29 January 2027**, revises residue limits for benomyl, carbendazim and thiophanate-methyl across herbs and spices — and, explicitly, *botanicals, concentrates and extracts*. The recital concedes what exporters have long argued: concentration during drying or extraction increases the commercial significance of residues. A raw herb passing at 0.05 mg/kg can yield a 10:1

extract reading several times higher; both certificates are honest, and only one clears the border. Extract-level testing is now the only defensible position.

1.3 Novel Food: two sides of the line

Gotu kola, southern ginseng, wild oregano and white sage are deemed novel in EU food supplements — unauthorised without a Novel Food authorisation — while **pomegranate extract** was confirmed non-novel and stays freely marketable. The Union's year to date: fifteen authorisations against twenty-one rejections.

1.4 Also on the radar

MOAH/MOSH — maximum limits adopted 11 June; relevant to oleoresins and lipid-based extracts.

Berberine — EFSA's public consultation on regulatory status remains open.

Monacolin K — a proposed ban decision is expected before the year closes.

PFAS — banned in EU food-contact packaging from 12 August 2026; check drum-liner specifications.

Border controls — Reg (EU) 2026/194 delisted Indian vanilla and cloves on sustained compliance and set *Piper*, *Capsicum* and *Pimenta* at 20 per cent checks; a June amendment (2026/1206) revised the annexes again. Verify the current listing for each commodity before shipment planning.

§ 2 Quality Desk

THE MORINGA OUTBREAK, READ PROPERLY

On 17 July the CDC and FDA closed their investigation into a multistate outbreak of extensively drug-resistant *Salmonella* traced to moringa leaf-powder capsules: thirty-four illnesses, eleven hospitalisations, and recalls spanning product distributed from December 2025 through July 2026.

The lesson is not *avoid moringa*. It is that powder and extract are different microbiological propositions. A leaf powder is dried plant material — often sun- or air-dried at origin —

carrying whatever bioburden the field and the drying yard left on it. A solvent-extracted, spray-dried extract has passed through steps hostile to vegetative pathogens. The buyer's protection is specification: total plate count, yeast and mould, pathogen absence per 10 g — and the question of *how* the material was treated, since steam sterilisation and ethylene-oxide treatment carry opposite consequences for an EU-bound shipment.

“Leaf powder and leaf extract are different microbiological propositions.”

QUALITY DESK · № 1

Thirty-four case files, eleven hospital admissions and one drug-resistance profile make the argument better than any

sales letter: microbiology belongs on the specification, not in the assumptions.

§ 3 Supply & Price Direction

FROM THE MANDIS

Turmeric is firming. Mandi prices moved from roughly ₹13,900 per quintal in early July to an August average near ₹14,900 — some six per cent inside a month, with Telangana and Andhra Pradesh holding half the planted area. Extract buyers should expect the move to reach curcumin 95% con-

tract pricing on the customary one-quarter lag: fourth-quarter quotations will carry it. The prudent response is neither panic nor pre-buying, but asking the supplier which crop the quote is written against.

§ 4 Knowledge Series

№ 1 — WHY TWO HONEST COAS CAN DISAGREE

Each number of this Brief takes one recurring buyer's question and treats it properly. First: the assay-method problem, the most common source of specification disputes we see. Ultraviolet spectrophotometry measures everything in a sample that absorbs at the target wavelength — the whole actives complex, and anything else absorbing beside it. High-performance liquid chromatography separates the individual compounds and counts only those it names.

Neither method is wrong; they answer different questions. But *forty per cent by UV* and *ten per cent by HPLC* can describe the same powder, and a purchase order that states a number without its method is an invitation to a rejected shipment. The rule that prevents the dispute costs one line: **write the method beside the number.**

Further reading in our library at svbotanica.com/insights: *How to Read a CoA* · *Silymarin: UV v. HPLC* · *Withanolides: Gravimetric v. HPLC* · *Bacosides: HPLC v. UV*.

FROM THE DESK
OF SV BOTANICA

MUMBAI · AUGUST
2026

Every quarter a buyer sends us a raw-material pesticide report and asks why the extract needs testing of its own. It is always asked in good faith — the herb passed; surely the extract passes too. But extraction is concentration, and residues concentrate along with the actives. We have spent years making that argument specification by specification; this month, Regulation 2026/1546 made it for us, in a legal text with a compliance date attached. If an incoming-QC protocol still tests the herb and not the extract, January 2027 is the deadline for changing it — and the supplier should be volunteering the extract-level data before being asked.

— The desk

§ 5 The Specimen: Milk Thistle

SILYBUM MARIANUM L. · ASTERACEAE



PLATE I (DETAIL) — FLOWERING HEAD

BOTANICAL	<i>Silybum marianum</i> L., seed (fruit)
ACTIVES	Silymarin — flavonolignan complex: silybin A&B, isosilybin A&B, silychristin, silydianin
TRADE GRADE	80% silymarin by UV (≈ 30% silibinin by HPLC)
MONOGRAPHS	Ph. Eur. & USP — both quantify on the HPLC / silibinin basis
WATCH FOR	Cutting with cheaper <i>Carduus</i> thistle seed; UV-only specifications

Milk thistle is one of the world's highest-volume liver-support ingredients, and the perfect illustration of this number's Knowledge Series. Silymarin is not one molecule but a complex of six flavonolignans, and the market's workhorse grade — **eighty per cent by UV** — describes the same powder that reads near thirty per cent when assayed by HPLC as silibinin. Two honest numbers; one material; endless disputes between buyers who compare one supplier's UV figure against another's HPLC figure.

Three things belong on every milk thistle specification. The **method beside the number**, first and always. An **HPLC identity fingerprint** second: silymarin content drives price, which makes the seed a known target for economically motivated cutting with cheaper thistles, and a UV-only specification is the easiest to fool. Third, the **EU-entry trio** — heavy metals by ICP-MS, residual solvents, and ethylene-oxide status — which decides whether the shipment clears customs at all.

Demand context: the liver-health category continues to grow on alcohol-alternative and metabolic-wellness positioning, and silymarin remains its anchor active. The full dossier — buyer's guide, regulatory notes, sourcing, adulteration — stands in our library at svbotanica.com/insights.

§ 6 Logistics & Trade

TARIFF · FREIGHT · CARBON

Tariff. United States duties on Indian goods now stand at 18 per cent, down from a peak of fifty, the February interim agreement having rescinded the punitive tranche. India's AYUSH and herbal exports to the American market ran to some \$689 million in FY 2024–25, led by ashwagandha and turmeric. **Freight.** The Drewry index fell four per cent to \$4,374 per forty-foot box in the week ending 23 July; India–

Europe spot sits at \$3,500–4,500 with Cape routing adding ten to fourteen days.

Carbon. The EU's emissions-trading scheme reached full cost recovery this year, adding an estimated \$150–400 per container on Asia–Europe lanes. The charge is structural, not cyclical; it belongs in every landed-cost model from here on.

§ 7 Dates for the Diary

DATE	WHAT HAPPENS
12 · VIII · 2026	PFAS ban in EU food-contact packaging applies
Q3 · 2026	EU consultation opens on harmonised vitamin & mineral maximum levels
Late 2026	Monacolin K ban decision expected
29 · I · 2027	Regulation (EU) 2026/1546 pesticide MRLs apply — extracts in scope



SV Botanica

PREMIUM NATURAL INGREDIENTS

SV Botanica is an India-based **B2B supplier and exporter** of standardised botanical ingredients for supplement, food and cosmetic manufacturers worldwide. Every material ships export-ready with full documentation — batch Certificate of Analysis, HPLC assay, specification sheet — and is tested against the standards our buyers' markets actually enforce: EU pesticide MRLs, heavy metals and ethylene oxide, with third-party laboratory verification.

HERBAL EXTRACTS · NATURAL OILS · SPICES & OLEORESINS · NUTRACEUTICALS

GMP · FSSAI · APEDA · ISO · ORGANIC — EU MRL · HEAVY METALS · ETO TESTED

Sourcing for 2027? Send the specification — grade, assay method, destination market — and we respond within twenty-four hours.

info@svbotanica.com · +91 8767675175 · svbotanica.com

№ 2 — AUGUST 2026

The regulatory month, the Knowledge Series continued, and a new specimen. Every number at svbotanica.com/insights.

The Extract Brief is published monthly by SV Botanica as a good-faith orientation for professional ingredient buyers. Regulatory items are summarised from public sources current at publication; always verify against the official text before acting. Nothing herein constitutes legal, regulatory or investment advice.

© 2026 SV Botanica, India · read this number online at svbotanica.com/insights/extract-brief-july-2026