

Stay ahead of regulatory change — before it affects your business.

PlainReg AI is a bespoke consultancy that monitors regulatory developments, structures obligations into actionable intelligence, and builds compounding knowledge bases — purpose-built for your product categories and jurisdictions. Every output cites its official source, so everything your team acts on is traceable and defensible.

RegChange Monitor

Continuous regulatory monitoring

- AI scores each item 0–100 for relevance to your products
- Deduplicated — the same alert never arrives twice
- Digest email with CSV attachment, one click to trigger

CAS Watch

Substance-level change monitoring

- Paste CAS numbers — matches highlight with an instant count
- SVHC listings, authorisation deadlines, and sunset dates
- One-click export of the matched view to CSV or JSON

Extraction Service

On-demand obligation analysis

- Every obligation, condition, and deadline in a filterable table
- Each item links to its exact source page
- Export to XLSX or JSON for your compliance workflow

RegWiki

A living regulatory knowledge base

- New regulations update the whole base — not just add a page
- Plain-English questions, cited and traceable answers
- Self-auditing: contradictions and gaps flagged automatically

A complete regulatory intelligence loop

From early signal to structured knowledge — your team focuses on decisions, not document triage.

1 Monitor detects

RegChange and CAS Watch surface relevant change early, scored against your profile.



2 Obligations structured

Extraction pulls every obligation and deadline into a cited, filterable table.



3 Knowledge compounds

RegWiki integrates it all — ask plain-English questions, get cross-referenced answers.

CAS Watch — your watchlist against live regulatory change

Paste your substance list. Matches highlight instantly — every row cites its official public source.

2 of your 5 watched substances have new regulatory activity

Export CSV · JSON

CAS #	Substance	Source	Change	Status
80-05-7	Bisphenol A (BPA)	SVHC Candidate List	Endocrine disruptor	Listed
117-81-7	DEHP (phthalate)	Annex XIV list	Authorisation req.	In force
7439-92-1	Lead	SVHC Candidate List	Reprotoxic	Listed
1310-73-2	Sodium hydroxide	C&L Inventory	No change	—
64-17-5	Ethanol	C&L Inventory	No change	—

Illustrative example — public ECHA data and public example CAS numbers only.

Extraction Service — EU MDR 2017/745 sample

Upload any regulatory document. Every obligation lands in a structured table, cited to the exact page.

Clause	Type	Topic	Page
Manufacturers shall establish, document, implement and maintain a system for risk management	Obligation	Risk management	p.34
Technical documentation shall be drawn up before the device is placed on the market	Requirement	Technical docs	p.41
Each device shall be assigned a unique device identifier (UDI)	Requirement	UDI / labelling	p.56
Post-market surveillance system shall be proportionate to the risk class	Obligation	Post-market	p.78

Exports to XLSX with all clauses, citations, and deadlines.

Every output cites its official source.

Traceable, validated, defensible — built for teams that can't act on AI output they can't verify.

Try the live extraction demos — no setup or login required

plainregai.com · SB 343 (California Recycling) · Oregon EPR (Plastic Pollution Act)

Talk to us about your challenge: jaz@riskaversetechcompany.com