

Schedule Classification Guide

Pharma, Chemicals & Life Sciences

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This guide explains what the tool produces, how to use the output, and what to watch for. The actual output (draft/report) is generated on the tool page.

Background

Explains drug schedule classification under the Drugs and Cosmetics Rules — Schedule H, H1, X and others, prescription and sale conditions — with the implications (labeling, sale, record-keeping) and how to determine your product schedule. Use this to know your drug schedule and its rules. Confirm the exact schedule and conditions for your product.

Purpose of this tool & output

Know your drug schedule and its rules

How the output is used

Use the guide to identify which drug schedule your product falls under and the sale, record and prescription rules that follow.

Important cautions

AI-generated guide; confirm the exact schedule and conditions for your product.

Companion documents

- CDSCO Regulatory Primer
- Drug Manufacturing Licence Guide
- Product Permission / Registration
- Regulatory Dossier Structure
- Drug Labeling Compliance
- Regulatory Strategy Planner

This is AI-assisted guidance. Verify with a qualified professional before any binding decision. Fees, authorities and processes vary by state and change over time — confirm with your state authority.