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# Import Licence & Registration

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 import licence and registration for India — the registration certificate and import licence, authorised agent, documents and CDSCO involvement — with the documents needed (manufacturer details, GMP, product info). Use this to import drugs or materials legally. Drug import is tightly regulated; engage a regulatory professional.

## Purpose of this tool & output

Import drugs and materials legally

## How the output is used

Use it to plan the registration certificate and import licence for a drug or material and to line up the Indian authorised agent and dossier before you apply.

## Who receives it

CDSCO (Central Drugs Standard Control Organisation) / DCGI; imported drugs need a registration certificate and an import licence.

## How to submit / file it

File through the CDSCO SUGAM portal with the manufacturing-site details, product dossier and an Indian authorised agent holding a wholesale licence; pay the prescribed fees.

## Fees & charges

Registration and import-licence fees as prescribed by CDSCO, per site and per product.

## Who should vet it

A CDSCO-experienced regulatory consultant and your Indian authorised agent.

## Important cautions

AI-generated guide; import of drugs is tightly regulated, engage a regulatory professional.

## 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.