

Loan Licence Agreement

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

Drafts a loan licence manufacturing agreement — a clause-by-clause draft covering scope and products, use of licence and premises, quality and GMP responsibility, materials, batch release, IP and confidentiality, liability, and term and termination — with a note on loan-licence regulatory responsibilities. Use this to manufacture under a loan licence. It has regulatory implications; have it reviewed by a lawyer and regulatory expert.

Purpose of this tool & output

Manufacture under a loan licence

How the output is used

Use the draft to set up loan-licence manufacturing with clear roles, quality responsibility and liability, then have a lawyer review it before signing.

Who receives it

The loan licensee (brand owner, you) and the manufacturer whose licence and premises are used — both sign; a copy for each party.

Fees & charges

Stamp duty per your state on the agreement; no separate government filing (the manufacturing licence itself is already held).

Who should vet it

A lawyer for the quality-responsibility, liability and termination clauses; QA on the manufacturing and batch-release responsibilities.

Important cautions

AI-generated template; loan-licence arrangements have regulatory implications, have it reviewed by a lawyer and regulatory expert.

Companion documents

- Third-Party Manufacturing Agreement
- Distribution / Stockist Agreement
- NDA Builder
- Quality Agreement Builder
- Testing / Service Agreement
- Pharma Compliance Calendar

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.