AISEVAK.IN · FREE GUIDE

Quality Agreement Builder

Pharma, Chemicals & Life Sciences

Document type: Contract / agreement · Generated 27 Aug 2026

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 quality agreement structure between a pharma company and a manufacturer or supplier — a responsibilities matrix (QC, release, deviations, changes, complaints, recall, audits), specifications, documentation and communication. Use this to define quality roles with partners. A quality agreement is a GMP expectation; have QA and legal review. Pair it with the Third-Party Manufacturing Agreement.

Purpose of this tool & output

Define quality roles with partners

How the output is used

Use the draft to define who is responsible for each quality activity between you and a manufacturing or testing partner, then have QA and a lawyer review it.

Who receives it

You and the manufacturing or testing partner — both sign; a copy for each party. It is usually annexed to the main supply/manufacturing contract.

Fees & charges

Stamp duty per your state if executed as a standalone deed; often signed as an annexure with no separate fee.

Who should vet it

QA to allocate every GMP responsibility correctly, and a lawyer for the liability and audit-rights clauses.

Important cautions

AI-generated structure; a quality agreement is a GMP expectation, have QA and legal review.

Companion documents

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

processes vary by state and change over time — confirm with your state authority.