

Manufacturing SOP Writer

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

Writes a GMP manufacturing SOP for a specific process — purpose, scope, responsibilities, step-by-step procedure, in-process checks, documentation (batch-record entries) and references. Use this to standardise a production process compliantly. For the batch record structure use the Batch Manufacturing Record (BMR) tool.

Purpose of this tool & output

GMP-compliant production procedures

How the output is used

Use the drafted SOP as the basis for a GMP-compliant production procedure, then adapt it to your equipment and process before use.

Important cautions

AI-generated SOP; align to your product, Schedule M and validated process.

Companion documents

- GMP Implementation Guide
- Batch Manufacturing Record (BMR)
- Cleaning Validation Protocol
- Process Validation Protocol
- Equipment Qualification (IQ/OQ/PQ)
- GMP Self-Audit Checklist

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.