



# Pharmaceutical Document Control Checklist

## Document Creation & Approval

Ensuring proper authorization and version control during document creation.

### Document Title

Write something...

### Document Purpose/Scope

Write something...

### Document Type (SOP, Form, Report, etc.)

- ☐ SOP
- ☐ Form
- ☐ Report
- ☐ Record
- ☐ Other

**Document Classification (Confidential, Public, etc.)**

- ☐ Public
- ☐ Confidential
- ☐ Restricted

**Date of Creation**

Enter date...

**Document Version Number**

Enter a number...

**Author Signature**

**Reviewer Signature**

**Document Review & Revision**

Verification of accuracy and completeness by designated personnel.

### Review Date

Enter date...

### Reviewer Role

- ☐ Subject Matter Expert
- ☐ Quality Assurance
- ☐ Regulatory Affairs

### Review Comments/Observations

Write something...

### Revision Number (if applicable)

Enter a number...

### Review Result

- ☐ Approved
- ☐ Requires Revision
- ☐ Rejected

### Reviewer Signature

### Approval Date (if approved)

Enter date...

## Document Distribution & Access

Controlled distribution and limited access to approved documents.

### Distribution Method

- ☐ Electronic (Controlled)
- ☐ Printed (Controlled)

### Recipients (Document Distribution List)

- ☐ QA Department
- ☐ Manufacturing Team
- ☐ Validation Team
- ☐ Regulatory Affairs

### Distribution Notes/Comments

Write something...

### Date of Distribution

Enter date...

Distribution Acknowledgement

Document Storage & Retention

Secure and compliant storage of all documents for the required retention period.

Total Number of Documents Stored

Enter a number...

Last Storage System Review Date

Enter date...

Physical Storage Location (if applicable)

 [Set My Current Location](#)



### Description of Storage Conditions (e.g., temperature, humidity)

Write something...

### Retention Period (Years)

Enter a number...

### Document Destruction Start Date (if applicable)

Enter date...

### Storage Media Type (Electronic/Physical)

☐ Electronic

☐ Physical

## Document Retrieval & Availability

Ensuring readily available access to necessary documents when required.

### Date of Last Document Retrieval

Enter date...

### Document Retrieval Purpose/Reason

Write something...

### Retrieval Method (e.g., Electronic, Paper)

☐ Electronic

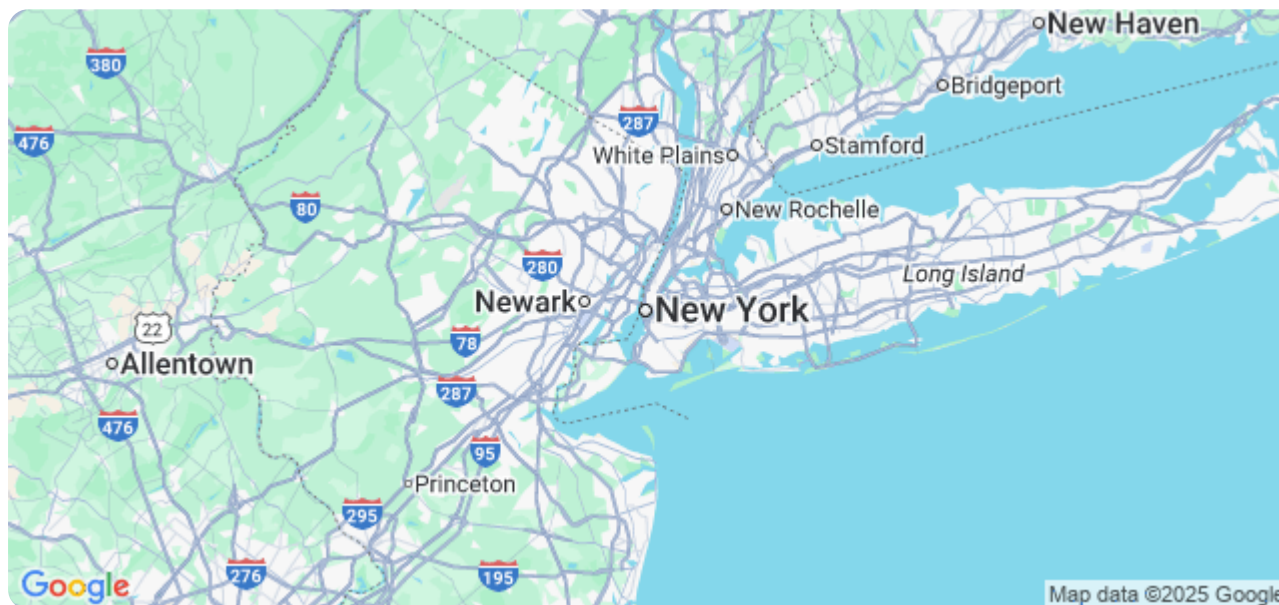
☐ Paper

### Number of Documents Retrieved

Enter a number...

### Location of Document Retrieval (e.g., Server, Archive)

[📍 Set My Current Location](#)



### Time of Retrieval

## Obsolete Document Control

Proper identification and removal of obsolete documents from circulation.

### Date of Document Identification as Obsolete

Enter date...

### Reason for Document Obsolescence

- ☐ Superseded by newer version
- ☐ Process Change
- ☐ Regulatory Change
- ☐ Other

### Description of the reason for obsolescence

Write something...

### Upload Document for Review (Optional)

 Upload File

### Document Status Post-Obsolescence

- ☐ Archived
- ☐ Removed from Active Distribution
- ☐ Scheduled for Destruction

### Reviewer Signature



### Document ID

Write something...

## Change Management (Documents)

Tracking and documenting all changes made to controlled documents.

### Change Request Date

Enter date...

### Reason for Change

Write something...

### Detailed Description of Change

Write something...

### Affected Document(s)

- ☐ SOP
- ☐ Manufacturing Record
- ☐ Validation Protocol
- ☐ Other (Specify)

### Document Revision Number (Before Change)

### Document Revision Number (After Change)

### Supporting Documentation

 Upload File

### Preparer Signature

### Reviewer Signature

## Document Master Record (DMR) Review

Verification and approval of DMRs for manufacturing processes.

### DMR Review Date

### DMR Version Number

**Review Status**

- ☐ Approved
- ☐ Rejected
- ☐ Requires Revision

**Review Comments/Observations**

Write something...

**Reviewer Signature**

**Deviation Identified?**

- ☐ Yes
- ☐ No

**Supporting Documentation (if applicable)**

 Upload File