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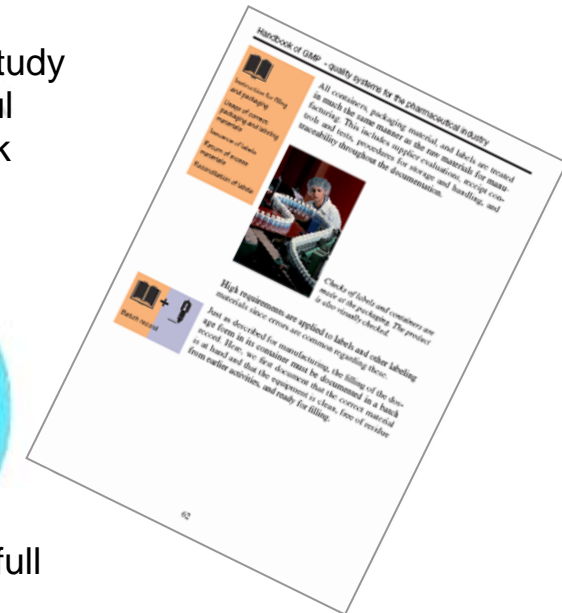
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- EU GMP Guideline Part I for medicinal products
- EU GMP Guideline Part II for manufacturing of active substances used as starting materials (corresponds to ICH Q7 "GMP for API")
- 21 CFR 210/211 "Current Good Manufacturing Practice for Finished Pharmaceuticals" (US / FDA)