

Date: 01/Dec./2022

EGYPTIAN DRUG AUTHORITY GUIDELINES ON GOOD MANUFACTURING PRACTICE OF PHARMACEUTICALS

(Under review and Technical consultation)

1. Background

This guidance document is intended to be applied in the processing of pharmaceutical drugs. It describes the current basic concepts on steps for manufacturing and controlling pharmaceutical products in order to guide manufacturers of pharmaceutical products and regulatory authorities responsible for pharmaceutical inspections on pharmaceutical facilities.

This guide to GMP shall be used as a standard to justify GMP status, and as a basis for the inspection of manufacturing facilities.

2. Consultation procedures:

The purpose of this consultation is to collect relevant comments and reviews from pharmaceutical manufacturers to ensure the development of Egyptian Drug Authority Guidelines on Good Manufacturing Practice of Pharmaceuticals in order to achieve the outmost benefit for the industry and ensure the delivering of Safe and effective pharmaceutical products.

The consultation procedure will take place in period from 01/Dec./2022 to 05/Dec./2022

Any comments should be sent to dfi.auditunit@edaegypt.gov.eg and dfi.devunit@edaegypt.gov.eg

