Change Control

MEGGLE Products:
- Lactose Monohydrate: CapsuLac® 60, FlowLac® 90, FlowLac® 90 MS, FlowLac® 100, FlowLac® 100 MS, FlowLac® 100 SD, GranuLac® 70, GranuLac® 70 MS, GranuLac® 80, GranuLac® 140, GranuLac® 140 S, GranuLac® 200, GranuLac® 200 MS, GranuLac® 200 S, GranuLac® 230, InhaLac® 70, InhaLac® 120, InhaLac® 140, InhaLac® 145, InhaLac® 150, InhaLac® 160, InhaLac® 180, InhaLac® 230, InhaLac® 251, InhaLac® 300, InhaLac® 400, InhaLac® 500, Lactose Monohydrate 200 Mesh IP, Lactose Monohydrate Impalpable, Lactose Monohydrate Low Endotoxin, PrismaLac® 40, SacheLac® 80, SorboLac® 400, SpheroLac® 100, Tablettose® 70, Tablettose® 80, Tablettose® 100, Tablettose® 100 MS
- Co-processed Excipients: Cellactose® 80, CombiLac®, MicroceLac® 100, RetaLac®, StarLac®

A Change Control procedure according to “The International Pharmaceutical Excipient Council Significant Change Guide for Pharmaceutical Excipients” published by IPEC Federation is implemented.

Quality Unit has the responsibility and authority for final approval of changes.

The procedure includes an assessment of the need to inform customers and authorities about the change.

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