EC Certificate Production Quality Assurance System: Certificate US15/842060

The management system of

Henkel Corporation

6050 West 51st Street,
Chicago, IL, 60638, United States

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

CO2 absorbent materials for use in anesthesia circuits and respiratory therapy equipment Including TM Brands - SODASORB® H Med, SODASORB® LF and PRE PAK®

For placing on the market of Class IIb or Class III devices covered by this certificate, an EC Type Examination Certificate according to Annex III is required.

This certificate is valid from 14 March 2018 until 20 February 2021
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 24 August 2018
Issue 4. Certified since 20 February 2015

Certification is based on reports numbered WW/MC 209646

Authorised by

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