

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

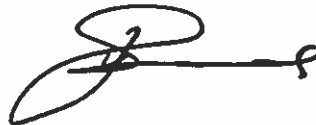
**CO2 absorbent materials for use in anesthesia circuits  
and respiratory therapy equipment Including  
TM Brands - SODASORB® H Med, SODASORB® LF and PRE PAK®**

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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