

dical Declaration of Conformity

Manufacturer CuraMedical B.V. Address Industrieweg 6B

1566 JP ASSENDELFT

The Netherlands

Product Type Oxidized Regenerated Cellulose Hemostat, Sterile, Absorbable, Single Use

Classification

Ш

(MDD, Annex IX)

GMDN code 38771

arterial bleeding when binding or other conventional methods of control are

impractical or ineffective.

Brand name CuraCel® Standard

Dye Un-dyed (natural, beige-caramel)

Insulation / Coating NA

Types and Sizes:

CC-501 50x12.5mm CC-507 50x75mm CC-538 100x100mm CC-503 25x25mm CC-535 50x350mm CC-540 100x200mm CC-505 CC-537 100x75mm 150x230mm 50x25mm CC-560

We herewith declare under our sole responsibility that the above mentioned products meet the provisions of the following EC Council Directives and Standards. The design of the products and the Quality System meets the requirements according to Annex II – Section 3 and 4 of the Council Directive 93/42/EEC of June 14th, 1993 concerning medical devices. All supporting documentations are retained under the premises of the manufacturer and the notified body.

DIRECTIVES

General applicable directives:

Medical Device Directive: COUNCIL DIRECTIVE 93/42/EEC of June 14^{th} , 1993 concerning medical devices (MDD 93/42/EEC), as last amended September 5^{th} , 2007 (Directive 2007/47/EC).

Standards:

All relevant Harmonized Standards as published in the Official Journal of the European Communities and listed in the latest version of document: QS-1530-01 List of applicable standards Curacel, are applicable to this type of product.

United States Pharmacopoeia, USP Monographs OXIDIZED REGENERATED CELLULOSE.

Notified Body MDC Medical Device Certification GmbH,

Kriegerstrasse 6, D-70191 Stuttgart, Germany

Notified Body number: 0483

Certificate no. D1169400030 Expiry Date 2024-05-26

Date CE mark was first affixed 21-03-2008

Place & Date ASSENDELFT, 18-05-2021

Signature

Name F. J. Hoogland

Position Director Marketing & Sales and Regulatory Affairs