



# Declaration of Conformity

Manufacturer CuraMedical B.V.  
Address Industrieweg 6B  
1566 JP ASSENDELFT  
The Netherlands

Product Type Oxidized Resorbable Cellulose Hemostat, Sterile, Absorbable, Single Use  
Classification III  
(MDD, Annex IX)  
GMDN code 38771  
Intended use To be used in surgical procedures to control capillary, venous and small arterial bleeding when binding or other conventional methods of control are impractical or ineffective

Brand names **CuraTamp® Standard**  
Dye Un-dyed (natural, beige-caramel)  
Insulation / Coating NA

## Types and Sizes:

|        |           |        |          |        |           |
|--------|-----------|--------|----------|--------|-----------|
| CT-101 | 50x12.5mm | CT-107 | 50x75mm  | CT-138 | 100x100mm |
| CT-103 | 25x25mm   | CT-135 | 50x350mm | CT-140 | 100x200mm |
| CT-105 | 50x25mm   | CT-137 | 100x75mm |        |           |

*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 14<sup>th</sup>, 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<sup>th</sup>, 1993 concerning medical devices (MDD 93/42/EEC), as last amended September 5<sup>th</sup>, 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/CuraTamp, are applicable to this type of product.

United States Pharmacopoeia, USP Monographs **OXIDIZED 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 10-02-2010  
Place & Date ASSENDELFT 18-05-2021

Signature

Name

F. J. Hoogland

Position

Director Marketing & Sales and Regulatory Affairs

