EU/EC Declaration of Conformity

Identification of the product: KinesioTape

Date: 22/06/2020

Object of the declaration:

Référence	Length	Width	Rolls/ box	Boxes/ carton
Z181A120002	5 m	5 cm	1	12

Name and address of the manufacturer and/or his authorised representative:
Sylamed
3 Square de Maubeuge
75009 Paris - FRANCE

This declaration of conformity is issued under the sole responsibility of the manufacturer.

The object of the declaration described above is in conformity with the relevant Community harmonisation legislation:
Directive 93/42/CEE - 14 June 1993

References to the relevant harmonised standards used or references to the specifications in relation to which conformity is declared:
European Pharmacopoeia
NF EN ISO13485:2012

Classe : I NON STERILE classification according to annexe IX

This statement leans on the technical documentation established according to:
ANNEX VII of the DIRECTIVE 93/42/EEC concerning Medical Devices, amended by Directive 2007/47/CE
Déclaration to French Ministry (ANSM) :
Ref. letter 18/09/2012

Sylvia LANDAU
Quality Manager

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S.A.R.L. au capital de 1 000 000 euros – RCS Paris B 338 933 351 – code APE 4646Z