565 Nucla Way, Unit B
Aurora, Colorado 80011 USA
Phone: (303) 366-1804
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EC Declaration of Conformity

We, the Manufacturer:

Weaver and Company
565 Nucla Way, Unit B
Aurora, Colorado 80011
United States of America
+001 800 525 2130

Declare under sole responsibility that the topical medical devices described below:

Nuprep® Skin Prep Gel

- 4oz Tube 3-pack (Item #10-30)
- 4oz Tube Kit (Item #10-30K)
- 25 g Tube Sample (Item #10-25S)
- 25 g Tube Kit (Item #10-25)
- 25 g Tube 6-pack (Item #10-61)

Ten20® Conductive Paste

- 4oz Jar 3-pack (Item #10-20-4)
- 8oz Jar 3-pack (Item #10-20-8)
- 4oz Tube 3-pack (Item #10-20-4T)
- 4oz Tube Kit (Item #10-20-4TK)
- 50 g Jar Sample (Item #10-20-2S)
- 50 g Jar 3-pack (Item #10-20-2)

Are medical devices subject to Council Directive 93/42/EEC of June 14, 1993, amended by Directive 2007/47/EC of September 5, 2007, concerning medical devices, are Class I devices according to the classification criteria of Annex IX of the directive, Rule 1;
Are non-invasive medical devices not intended as sterile devices;
Do not perform a measuring function;
Are therefore eligible for, and conform with, the conformity assessment procedure described in Annex VII of the directive; and
Meet the essential requirements of Annex I of the Council Directive.

In addition, the medical devices described above conform with Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances.

The initial date of the CE mark on Nuprep Skin Prep Gel and Ten20 Conductive Paste was June 1, 1998.

Authorized Representative:

EC REP **Emergo Europe**
Prinsessegracht 20
2514 AP The Hague
The Netherlands
+31 70 345 8570

WEAVER AND COMPANY, INC. CORPORATE SEAL 1978 COLORADO

Chris Cooper, CEO
Management with Executive Responsibility
Weaver and Company

1 August 2017
Date