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## EC Declaration of Conformity

We, the Manufacturer:

**D.O. Weaver and Company (doing business as Weaver and Company)**  
**565 Nucla Way, Unit B**  
**Aurora, Colorado 80011 USA**


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


**Nuprep<sup>®</sup> Gel**  
**Ten20<sup>®</sup> Conductive Paste**

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.

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

Authorized Representative:  
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**Molenstraat 15**  
**2513 BH The Hague**  
**The Netherlands**  
**+31 70 345 8570**

  
\_\_\_\_\_  
Management with Executive Responsibility  
D.O. Weaver and Company

  
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Date