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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® Skin Prep Gel
Ten20® 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 I, 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 Skin Prep Gel and Ten20 Conductive Paste was June 1, 1998.

Authorized Representative:

Emergo Europe
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Management with Executive Responsibility
D. O. Weaver and Company

3/3/2011
Date