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2011

DECLARATION OF CONFORMITY

Manufacturer:	Ferno-Washington, Inc.
Manufacturer's Address:	70 Weil Way Wilmington, Ohio USA
Name of Device(s):	Model 125 Kendrick Extrication Device
Class of Device(s):	I
Intended Use:	Immobilizing and Extricating Patients

In accordance with European Community Council Directive 93/42/EEC, Annex VII, Ferno-Washington Inc. declares the products that bear the CE mark conform with all the provisions of this Directive that apply to them.

Ferno-Washington Inc. products have been determined to be a Class I device according to Article 9.1 and using classification criteria from Annex IX of the Directive.

Ferno-Washington Inc. or representative will make available upon request all applicable technical documentation to allow assessment of conformity of its products in accordance with Annex VII, Sections 2 and 3.

Ferno-Washington Inc. has a systematic procedure in place to review experience gained from its devices in the post-production phase, taken appropriate corrective action based on this experience and report malfunctions to the competent authorities in accordance with Annex VII, Section 4.

Thomas E. Livingston
Director of Quality