CE Declaration of Conformity

Company: **Cascade Designs Inc.**
4000 1st Ave. So.
Seattle, WA 98134
(206)505-9500

Declares that the following device:

Device: **VARILITE® ProForm NX™ Seating System**

Model numbers: **72411, 72412, 72421, 72422, 74411, 74412, 74421, 74422, 74611, 74612, 74621, 74622, 75511, 75512, 75521, 75522, 75711, 78712, 75721, 75722, 76611, 76612, 76621, 76622, 76811, 76812, 76821, 76822, 76011, 76012, 76021, 76022, 77711, 77712, 77721, 77722, 78611, 78612, 78621, 78622, 78811, 78812, 78821, 78822, 78011, 78012, 78021, 78022, 70611, 70612, 70621, 70622, 70811, 70812, 70821, 70822, 70011, 70012, 70021, 70022**

Conforms to Directive 93/42/EEC, Official Journal of the European Communities, and EN 12182:2012, all parts applicable to Class I medical devices.

EU Representative: **Cascade Designs Limited**
Dwyer Road, Midleton, Co. Cork
Republic of Ireland
+353-21-632399

CE Certification: **February 11, 1998**

Drawn under the supervision of and approved by:

Susan Cwiertnia

November 28, 2014

Susan Cwiertnia **Date**
Acting Director of Medical
Cascade Designs, Inc.