

**Klarity Medical Products**

**EC DECLARATION OF CONFORMITY**

According to annex VII of the Council Directive 93/42/EEC (amended 2007/47/EC) concerning medical devices:

We: Klarity Medical Products  
80 Westgate Drive  
Newark, OH 43055 USA

Declare that the following non-sterile medical devices under class I (according to rule 1 of annex IX of the Council Directive 93/42/EEC)

**KLARITY thermoplastic materials for radiation therapy patient stabilization**  
**Klarity R<sup>™</sup>**  
**Klarity Green<sup>®</sup>**

fulfill the basic requirements according to annex I no. 1-14 of the Council Directive 93/42/EEC (amended 2007/47/EC)

Conformity assessment was performed according to Annex VII.

The above products were manufactured under the following quality management systems:

EN ISO 9001:2000 Certificate No.: 20832  
EN ISO 13485:2003 Certificate No.: Q2N 090349007002



Peter M. Larson  
President  
Klarity Medical Products

Newark, Ohio USA  
September 15, 2010

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