Kerr™ EC-Declaration of Conformity

Issuer's Name: Kerr Corporation

Issuer's Address: 1717 West Collins Avenue, Orange, CA 92867, USA

EU Representative: Kerr Italia S.r.l. – Via Passanti, 332 I-84018 Scafati (SA), Italy

Object(s) of the Declaration: Herculite XRV Ultra Flow
(Class IIa)

Finished Article Number	Description
35407	Refill - Herculite XRV Ultra Flow (A1)
35408	Refill - Herculite XRV Ultra Flow (A2)
35409	Refill - Herculite XRV Ultra Flow (A3)
35410	Refill - Herculite XRV Ultra Flow (A3.5)
35411	Refill - Herculite XRV Ultra Flow (A4)
35412	Refill - Herculite XRV Ultra Flow (B1)
35413	Refill - Herculite XRV Ultra Flow (B2)
35414	Refill - Herculite XRV Ultra Flow (C2)
35415	Refill - Herculite XRV Ultra Flow (D2)
35416	Refill - Herculite XRV Ultra Flow (XL1)
35417	Refill - Herculite XRV Ultra Flow (XL2)
35418	Refill - Herculite XRV Ultra Flow (Universal Opaque)
35404	Sample - Herculite XRV Ultra Flow (A2)
36174	Sample - Herculite XRV Ultra Flow (A3)

By virtue of this declaration of conformity,
Kerr Corporation,
ensures and declares
that the object(s) described above are Class IIa products according to Annex IX of Directive 93/42/EEC and are manufactured in conformity with the provisions of the Council Directive 93/42/EEC and the technical documentation referred to in Annex VII, Section 3 and Annex V as amended by 2007/47/EC,
thus, meeting the requirements of the Medical Device Directive as demonstrated by certificate CE 00847, issued by BSI, Notified Body no. 0086.

Name: Mark Dzendzel

Function: Senior Quality Assurance Manager

Signature: 

Date: 27 March 2015