Kerr™

Annex IV EU Declaration of Conformity

Manufacturer Name and Address:	Kerr Corporation also trading as Pentron Clinical 1717 West Collins Avenue Orange, California 92867 USA
Authorized Representative Name and Address:	Kerr Italia S.r.l. Via Passanti, 174, 84018 Scafati (SA) Italy
Single Registration Number (SRN)	Not available
Technical File Name/Number:	OptiView/ R111
Basic UDI-DI:	See Attachment 1
Product Tradename(s):	OptiView™
Device Identification:	See Attachment 1
Classification and Rule(s):	Class I, Rule 5
Common Standards:	Not available
Notified Body:	Not applicable
Notified Body Number:	Not applicable
Conformity Assessment Procedure & Certificate issued:	Annex IV of EU MDR 2017/745 CE Certificate: Not applicable

Declaration Statement:
This declaration of conformity is issued under the sole responsibility of Kerr Corporation. We hereby declare that the above-mentioned device(s) comply with EU MDR 2017/745.

Signed for and on behalf of Kerr Corporation:

Orange, California USA	22 October 2021	
Place	Date of Issue	Name: Mark Dzendzel Title: Director, Quality Assurance Systems

OptiView/ R111 Attachment 1 to Annex IV EU Declaration of Conformity		
REF	**Basic UDI-DI**	**Description**
5500	084139611000080A2	OptiView™ Standard Kit
5501		OptiView™ Refill
5502		OptiView™ Small Kit