Kerr™

Annex V EC Declaration of Conformity

Manufacturer Name and Address:	Kerr Corporation 1717 West Collins Ave. Orange, CA, 92867 (USA)
Authorized Representative Name and Address:	Kerr Italia S.r.l. Via Passanti 174, 84018 Scafati (SA) Italy
Technical File Name/Number:	Project #3011 and Technical File #R022
Product Tradename(s):	NX3
Device Identification:	See Attachment 1
Classification and Rule(s):	Class IIa – Rule 8
Notified Body:	BSI Group The Netherlands B.V.
Notified Body Number:	2797
Conformity Assessment Procedure & Certificate Issued:	Annex V – Production Quality Assurance CE certificate: CE 00847

Declaration Statement:
We hereby declare that the above-mentioned device(s) comply with Council Directive 93/42/EEC.

Regulatory Affairs Signature:

16 June 2021
Issue date

Mark Dzendzel
Director, Quality Assurance Systems

NX3 Project #3011 and Technical File #R022 Attachment 1 to Annex V EU Declaration of Conformity		
REF	**Description**	
33643	NX3 Automix Dual-Cure Syringe Clear	Orange
33644	NX3 Automix Dual-Cure Syringe White	Orange
33645	NX3 Automix Dual-Cure Syringe Yellow	Orange
33646	NX3 Automix Dual-Cure Syringe Bleach	Orange
33647	NX3 Automix Dual-Cure Syringe Opaque	Orange
33648	NX3 LIGHT-CURE REFILL CLEAR	Orange
33649	NX3 LIGHT-CURE REFILL WHITE	Orange
33650	NX3 LIGHT-CURE REFILL YELLOW	Orange
33651	NX3 LIGHT-CURE REFILL BLEACH	Orange
33652	NX3 LIGHT-CURE REFILL WHITE Opaque	Orange