

## Annex VII EC Declaration of Conformity

<b>Manufacturer Name and Address:</b>	Kerr Corporation also trading as Pentron Clinical 1717 West Collins Avenue Orange, California 92867 USA
<b>Authorized Representative Name and Address:</b>	Kerr Italia S.r.l., Via Passanti, 174 84018 Scafati, (SA), Italy
<b>Technical File Name/Number:</b>	Endodontic Root Canal Sealers Endo-010
<b>Product Tradename(s):</b>	Sealapex & Sealapex Xpress
<b>Device Identification:</b>	see Attachment 1
<b>Classification and Rule(s):</b>	Class IIa, Rule 8
<b>Notified Body:</b>	BSI Group The Netherlands B.V.
<b>Notified Body Number:</b>	2797
<b>Conformity Assessment Procedure &amp; Certificate Issued:</b>	Annex V CE Certificate 00847

### Declaration Statement:

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

### Regulatory Affairs Signature:

31 March 2021

**Issue date**



**Name:** Mark Dzendzel

**Title:** Director, Quality Assurance Systems

Sealapex & Sealapex Xpress - Attachment 1 to Annex VII EC Declaration of Conformity	
REF	Description
18432	Sealapex
18598	Sealapex Bulk Pack
33639	Sealapex Xpress