

# Sterimedix Ltd

Quality System Document QD 2040 Issue P  
EC Declaration of Conformity

## Declaration of Conformity for Capsule Polishing Cannulae

Ref: DOC03

30/05/2018 - Issue 4

### Council Directive 93/42/EEC Concerning Medical Devices

The undersigned declares that the product named in this document meets the applicable Council Directive provisions that apply and the CE Mark may be affixed.

<b>General Product Name</b>	Capsule Polishing Cannulae	
<b>Product Reference</b>	<b>Capsule Polishing Cannulae</b> M8603; M8604; M8604A; M8604B; <b>Silicone Tipped Polishers</b> M8605; M8605A; M8608; SD5005; SD1605; <b>Cannulae Olive Tipped Cannulae</b> M8620; M8620C; M8621; M8622; SD4815; <b>Posterior Scrapers</b> M8645;	
<b>Manufacturer</b>	Sterimedix Ltd 1 Madeley Road, North Moons Moat, Redditch Worcestershire, B98 9NB, UK	
<b>Notified Body</b>	Notified Body 0120 SGS United Kingdom Ltd 202b Worle Parkway, Weston-super-Mare, BS22 6WA	
<b>Intended Use</b>	Surgically invasive device for polishing and cleaning the residual cortex from the capsular bag. Silicone tipped polishers provide for gentle polishing and atraumatic insertion and removal. Micro etched tips are slightly roughened to facilitate gentle polishing, and are intended for transient use.	
<b>Sterile</b>	Sterile - EO. Approved by SGS United Kingdom Ltd	
<b>Measuring Function</b>	No	
<b>Conforming to Current Standards</b>	EN 556-1 ISO 594-1, 2 BS EN 980 EN 1041 EN 1707 EN ISO 10993-1, 3, 4, 5, 6, 7, 11 EN ISO 11135-1	EN ISO 11607-1, 2 EN ISO 11737-1, 2 EN ISO 13485 EN ISO 14155-1 EN ISO 14937 EN ISO 14971 EN ISO 15223-1
<b>Classification</b>	Class IIa in accordance with Annex IX Rule 6	
<b>CE Marking</b>	Notified Body and Number: Initial Date of CE Marking: Certified to: Certificate Number: Certificate Issue Date: Certificate Expiry Date:	SGS 0120 21/05/1997 93/42/EEC GB97/9964 12/08/2015 06/02/2020
<b>Assessment Route</b>	By Annex II excluding section 4 and Article 11 of the European Communities Council Directive 93/42/EEC amended by Directive 2007/47/EC, and the application of BS EN ISO 13485:2016.	

This product does not contain human blood derivatives and is not manufactured utilising tissues of animal origin.

Signed  Date 31/5/18

Name: Andrew Ellis

Position: QA/RA Manager