

# EC Certification



## EC DESIGN EXAMINATION CERTIFICATE Directive 93/42/EEC for Medical Devices, Annex II (4) OBL

We hereby declare that a design examination has been carried out on the device(s) listed hereafter following the requirements of the UK national legislation to which the undersigned is subjected, transposing Annex II Section 4 of the Directive 93/42/EEC on medical devices via an Own Branding arrangement. We certify that the design of the device(s) listed hereafter conforms with the relevant provisions of Annex II Section 4 of the aforementioned legislation, and the result entitles the organization to use the CE 0473 marking on those products.

## JRI ORTHOPAEDICS LTD

18 Churchill Way, 35A Business Park, Chapeltown, Sheffield, S35 2PY

<i>Repros® Paste</i>	0.5cc
<i>Repros® Paste</i>	1cc
<i>Repros® Paste</i>	2cc
<i>Repros® Paste</i>	2.5cc
<i>Repros® Paste</i>	3cc
<i>Repros® Paste</i>	5cc
<i>Repros® Paste – (multipack 2 x 5 cc)</i>	10cc
<i>Repros® Paste – (multipack 3 x 5 cc)</i>	15cc

\*For CE marking the class III devices covered by this certificate, an EC certificate according to Annex II (3) is also required.

**Certificate Number:** 070 E DE  
**Initial Certification Date:** 23 May 2013  
**Certificate Effective Date:** 31 October 2013  
**Certificate Expiry Date:** 22 May 2018

Brian Johnson ~ Authorised Signatory

AMTAC Certification Services Limited, Milton Keynes, UK  
This certificate is the property of AMTAC Certification Services Ltd a wholly owned subsidiary of Intertek Holdings Ltd



In the issuance of this certificate, Intertek assumes no liability to any party other than to the Client, and then only in accordance with the agreed upon Certification Agreement. This certificate's validity is subject to the organization maintaining their system in accordance with Intertek's requirements for systems certification. Validity may be confirmed via email at [certificate.validation@intertek.com](mailto:certificate.validation@intertek.com) or by scanning the code to the right with a smartphone.

This Certificate is for the exclusive use of AMTAC's client and is provided pursuant to the agreement between AMTAC and its Client. AMTAC's responsibility and liability are limited to the terms and conditions of the agreement. AMTAC assumes no liability to any party, other than to the Client in accordance with the agreement, for any loss, expense or damage occasioned by the use of this Certificate. Only the Client is authorized to permit copying or distribution of this Certificate. Any use of the AMTAC name or one of its marks for the sale or advertisement of the tested material, product or service must first be approved in writing by AMTAC.

The certificate remains the property of Intertek, to whom it must be returned upon request.

The certification is subject to the organization maintaining their system in compliance with the regulations stated in this certificate, allowing regular assessments and following the contracted requirements of the Notified Body.

AMTAC Certification Services Limited is a Notified Body according to Directive 93/42/EEC for medical devices, with identification number 0473.