

*Copy given to Dr S Zilberman*

**Specialists in Orthopaedics**

10<sup>th</sup> November 2011

File Reference 2011/011177

Attention: [REDACTED]

Head, Monitoring and Compliance Group  
Therapeutic Goods Administration  
PO Box 100, Wooden  
ACT 2606

Dear [REDACTED]

**Removal of the Eska Adapter Femoral Stem Prosthesis from the register**

**Device Name:** Eska Adapter Femoral Stem Prosthesis  
**ARTG Number – Name:** 118441 – Eska Australia – Prosthesis, internal, joint, hip, femoral component

We refer to your letter of the 25<sup>th</sup> October, file reference 2011/011177 informing us of your decision with respect to the above products listed under ARTG code 118441.

I confirm that in accordance with section 17 of your letter that Eska Australia will cease the supply of the Adapter Stem Femoral Prosthesis on or before the 26<sup>th</sup> November 2011, this being in line with the minimum period advised in this clause.

Eska Australia hereby declares that in accordance with sections 10 and 18 of your letter other medical devices are registered and are being supplied under this ARTG number.

These products are being placed on the market under the following descriptions.

1. GHE Femoral Stem Cementless Standard Offset
2. GHE Femoral Stem Cementless Lateral Offset
3. GHE/s Femoral Stem Cementless
4. GHE Femoral Stem Cementless for Extension
5. Femoral Cone Cementless - Revision

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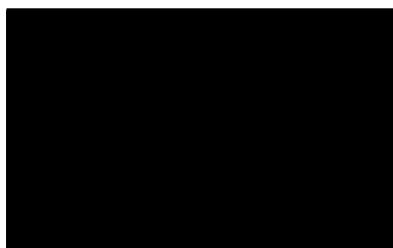
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com.au



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Can you please confirm by return that the supply of the products under the descriptions listed above remain unaffected by the removal of the Adapter Stem Femoral Prosthesis?

I have also advised the manufacturer Orthodynamics GmbH of your decision and its implications. Should you require any additional information please contact Eska Australia on 02 9684 4798 or email [customerservice@eskaustralia.com.au](mailto:customerservice@eskaustralia.com.au).



ESKA Australia Pty Ltd

cc [redacted]



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