



# Medical Device Application

## Class IIb Status : Approved

### Application Change history

#### Application Progress Date

Date received: 22/12/2009

#### Review Information

Review flag:

Auto review required: No

#### ARTG & Product ID

ARTG ID 168581

Product ID 282699

#### Application Details

Application identifier: DV-2009-DA-14567-3

Submission identifier: DA-2010-00047-7

Sponsor's own reference: Lima Orthopaedics - SMR System glenoid components

Application for: Medical Device - Included

Will you be applying for listing of this product or procedure in the Medicare Benefit Schedule (MBS)?  Yes  No

Will you be applying for listing of this product on the Prosthesis List?  Yes  No

Will you be applying for listing of this product on the Co-dependent or hybrid technology application list?  Yes  No

Cancel ARTG - product:

#### Sponsor Details

Agent name: [REDACTED]

Sponsor name: Lima Orthopaedics Australia Pty Ltd

Contact details: [REDACTED]

Contact email: [REDACTED]

### Class Details

Class: Class IIb

Intended purpose: A component of a shoulder joint prosthesis that is affixed to the glenoid cavity bone to replace or repair the articulating surface. The device may be designed to be used with or without bone cement.

### Device Product Characteristics

Is the device, or any form of the device, supplied sterile: Yes

Sterilisation Method:

Is the device intended to be invasive: Yes

Is the device, or any form of the device, intended for single use: Yes

Is the device an active device: No

Does the device contain material or ingredients of microbial origin: No

Does the device contain material or ingredients of recombinant origin: No

Does the device contain material or ingredients manufactured or formulated using a genetically modified organism: No

Does the device contain material or ingredients of Human Origin: No

Does the device contain Human Blood or its components: No

Does the device consist of: Single product only

Does the device contain material or ingredients of Animal Origin rendered non-viable: No

Animal Species:

Country of Origin:

Does any component in the procedure, kit or system contain material or ingredients of Animal Origin rendered non-viable: No

Is the device medicated: No

Is the device formulated: No

Does the product contain a medicine that is supplied separately in the Australian Market: No

Does the product contain a medical device which incorporates a medicine as an integral part and that has an action ancillary to the device: No

Does the device contain a metal on metal bearing:

I declare that this device contains only components that are medical devices which have been individually certified. No

### Manufacturer Details

Manufacturer evidence number: DV-2009-MC-13143-3 :Lima Ortho Aust - Lima Lto Italy

Manufacturer name: Lima Lto SPA (Italy)[28937]

Manufacturer address as on evidence: Via Nazionale 52 Villanova di San Daniele UDINE 33030 Italy S [ 103905]

**GMDNS Code and Description**

GMDNS code and description: Prosthesis, internal, joint, shoulder, glenoid component[36259]

**Device Category Terms**

Device category 1: Non-active implantable devices

Device category 2:

Device category 3:

**Product Details**

UPI (Unique product identifier):

Total number of devices covered:

Functional description:

**Variant List**

#	Variant type	Variant range
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**Standard Conditions**

- The inclusion of the kind of device in the ARTG is subject to compliance with all conditions placed or imposed on the ARTG entry. Refer Part 4-5, Division 2 (Conditions) of the Therapeutic Goods Act 1989 and Part 5, Division 5.2 (Conditions) of the Therapeutic Goods (Medical Devices) Regulations 2002 for relevant information.
- Breaching conditions of the inclusion related to the device of the kind may lead to suspension or cancellation of the ARTG entry; may be a criminal offence; and civil penalties may apply.

**Non Standard Conditions**

Note: A non standard conditions must not contain semi colons.

To remove, enter item #

**Declaration**

- (a) devices of the kind in question are medical devices; and
- (b) devices of that kind are intended for a specified purpose, as ascertained under The definition

of a medical device; and

(c) the kind of device is correctly classified according to the medical device classifications; and

(d) devices of that kind comply with the essential principles; and

(e) I:

(i) have available sufficient information to substantiate that compliance with the essential principles; or

(ii) have procedures in place, to ensure that such information can be obtained from the manufacturer within the period specified in the regulations; and

(f) an appropriate conformity assessment procedure has been applied to devices of that kind; and

(g) I:

(i) have available sufficient information to substantiate the application of those conformity assessment procedures; or

(ii) have procedures in place, to ensure that such information can be obtained from the manufacturer within the period specified in the regulations; and

(h) devices of that kind comply with every requirement (if any) relating to advertising applicable under the regulations; and

(i) devices of that kind do not contain substances that are prohibited imports for the purposes of the Customs Act 1901; and

(ia) devices of that kind are not to be used exclusively for one or more of the purposes specified under section 41BEA; and

(j) the information included in or with the application is complete and correct.

I understand the consequences of making a false declaration, as outlined below.

In electronically submitting this application to TGA, I hereby declare that in relation to this medical device the information given in this application and the above statements on this declaration form are current and correct.

**PLEASE NOTE:**

A false declaration will result in the device entry being removed/cancelled from the ARTG.

Signatory name of the person submitting the application.:

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History	
2/02/2010 1:00:29 PM Approved.	
Review Completed - Accepted, 2/02/2010)	

Record	Date
Fee 790	Date Paid 04/01/2010
	Date Decision 02/02/2010

Start Dates	Finish Dates	Working Days
Application Received 22/12/2009	Payment Received 04/01/2010	9
Payment Received 04/01/2010	Application Decision 02/02/2010	30
Total Working Days		39