



Place **PATIENT DETAILS** label here

**and/or**

if any patient details are not available on the hospital label please complete below

Surname: ..... Female:  Male:   
 First Name: ..... Middle Initial: .....  
 Address: .....  
 ..... Post Code: .....  
 Hospital Patient No: ..... DOB: ...../...../.....  
 Medicare No: ..... DVA No. ....  
 (If applicable)  
 Name of Hospital: ..... State: .....  
 Consultant Surgeon Code: .....

Weight (kg) ..... Height (cm) ..... ASA .....

### PLEASE COMPLETE THIS SECTION IN FULL

(If bilateral use **TWO** forms)

**OPERATION DATE** ...../...../.....

**L**  **R**

**PRIMARY**






**REVISION/RE-OPERATION**

- Osteoarthritis.....
- Post Traumatic Arthritis .....
- Rotator Cuff Arthropathy.....
- Rheumatoid Arthritis.....
- Other Inflammatory Arthritis.....
- Fracture *specify* .....
- Osteonecrosis/Avascular Necrosis.....
- Dislocation .....
- Instability.....
- Tumour *specify* .....
- Other *specify*.....

- (includes removal, exchange or addition of one or more components)
- Loosening .....
  - Lysis .....
  - Infection.....
  - Implant Breakage *specify* .....
  - Instability .....
  - Dislocation.....
  - Component Dissociation.....
  - Fracture *specify* .....
  - Other *specify* .....

### GLENOID MORPHOLOGY\*

**CT Scan** Yes  No

**A1**   **B1**   **C**    
**A2**   **B2**   **Tick one diagram box only**

\* Reprinted from The Journal of Arthroplasty, Vol 14(6), G Walch, R Badet, A Boulahia, & A Khoury, Morphologic study of the Glenoid in primary glenohumeral osteoarthritis. Figure 2: Different morphologic types of the glenoid in primary genohumeral osteoarthritis, pg. 757, (1999), with permission from Elsevier.

### ROTATOR CUFF

- GRADE 0  Normal Tendon
- GRADE I  Tendinopathy / Partial Thickness
- GRADE 11A  Full Thickness (<1CM)
- GRADE IIB  Full Thickness (>1CM)
- GRADE III  Large (>2 tendons / decentred head)

**Rotator Cuff Repair Undertaken**

Yes  No

### GLENOID COMPONENTS

(Mark relevant box, place company labels on coloured areas or complete details by hand)

Company .....  
 Prosthesis Name .....  
 Cat/Ref No. ....  
 Lot No. ....

Company .....  
 Prosthesis Name .....  
 Cat/Ref No. ....  
 Lot No. ....

Company .....  
 Prosthesis Name .....  
 Cat/Ref No. ....  
 Lot No. ....



### CEMENT

(Use company label or complete details: if more than one mix is used, use only 1 label)

Humeral **NO**  **YES**

Cement Name .....

Glenoid **NO**  **YES**

Cement Name .....

### HUMERAL COMPONENTS

(Mark relevant box, place company labels on coloured areas or complete details by hand)

Company .....

Prosthesis Name .....

Cat/Ref No. ....

Lot No. ....

Company .....

Prosthesis Name .....

Cat/Ref No. ....

Lot No. ....

Company .....

Prosthesis Name .....

Cat/Ref No. ....

Lot No. ....

Company .....

Prosthesis Name .....

Cat/Ref No. ....

Lot No. ....

### ADDITIONAL COMPONENTS

Company .....

Prosthesis Name .....

Cat/Ref No. ....

Lot No. ....

### RE-OPERATION

This is an additional operation on a joint that has previously received a prosthesis. A re-operation however, is not a revision. (i.e) IT DOES NOT involve removal, exchange or addition of one or more components. It is usually an isolated soft tissue and/or bony procedure.

Re-operation performed .....

Reason for re-operation .....

Comments (If required) .....

### TECHNOLOGY ASSISTED

*tick all that apply*

Computer Navigated ..... **NO**  **YES**

*System used:* .....

Image Derived Instrumentation (IDI)..... **NO**  **YES**

*System used:* .....

Robotic Assisted ..... **NO**  **YES**

*System used:* .....

Other ..... **NO**  **YES**

*System used:* .....

Affix label here if available:

### ADDITIONAL COMMENTS (or Extra Labels)

**ALL SECTIONS of this form MUST be COMPLETED**