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Rationale of the AEQUALIS™ Press-Fit Prosthesis

Glenohumeral arthritis is characterized by persistent pain, loss of glenohumeral motion and loss of glenohumeral joint space.

Prosthetic replacement of the glenohumeral joint seeks to relieve pain and restore normal kinematics and biomechanics.

Due to the relatively fixed geometry of first and second generation shoulder prosthesis, the possibilities to reproduce the normal kinematics of the glenohumeral joint are limited and often result in abnormal or reverse scapulo-humeral rhythm.

By anatomically reconstructing the proximal humerus, normal kinematics and biomechanics can be restored allowing for normal scapulo-humeral rhythm. Due to the highly variable anatomy of the proximal humerus, it is necessary to utilize a prosthesis, which takes into account the normal spherical shape of the humeral head along with the variable inclination, retroversion and combined medial and posterior offsets of the humeral head replacement in order to adapt the prosthesis to the individual patient’s anatomy.

The AEQUALIS™ prosthesis is the first third generation shoulder prosthesis. It is not only modular without the gap between the collar and the head, but also adaptable to allow restoration of the patient’s healthy anatomy and maintain the normal superior profile*.

The AEQUALIS™ Press-Fit Shoulder Prosthesis is specifically designed for long term stability of the humeral component. Its stepped surface geometry maximizes bone support for an optimal primary fixation and secondary fixation. The unique instrumentation allows compacting of the metaphyseal bone to give added support to the prosthesis and to prevent distal migration. Compaction grafting along the calcar increases the press-fit helping to avoid postoperative subsidence.

The stem is available in five diameters. 7, 9, 11, 13 and 15 mm. Four neck angles of 125, 130, 135 and 140 degrees allow the humeral implant to be better adapted to the humeral resection surface.

Ten articular heads are available, corresponding to the anatomical variations encountered in practice. Diameter and thickness are related, each corresponding to the anatomical variations.

For the largest three sizes, 50, 52 and 54 millimeters diameters, two thicknesses are available.

It is possible to reproduce the combined offset of the articular surface, medial and posterior offsets, by means of an original eccentric dial system.

* Adaptability and modularity of shoulder prosthesis. P. Boileau and Gilles Walch
The AEQUALIS™ Press-Fit Prosthesis
Designed to reproduce the anatomy.

- Unique stair-step metaphyseal geometry for strong primary and long term fixation
- Grit blasted surface
- Widebody metaphyseal segment
- Adaptable humeral head
- Cobalt chromium alloy
- Titanium polished stem: 7, 9, 11, 13, 15 ø mm.
- Eccentric dial heads system
- Variable inclination neck
Surgical Technique

1. Preoperative planning
   • Four plain shoulder A-P x-rays: neutral, internal rotation, external rotation and axillary.
   • CT Scan for the most accurate assessment of glenoid version.
   • Arthrography with contrast to confirm integrity of the rotator cuff if there is a clinical question about the cuff integrity.

2. Patient positioning
   General anesthesia modified beach chair position (Figure 1).

Figure 1: The whole arm is draped free and prepared under sterile conditions.
3. Delto-pectoral approach

**An incision** is made from the coracoid along the Deltopectoral groove, lateral to the axillary fold to avoid a postoperative contracture (Figure 2).

The Delto-pectoral interval is developed (Figure 3).

Figure 2: The incision along the coracoid and Deltopectoral groove.

Figure 3: As necessary, a partial release of the anterior deltoid insertion facilitates deltoid retraction.
The upper border of the subscapularis is identified. The clavipectoral fascia is released up to the coracoid between the coracoacromial ligament and the conjoined tendon of the coracobrachialis and short head of the biceps brachii muscles (Figure 4).

Inspect the bicipital groove to remove any osteophytes, loose bodies or excessive biceps tenosynovian. Move the biceps laterally (on the left) to check the groove.

Figure 4: Arm in abduction, rotated externally with an angled retractor placed above the coracoid process.

Figure 5: Arm abducted and internally rotated. The long head of the biceps is exposed. The pectoralis insertion occasionally needs to be incised 1 or 2 cm.
Identification of the Axillary Nerve (Figure 6)

**Subscapularis Tendon Release**

- **The upper limit of the subscapularis tendon**, which is often covered by an extension of the subcoracoid serous bursa, lies immediately below the tip of the coracoid process (Figure 7).

- **The inferior border** is defined by the anterior circumflex vessels.
A modified Mason-Allen interlocking suture is placed at the top corner of the subscapularis adjacent to the rotator interval (Figure 8).

After opening the rotator interval, the anterior subscapularis is incised 1 cm medial to its insertion on the lesser tuberosity. Then, with a progressive rotation of the arm, the inferior capsule is released up to the posterior aspect of the humeral head (Figure 9). The release of the upper pectoralis and latissimus dorsi tendons facilitates the exposure and release of the inferior glenohumeral capsule.

Superior arthrotomy: the coraco-humeral ligament is preserved.

The inferior release is a key step for adequate glenoid release.
An anterior capsulotomy is performed with a release of the middle and inferior glenohumeral ligament to mobilize the muscle and produce a long flap of subscapularis which allows tension-free reinsertion following the procedure, regardless of the position of the arm (Figure 10).

With these adequate releases, the humeral head is dislocated into the delto-pectoral interval by abduction of the arm progressive external rotation extension. In cases of severe restriction of external rotation (0° or less), it is recommended to release more of the pectoralis insertion.

4. Humeral head osteotomy

All osteophytes are carefully trimmed from the humeral head. It is recommended to use a curved osteotome to separate the interval between the osteophyte and the cortical bone.

The osteotome is placed inferiorly and directed upwards. The goal is to identify the exact capsular insertion at the anatomic neck.

If additional exposure is necessary to visualize the superior articular margin, a curved awl or sharp Hohmann is placed under the long head of the biceps and through the supraspinatus attachment to retract both. The cut is then marked with a bovie.
Superiorly and anteriorly, the anatomical neck contains the tendon insertions of the rotator cuff (supraspinatus and subscapularis), and inferiorly it is entirely continuous with the cartilage of the head and inferior cortical surface of the humerus. Posteriorly however, there is a 6 to 8 mm area which does not contain cartilage or tendon insertions; the cut should be made through the rim of the cartilage in a manner to preserve the posterior nonarticular bare area.

• The humeral head may be cut freehand (Figure 11):

• Or with a cutting guide using an oscillating saw along the limit of the anatomical neck (Figure 12 & 13):

Figure 11: The cutting guide is positioned around the anatomical neck. The two fixation hooks are securely fixed around the neck by first tightening the adjustment knob 1 followed by knob 2. Firm three point fixation is then achieved by inserting the 3 mm guide pin through the guide into the humeral head 3.

Figure 13: The oscillating saw is then used flat on the cutting guide.
5. Reaming the humeral shaft

Accurate visualization of the plane through which the humerus is cut (Figure 14) allows the hinge point to be located. Typically, the entrance point into the humeral canal is 3 mm medial from this point (to prevent damage to the greater tuberosity). The hinge point is defined as the intersection of the proximal metaphyseal humeral axis and the highest point of the cut described previously (Figure 15).

The arm is placed in extreme external rotation, with the elbow towards the body utilizing careful progressive reaming to avoid the risk of a spiral fracture.

In case the cut is too superior, the re-cutting guide may be used to remove an additional 2 mm.

1. Apply the guide on the humerus as shown.
2. Insert the pins.
3. Remove the cutting guide and rotate 180°
4. Reposition the guide on the pins.
5. Make the recut with the oscillating saw placed flat on the guide.

Figure 14

Figure 15
The humeral shaft is progressively reamed using reamers of increasing diameter from 7, 9, 11, 13 to 15 mm (Figure 16).

Reaming is completed once initial contact is made with the cortical bone of the diaphysis.

6. Choice of humeral inclination and retroversion

Once reaming of the humeral shaft has been completed, the final reamer used is left in the humerus. The inclination is then measured using the inclination guide (Figure 17). The inclination guide is then slid downwards and seated in the slots along the reamer.
The inclination is then assessed by reading the graded scale on the guide (template) (Figure 18). There are 4 possible inclinations, 125°, 130°, 135° and 140°.

When the angle lies between two values, the lower should be selected for the prosthesis. The mobile plate must sit flat on the anatomical cut. It may be necessary to slightly reorient the reamer using the handle to obtain optimal positioning of the inclination guide.

Humeral retroversion is marked with the inclination guide by making a groove into the greater tuberosity with the retroversion osteotome (Figure 19). This groove will serve to orient the compactor fin.

Remove the reamer and the inclination guide. Using a small rongeur, notch the cortical bone at the level of the groove to allow passage of the compactor fin (Figure 20).
7. Metaphyseal preparation

The monobloc compactors are used to prepare the metaphyseal bone (Figure 20).

Start with the compactor of the smallest size and progress sequentially to the desired size. Retroversion is observed by aligning the fin of the compactor with the slot created by the osteotome described previously. Compaction is completed when metaphyseal stability is obtained.

It is important to avoid varus prosthetic placement. This is accomplished by positioning the compactor fin as far laterally as possible into the greater tuberosity.

The compactor should be seated to the neck angle measured previously with the inclination guide. For example: for a 135° angle, stop when its ridge is reached (Figure 21).
8. Trial humeral implant positioning and choice of the humeral head offset

The humeral trial stem and trial neck are assembled. The trial neck chosen corresponds to the inclination determined with the inclination guide. The trial neck is slid onto the rail of the trial humeral stem (Figure 22). The system is secured by tightening the fixation screw with the 3.5 mm hexagonal screwdriver (Figure 23).

The trial stem-neck assembly is introduced into the prepared humeral shaft using the T-handle, taking care to observe the correct orientation of the fin (Figure 24). The T-handle is then removed.
It may be necessary to use the trial stem impactor to firmly seat the collar of the trial stem on the cut (Figure 25).

If the humeral trial prosthesis fits too low on the cutting surface or if there is insufficient bony support, it is recommended to either use an impaction grafting technique with bone from the humeral head or cement the prosthesis to the proper height.

In case of the necessity to cement the AEQUALIS™ stem, it is recommended that the same trial stem diameter as the optimal compactor be used.

If the trial stem of the chosen size is unstable it is recommended to cement.

Use the broach of the smaller diameter (7 mm) and progress sequentially to the desired size (Figure 26).

After positioning the trial head in an anatomic position, the entire trial prosthesis is then removed. Cement is injected into the canal after diaphyseal obturation and drying.

The definitive humeral implant is positioned and then impacted taking care to align the prosthetic fin with its slot in the tuberosity.
9. Determination of the trial head size

- Caliper or clamshell measurement of the diameter of the resected head (Figure 27).

- Directly on the trial head template (Figure 28).

Figure 27: 2 thicknesses are available for a 50, 52, or 54 mm diameter head.
The next step is to reproduce the articular surface offset using the eccentric index system.
The trial head is held with the trial head clamp and placed over the neck trunion (Figure 29A).
The head is then rotated until the ideal coverage of the humeral neck cut is obtained (Figure 29B).

Figure 29A: Posterior offset is preserved by choosing the indexed position which allows perfect coverage of the humeral surface cut.

Figure 29B
The entire trial prosthesis is then removed using the extractor-hammer. The posterior face of each trial head is marked from 1 to 8, corresponding to 8 possible index positions. The appropriate figure is read from the mark on the superior pole of the neck, to give the selected anatomical index (Figure 30).

IMPORTANT NOTE:
Glenoid replacement must be performed at this point, if necessary. Please refer to the AEQUALIS™ Glenoid Surgical Technique.

Figure 30: Direct reading of anatomical index after removal of trial prosthesis (in this case index n° 4).
10. Humeral implant assembly and positioning

After removing the trial humeral prosthesis, the final stem and head are chosen using the same parameters for diameter and inclination of the stem size and offset of the head.

**The head is positioned on the stem,** aligning the offset number with its position, using the fin of the implant stem as a reference (Figure 31).

**IMPORTANT NOTE:**
The head should be assembled to the implant stem with clean gloves in a dry environment.

Should a humeral head ever have to be removed, position the Humeral Head Removal instrument below the inferior part of the head and firmly tap with the mallet to loosen the head.
The head is impacted onto the stem using the impaction support (Figure 32) with the head impactor and the mallet. Take care not to damage the articulating surface of the implant head.

At this stage, the long head of the biceps is cut from the superior glenoid and tenodesed into the groove. Three subscapularis sutures are preplaced through the lateral subscapularis and lesser tuberosity (Figure 33).

The prosthesis is then inserted. Proper orientation of the prosthesis during insertion is assured by positioning the fin into the groove prepared with the retroversion osteotome (Figure 34).

The prosthesis has to be impacted carefully into the medullary canal to avoid metaphyseal fracture.

**Note:** In case of impingement of the stem in the medullary canal, the stem should be removed and a stem of a smaller diameter should be cemented.

### 11. Reduction of the glenohumeral joint and closure

Following reduction of the glenohumeral joint, the stability and mobility of the shoulder are tested. The joint is closed by reinsertion of the subscapularis to the coracohumeral ligament, and to the subscapular remnant, allowing slight slippage of the subscapularis upwards. The wound is closed in layers over an aspiration drain.

**Postoperatively** the arm is immobilized in a simple sling.
Instruments

50 cm Ruler
Ref. MWA123

Awl
Ref. MWA024

Humeral Cutting Guide
Ref. MWB065

3 mm Guide Pins
Ref. MJU051

2 mm Humeral Re-Cutting Guide
Ref. MWB066

Cylindrical Reamers

| Ø 7    | Ref. MWB059 |
| Ø 9    | Ref. MWB060 |
| Ø 11   | Ref. MWB061 |
| Ø 13   | Ref. MWB062 |
| Ø 15   | Ref. MWB063 |

Inclination Guide
Ref. MWB058

Retroversion Osteotome
Ref. MWA101

Monobloc Compactors

| Ø 7    | Ref. MWB050 |
| Ø 9    | Ref. MWB051 |
| Ø 11   | Ref. MWB052 |
| Ø 13   | Ref. MWB053 |
| Ø 15   | Ref. MWB054 |
Instruments

Trial Stems

| Ø 7   | Ref. MWB020 |
| Ø 9   | Ref. MWB021 |
| Ø 11  | Ref. MWB022 |
| Ø 13  | Ref. MWB023 |
| Ø 15  | Ref. MWB024 |

Trial Necks

| 125°  | Ref. MWB016 |
| 130°  | Ref. MWB017 |
| 135°  | Ref. MWB018 |
| 140°  | Ref. MWB019 |

Hexagonal Screwdriver 4.5 mm
Ref. MWB012

T. handle Trial Stem Holder
Ref. MWA106

Trial Stem Impactor
Ref. MWA109

Trial Heads

| 37 x 13.5 mm | Ref. MWA237 |
| 39 x 14 mm   | Ref. MWA239 |
| 41 x 15 mm   | Ref. MWA241 |
| 43 x 16 mm   | Ref. MWA243 |
| 46 x 17 mm   | Ref. MWA246 |
| 48 x 18 mm   | Ref. MWA248 |
| 50 x 16 mm   | Ref. MWA250 |
| 50 x 19 mm   | Ref. MWA251 |
| 52 x 19 mm   | Ref. MWA252 |
| 52 x 23 mm   | Ref. MWA253 |

Trial Head Template

Caliper
Ref. MWA102

Trial Head Clamp
Ref. MWA103
Instruments

Humeral Broaches
(for cemented use only)

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Humeral Broach Handle
(for cemented use only)
Ref. MWB015

Humeral Prosthesis Impactor
Ref. MWA108

Impaction Support
Ref. MWB013

Mallet
Ref. MWA122

Humeral Cut Protectors

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Extractor Hammer
Ref. MWA118

Head Removal Instrument
Ref. MWB067
**Implants**

**Heads**

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**Humeral Stems**

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**Extended Sizes** (available upon request only)

**Heads**

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The length is given from the top of the fin to the distal end of the stem.
Proper surgical procedures and techniques are the responsibility of the medical professional. This material is furnished for information purposes only. Each surgeon must evaluate the appropriateness of the material based on his or her personal medical training and experience. Prior to use of any Tornier implant system, the surgeon should refer to the product package insert for complete warnings, precautions, indications, contraindications, and adverse effects. Package inserts are also available by contacting Wright.

Contact information can be found in this document and the package insert.

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