## Sidus™ Stem-Free Shoulder

Surgical Technique





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#### **Enhanced Simplicity**

### Bone-preserving with a secure fixation

The *Sidus* Stem-Free Shoulder arthroplasty system offers an anatomical shoulder joint reconstruction without procedural complexity.

Consideration of the long term care of the patient is essential. The *Sidus* Stem-Free Shoulder "Humeral Anchor", by virtue of its minimal profile, enables future options for revision or upgrade to an inverse/reverse solution, while also eliminating the potential for many of the intra-operative problems reported for traditional stemmed designs.

The position of the Humeral Anchor in the *Sidus* Stem-Free system is entirely independent of the location of the humeral canal, allowing for an anatomically driven reconstruction, and a straightforward strategy for challenging pathologies.

The Sidus Stem-Free Shoulder arthroplasty system uses a straightforward timesaving surgical technique that enables placement of the device in just a few steps — saving valuable OR time (versus traditional solutions) without compromising functional outcomes.



#### **Sidus Stem-Free Shoulder Humeral Heads**

- Proven CoCr alloy (CoCr28Mo6) *Protasul*®-21WF material construction, mirror finish
- Head sizes and head dimensions derived from the human anatomy for the best possible replacement
- Humeral Heads in 11 sizes to cover the resected humeral surface and optimally position the humeral head replacement with respect to the greater tuberosity.
- Proven female taper connection
- Compatible with *Anatomical Shoulder™* Glenoids



#### **Sidus Stem-Free Shoulder Humeral Anchor**



Four windows for free view of humerus and facilitating revision

Proven male taper connection with the possibility to use a wide range of additional head dimensions from the proven *Bigliani/Flatow®* Shoulder System



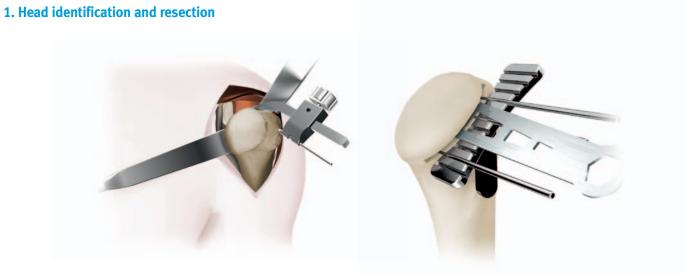
Anti lever-out surfaces on each fin designed to resist shear loads

Four open-fin press-fit design delivering protected anchoring and rotational stability while allowing for bone throughgrowth

Rough blasted surface structure fixation to promote bone on-growth, as well as to increase the friction between the implant and bone and thereby improve primary stability.

- Proven osteophilic Titanium Alloy (TiAl6V4), Protasul-64WF construction promotes long lasting osseointegration
- 3 Humeral Anchors diameters, 24, 28 and 32mm for an optimal fit
- 4 thin open star flange anchors designed to preserve bone
- One piece design for the minimization of disassociation risk
- Osteotome slots to minimize impact on bone during revision surgery
- Compatible with the *Bigliani/Flatow* Standard Humeral Heads, therefore compatibility with the proven *Bigliani/Flatow* and *Trabecular Metal*™ Glenoids is possible

### **Key Surgical Steps**

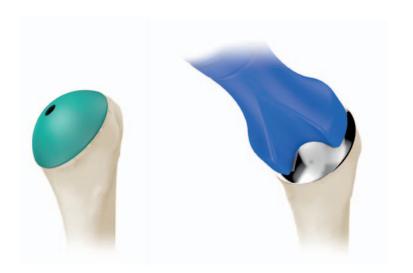


#### 2. Head size determination and positioning





#### 4. Head preparation and implantation





### **Key Revision and Intra-operative Correction Surgical Steps**

#### 1. Head removal





#### **Indications and Contraindications**

#### **General Information**

The *Sidus* Stem-Free Shoulder is indicated for uncemented use in hemi or total shoulder arthroplasty, replacing the shoulder joint of patients suffering severe pain or disability. Patient conditions of non-inflammatory degenerative joint disease (NIDJD), such as avascular necrosis and osteoarthritis and inflammatory joint disease (IJD), such as rheumatoid arthritis (conditions consequent to earlier operations), provided there is an adequate bone stock to support the fixation of the implant.

#### **Indications**

- Osteoarthritis
- Posttraumatic arthrosis
- Rheumatoid arthritis without humeral metaphyseal defects
- Focal Avascular Necrosis of the Humeral Head
- Previous surgeries of the Shoulder that do not compromise the fixation

#### **Contraindications**

- Soft or inadequate humeral bone (including Osteoporosis and extensive Avascular Necrosis or Rheumatoid arthritis) leading to poor implant fixation
- Metaphyseal bony defects (including large cysts)
- Post-traumatic tuberosity nonunion
- Signs of infection
- Irreparable cuff tear
- Revision from a failed stemmed prosthesis
- Charcot's shoulder (neuroarthropathy)

#### **Preoperative Templating**

Preoperative evaluation of the Humerus using the *Sidus* Stem-Free Shoulder templates helps determine the size of the prosthesis and level of the head resection. The goal is to make a resection that matches the anatomy of the patient.







X-Ray templates: Sidus Stem-Free Shoulder Humeral Anchor, REF 06.02281.000 Sidus Stem-Free Shoulder Humeral Head, REF 06.02282.000

#### **Patient Positioning and Exposure**

#### **Patient Positioning**

The patient should be placed in a beach chair on the edge of the operating table (Fig. 1).

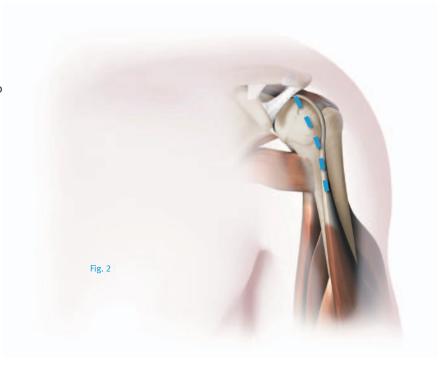
The involved shoulder should extend laterally over the edge of the table so the arm can be brought into full extension and adduction. An armrest is optional.

The upper part of the operating table has to be open on the homolateral side to allow shoulder extension (shoulder table).



#### **Initial Incision**

The incision should start in front of the ac joint 1 to 2cm lateral from the tip of coracoid running straight downwards to the humeral delta insertion (Fig. 2).



# 1. Humeral Head Identification, Preparation and Resection

With the humeral head fully exposed remove any unwanted osteophytes to restore the Humerus to near native anatomy. The humeral head should be resected exactly at the level of the anatomical neck.

Two humeral head resection techniques are possible with *Sidus* Stem-Free Shoulder Instrumentation: a freehand cut and an alignment guide "Supraspinatus" resection technique (Fig. 3).

In the superior and anterior superior aspects, the anatomical neck corresponds to the insertions of the tendons of the cuff supraspinatus and uppermost section of the subscapularis.

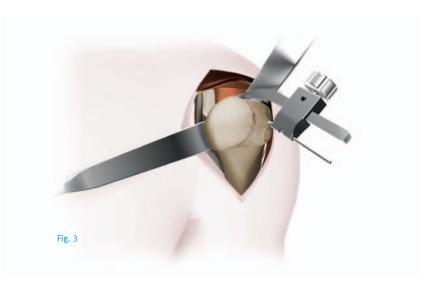
In the inferior aspect, there is a smooth transition between the cartilage of the head and the cortical bone of the Humerus.

In the posterior aspect, in the region of the infraspinatus and teres minor, is the sulcus, which is a groove of 6 to 8 mm in length, without cartilage or attached tendons. The resection must start exactly on the cartilage.

**Note:** Do not resect the cartilage-free area.

# Positioning of the Alignment Guide "Supraspinatus"

The superior part of the cut should be at the medial border of the insertion of the supraspinatus tendon #2 (Fig. 5). The Kirschner Wire (K-Wire) should exit at the posterior edge of the cartilage medial to the bare area #1 (Fig. 4). After defining the position, the displaceable part of the Alignment Guide "Supraspinatus" can be fixed with the screw #3 (Fig. 5).

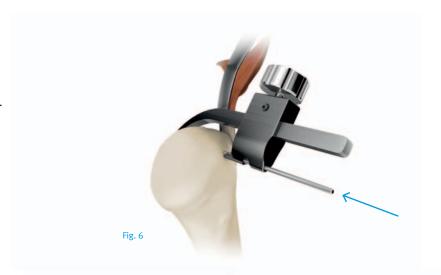


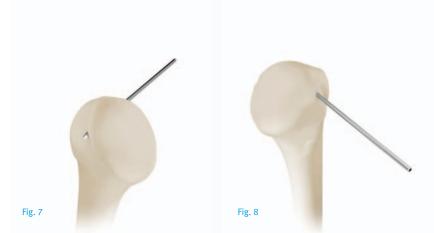




Place now the K-Wire 2x100 mm into the Alignment Guide "Supraspinatus" (Fig. 6).

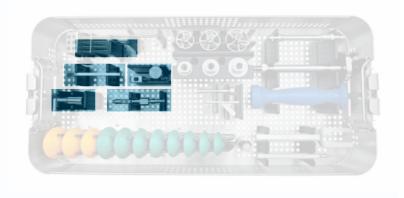
Use the Pin Retractor 2 mm to place the K-Wire.



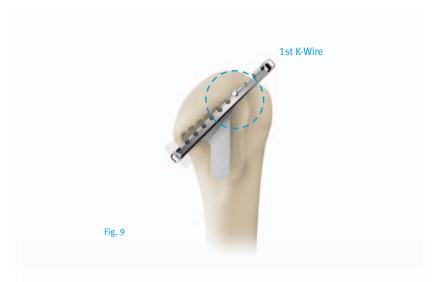


Remove the Alignment Guide "Supraspinatus".

The first K-Wire 2x100 mm, the Supraspinatus-Pin, defines the retroversion (Fig. 7 and Fig. 8).



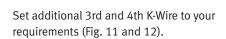
The "Resection Guide" is mounted over the 1st K-Wire (Fig. 9), the "Supraspinatus-Pin" < which identifies the retroversion >.



#### **Inclination angle identification**

Align the position of the 2nd K-Wire by identifying the correct inclination angle. Due to patient requirements, there are 2 mm K-Wires in 100 mm and 70 mm lengths.

The "Supraspinatus attachment" will additionally guide you laterally. Use the Pin Retractor 2 mm to place the K-Wire.



Use the Pin Retractor 2 mm to place the K-Wire.



Fig. 10



Fig. 11

**Note:** If you have any problems to remove and unlock the Pin Retractor 2 mm, use the "two hex key" from the Head Distractor.



Resect the Humeral Head, guided by Resection Guide and the K-Wires (Fig. 13 and 14).

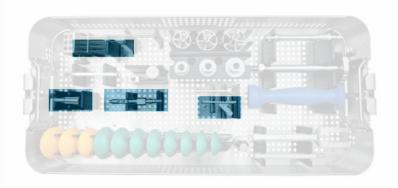
**Note:** Do not use a saw blade with downward facing teeth.



Remove the Resection Guide and the K-Wires, clean the resection area. Use again the Pin Retractor 2 mm.

**Note:** For TSA (Total Shoulder Arthroplasty) please continue on page 28.





# 2. Head Size Determination and Positioning

Choose a Trial Head size (38–52) that covers the resected humeral osteotomy (Fig. 15).

Table 1 lists the range of *Sidus* Stem-Free Shoulder Humeral Heads and Humeral Anchors with the optimal combination between head and anchor (Tab. 1).



Sidus Stem-Free Shoulder <b>HEAD</b>	<b>Diameter</b> Height	<b>38</b> 13	<b>40</b> 14	<b>42</b> 15	<b>44</b> 16	<b>44</b> 16	<b>46</b> 16	<b>48</b> 17/20	<b>50</b> 18/21	<b>52</b> 19/23
Sidus Stem-Free Shoulder ANCHOR		s				M			L	

Tab. 1 \* For compatibility between Bigliani/Flatow Standard Humeral Heads and the Humeral Anchor please refer to page 34

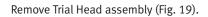
Insert the Central Pin Positioner for Trial Head through the selected Trial Head (Fig. 16).



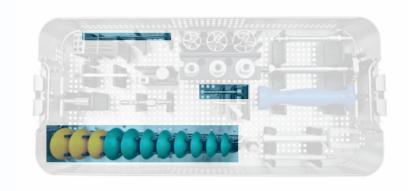
Place the Trial Head assembly back to the osteotomy (Fig. 17).



Insert the Central Pin 3.2 mm (Fig. 18).







# 3. Anchor Size Determination, Preparation and Implantation

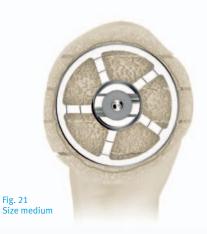
Determine first the size of the Anchor by using one of the Anchor sizer and countersink S/M/L. Place them over the 3.2 mm Central Pin.



To determine the size, the inner open ring should cover most of the cancellous bone (blue) without involving the cortex, the outer ring (orange) should be centric to the resected humeral surface (green) (Fig. 21a).

In this example, Fig. 21 with size medium Anchor sizer, would be the ideal solution.

Note: Undersizing of the anchor must be avoided as this can contribute to unfavourable conditions for long term fixation.



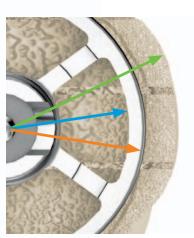
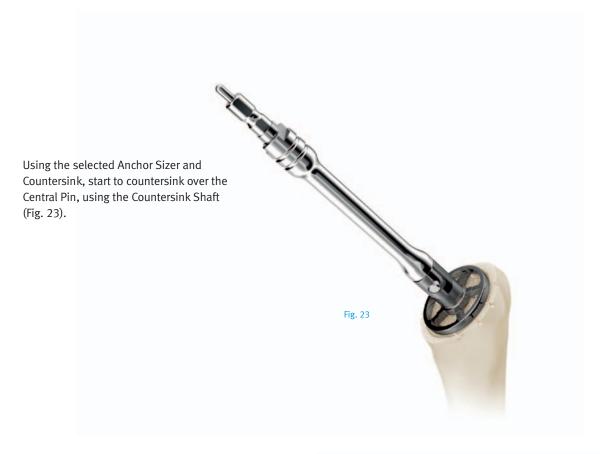


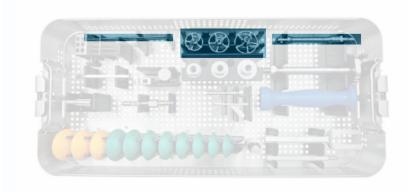
Fig. 21a



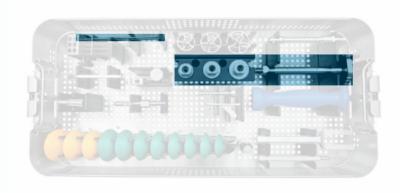


The counter bore depth of cut is limited and the collar counter bore will bottom out (Fig. 24).









Stop impacting when the Humerus Puncher sits flush within the countersunk area (Fig. 28 and 28a).

With controlled force, use a hammer to remove the Humeral Puncher over the Central Pin (Fig. 28b).

**Note:** If you have difficulty to remove the Humerus Puncher from the Impactor/ Extractor, use the two hex key from the Head Distractor.

The Humerus is now finally prepared for the implantation of the Humeral Anchor (Fig. 29).

**Note:** Now, check the proximal Humerus bone quality.

To achieve a good outcome, the patient must have adequate bone stock to support the fixation of the implant. If you have any doubt about bone quality affecting the stable fixation of the anchor, you must use a stemmed shoulder prosthesis. The *Anatomical Shoulder* System may offer the appropriate solution.

Align the Humeral Anchor S/M/L over the 3.2 mm Central Pin. Press, by hand, the Humeral Anchor into the previously prepared cuttings (Fig. 30).







Start impaction, by using the Anchor and Head Pusher (Fig. 31).

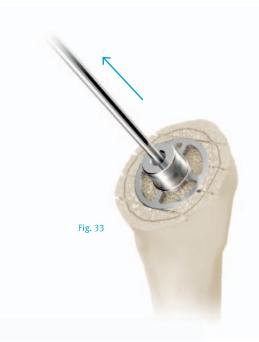


Due to the open areas of the Anchor and Head Pusher you are at every time aligned with the current position of your Humeral Anchor (Fig. 32).

The Humeral Anchor S/M/L should be flush with the countersunk area after impacting.

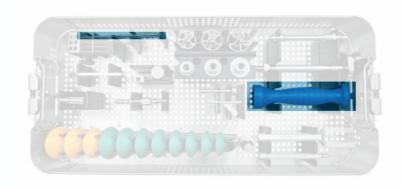


When the Humeral Anchor is flush with the countersunk area, remove the Central Pin (Fig. 33).



The Humeral Anchor is now in place (Fig. 34).





# 4. Head Preparation and Implantation

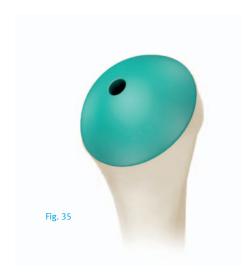
Place the Trial Head on the Humeral Anchor and conduct the trial reduction (Fig. 35).

Now use a burr or a rongeur to remove any residual osteophytes or any bone extending beyond the periphery of the humeral head. This relieves impingement with the Scapula.

It is important to balance soft tissue tension with the appropriate trial humeral head in place. It should be possible to fully internally rotate the arm across the chest so that the hand of the involved shoulder easily rests on top of the opposite shoulder, without elevating the involved shoulder off the table. It should also be possible to externally rotate the arm and still reapproximate the subscapularis tendons to the cut surface of the neck of the humerus. If the fit of the Humeral Head is so tight that the functional internal or external rotation or posterior subluxation cannot be obtained, then further soft tissue release posteriorly is required.

**Note:** All taper surfaces must be dry, clean and free from surface damage to achieve a good taper connection.

Remove the Trial Head and place (Fig. 36) the definitive Humeral Head.





Impact the definitive Humeral Head, by using the Anchor and Head Pusher (Fig. 37). The Head sits flush on the osteotomy plane.

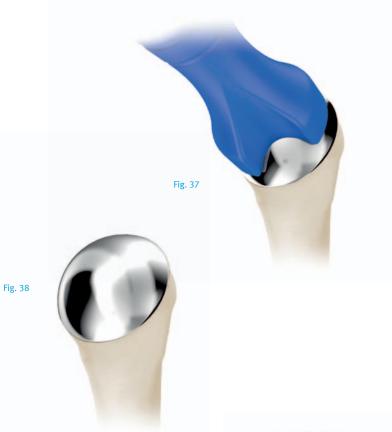
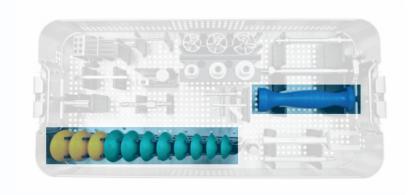


Fig. 39

The *Sidus* Stem-Free Shoulder is now implanted (Fig. 38 and 39).





# **Revision/Intra-operative Correction Surgical Steps**

#### 1. Head removal

To remove the Humeral Head (Fig. 40), slide the Head Distractor between the collar of the Humeral Anchor and the undersurface of the Humeral Head (Fig. 41).

Firmly tap the end of the instrument to loosen the Head (Fig. 42).

This instruments can be used to remove either Trial Head or the Implant Head.







#### 2. Anchor removal

To remove the Shoulder Humeral Anchor (Fig. 43):

1. Use the Fixation Release (Chisel) in any of the four (4) osteotome slots on the

Humeral Anchor to reduce or release any bone connected to the Anchor (Fig. 44 and 44a).

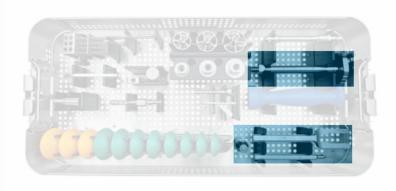
2. Thread the Revision Connector to the Humeral Anchor. Attach the Impactor/ Extractor to the Revision Connector. Gently tap the Impactor/Extractor with a hammer to remove the Humeral Anchor from the bone (Fig. 45).







The Implant is now removed (Fig. 46).



# **Glenoid Preparation Surgical Steps**

#### **Anatomical Shoulder Glenoid**

#### **Glenoid Preparation**

Sidus Stem-Free Shoulder in combination with the Anatomical Shoulder Glenoid (Fig. 47) (Tab 2). After removal of all the glenoid osteophytes (Fig. 48), a ring retractor is inserted and the proximal humerus levered out posteriorly. The inferior capsule must be incised carefully preserving the axillary nerve. Attention should be paid that the axillary nerve is protected. The caudal capsule is incised and the glenoid exposed.

Set the correction of version determined from the preoperative CT scan on the positioning guide. Identify the optimal position for the guide pin and introduce the Guiding Pin/Kirschner Wire

Introduce the Guiding Pin/Kirschner Wire into the Guiding Instrument for Glenoid.

The laser marking on the 3 mm Kirschner Wire (a) must disappear slightly into the eyelet of the positioning guide (Fig. 49).

After this, remove the positioning guide over the 3 mm Kirschner Wire.

The 3 mm Kirschner Wire is now perpendicular to the required alignment of the articulating surface, which was determined preoperatively.

The reamer and then the handle are mounted on the guide wire. For sclerotic glenoids the separate reamer (Fig. 49a) may be used to start the reaming process.



Humerus head-	glenoid "mismatch" in m				
	ملر				
Ø	S	М	L		
38	6.5	8	10		
40	6.5	8	10		
42	6	7.5	9.5		
44	5	6.5	8.5		
46	3.5	5	7		
48	2.5	4	6		
50	2	3.5	5.5		
52	1	2.5	4.5		

preferred combination

Tab. 2





If it is not impossible to insert the Kirschner Wire, drill a hole in the center of the glenoid with the glenoid drill and then machine the glenoid using the handle with the center locator (Fig. 50).

Now ream the glenoid on the basis of the reamer size and ream in the new alignment of the articulating surface (Fig. 50).

The size of the last-used reamer corresponds to the size of the glenoid.

If selection of the prosthetic components is difficult, it is generally preferable to err towards smaller heads and towards larger glenoids.

#### Note:

All sizes of Glenoids can be combined with all sizes of Humeral Heads (see "the Glenoid Component" Tab. 2).

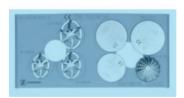
#### Optional:

If the need of an exact inferior/superior and/or retroversion correction is identified during pre-operative planning with CT-scans, the optional 3D Guiding Instrument (with 5° steps laser marks) can be used.

Continue with the pegged glenoid preparation on page 30 or with the Keeled Glenoid preparation on page 32.







### **Glenoid Components with 4-Peg Anchoring**

Now guide the Glenoid Drill Guide (left and right version) with the central hole along the 3 mm Kirschner Wire and place it on the surface of the glenoid.

The Glenoid Drill Guide can be secured in place by means of a drilled centering peg (Fig. 51).

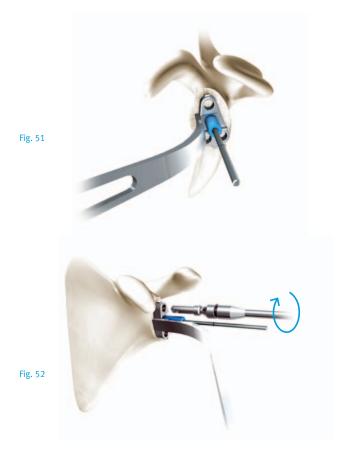
The proximal hole, with a diameter of 6.2 mm, is now drilled with the glenoid drill.

Care should be taken to ensure that drilling is continued as far as possible with the Glenoid Drill Guide (Fig. 52).

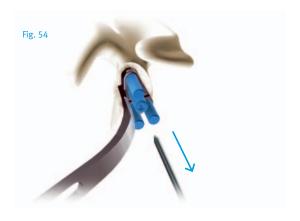
The drill can be used with either the flexible or the rigid shaft.

The proximal drilled hole is then fixed with a centering pin and one of the two distal holes is drilled. The second hole is then fixed with a centering pin while the second distal hole is drilled (Fig. 53).

The third drilled hole is now fixed and the 3 mm Kirschner Wire and the centering pins are removed (Fig. 54).







Now drill the central hole (Fig. 55).



Engage all the holes with the countersink in order to enable seating of the glenoid component (Fig. 56).



Fig. 56

Now insert the Glenoid Trial prosthesis into the prepared glenoid, using the holding forceps (Fig. 57).





#### **Glenoid Component with Keel**

Now guide the Glenoid Drill Guide with the central hole along the 3mm Kirschner Wire and place it on the surface of the glenoid.

The Glenoid Drill Guide can be secured in place by means of a drilled centering peg (Fig. 58).

**Note:** The Glenoid Drill Guide is available in two sizes: S/M for the Keeled Glenoid sizes S and M, and size L for the Keeled Glenoid size L. Both sizes are available in left and right version.

The proximal drilled hole is then fixed with a centering pin and the distal hole is drilled (Fig. 58). The second hole is then also fixed with a centering pin. The 3 mm Kirschner Wire and the centering pin are removed.

Now drill the central hole approximately 5mm into the glenoid (Fig. 58). Remove the two pins afterwards. Now drill the central peg until you have reached the Glenoid Drill Guide stop.

To help ensure optimal preparation of the bone to the keel glenoid, as a final step the bone is processed into a keel with a rasp (Fig. 59). This step gives optimal concentration of the bone.

If drill guide of sizes S or M were used, use Rasp size S and M. The rasp of size L is used, if the size L drill guide was used.

The glenoid test prosthesis is inserted into the prepared glenoid using the holding forceps (Fig. 60).



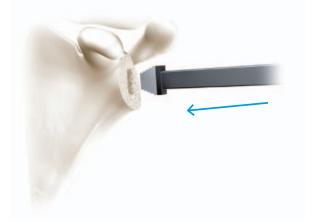




Fig. 60

Fig. 58

Fig. 59

#### **Cementing**

The glenoid trial prosthesis is used to test whether the seating is stable (Fig. 61).

The glenoid surface and the anchoring holes are now carefully cleaned and dried. The anchoring holes are filled with bone cement and the cement is pressed into the anchoring holes with a clean compress or using the instruments shown in in (Fig. 62 and 63). Glenoid Cement Setting Instrument for pegged glenoid can be used for all three sizes S, M and L (Fig. 62).

The Glenoid Cement Setting Instrument for a Keeled Glenoid consists of a handle and two pressurization adapters in size S/M and L (Fig. 63).

The anchoring holes are then filled where necessary and cement is applied to the glenoid surface and the back side side of the implant.

The implant is then cemented into place using the glenoid impactor (Fig. 64).

**Note:** Do not use the Impactor on the implant while the cement is hardening.

The excess cement is immediately and carefully removed with a knife blade.

**Note:** Continue the Surgical Steps with chapter "Head Size Determination and Positioning" on page 16.

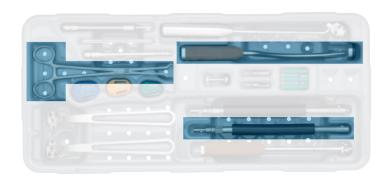






Fig. 64

Fig. 61





# Optional – Further Combination Possibilities

#### **Compatibility Matrix**



#### **Sidus Stem-Free Anchor**

REF Description/Size
01.04555.110 S
01.04555.120 M
01.04555.130 L



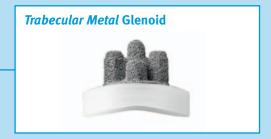
#### Bigliani/Flatow Shouler Standard Heads

REF	Description/Size	REF	Description/Size
00-4300-056-42	42 mm x 56 mm	00-4300-046-33	33 mm x 46 mm
00-4300-056-39	39 mm x 56 mm	00-4300-046-30	30 mm x 46 mm
00-4300-056-36	36 mm x 56 mm	00-4300-046-27	27 mm x 46 mm
00-4300-056-33	33 mm x 56 mm	00-4300-046-24	24 mm x 46 mm
00-4300-056-30	30 mm x 56 mm	00-4300-046-21	21 mm x 46 mm
00-4300-052-36	36 mm x 52 mm	00-4300-046-18	18 mm x 46 mm
00-4300-052-33	33 mm x 52 mm	00-4300-046-15	15 mm x 46 mm
00-4300-052-30	30 mm x 52 mm	00-4300-040-27	27 mm x 40 mm
00-4300-052-27	27 mm x 52 mm	00-4300-040-24	24 mm x 40 mm
00-4300-052-24	24 mm x 52 mm	00-4300-040-21	21 mm x 40 mm
00-4300-052-21	21 mm x 52 mm	00-4300-040-18	18 mm x 40 mm
00-4300-052-18	18 mm x 52 mm	00-4300-040-15	15 mm x 40 mm



#### **Bigliani/Flatow Shouler cemented Glenoids**

REF	Description/Size
00-4300-080-00	BF Glenoid with Keel 40 mm
00-4300-081-00	BF Glenoid with Keel 46 mm
00-4300-082-00	BF Glenoid with Keel 52 mm
00-4300-085-00	BF 40 mm Pegged Glenoid
00-4300-086-00	BF 40 mm Pegged Glenoid
00-4300-087-00	BF 40 mm Pegged Glenoid



#### Trabecular Metal Glenoid

KEF	Description/ Size
00-4326-040-00	40 mm
00-4326-040-46	40 x 46 mm Articular Surface
00-4326-046-00	46 mm
00-4326-046-40	46 x 40 mm Articular Surface
00-4326-046-52	46 x 52 mm Articular Surface
00-4326-052-00	52 mm
00-4326-052-46	52 x 46 mm Articular Surface
00-4326-052-56	52 x 56 mm Articular Surface

# Additional Surgical Technique and Instrument Sets

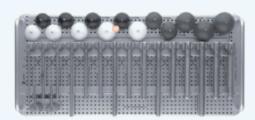
#### **Surgical Technique**

For the use of the *Bigliani/Flatow* Humeral Standard Head in combination with the *Sidus* Stem-Free Anchors, all surgical steps are identical aside from the use of the B/F Humeral Heads Provisionals.

Use the *Bigliani/Flatow* Humeral Head provisionals without Central Pin Positioner for Head Size Determination and Positioning Page 16.

#### **Instrument Sets**

*Bigliani/Flatow* Humeral Instrument Set (KT-4300-000-02),



Optional Set, Expanded Humeral Head Provisionals Expanded Head Provisional Set (00-4300-000-24)



Sidus Stem-Free Shoulder Anchor			<b>Small</b> 01.04555.110				<b>Medium</b> 01.04555.120			<b>Large</b> 01.04555.130						
Bigliani/Flatow	Ø 40	15	18	21	24	27										
Humeral Head	Ø 46	15	18	21	24	27		24	27	30	33					
	Ø 52		18	21	24		21	24	27	30			33	36		
	Ø 56											30	33	36	39	42

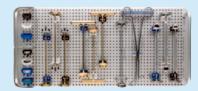
Tab. 3



Lit.No.: 97-4301-106-00

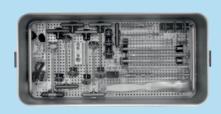


*Bigliani/Flatow* Glenoid Tray KT-4300-0001





Lit.No.: 97-4301-204-00



*Trabecular Metal* Glenoid Tray 00-4327-000-02

#### **Postoperative Treatment**

It is the responsibility of the doctor to decide which postoperative treatment is appropriate depending on each patient's health condition with the understanding that an uncemented implant will generally perform better when load activities in the first few weeks are not too aggressive. The following outlines recommendations which are generally made by surgeons. From the first day after the operation the patient may take the arm out of the immobilizing dressing several times a day to stretch his elbow. On the day of the operation pendulum exercises may be started, on the first day with passive flexing exercises, best performed using a cord passed over a roller. Depending on the intra-operative findings, active exercises may be started from the third week. If the rotator cuff was sutured or reconstructed, an abduction splint may be necessary for 4 to 6 weeks.

### **Implant Overview**

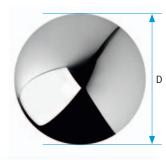
# **Sidus Stem-Free Shoulder Humeral Anchor**





Size	D mm	H mm	REF
S	24	16	01.04555.110
M	28	19	01.04555.120
1	32	22	01 04555 130

# **Sidus Stem-Free Shoulder** Humeral Head





D mm	H mm	REF
38	13	01.04555.380
40	14	01.04555.400
42	15	01.04555.420
44	16	01.04555.440
46	16	01.04555.460
48	17	01.04555.480
48	20	01.04555.485
50	18	01.04555.500
50	21	01.04555.505
52	19	01.04555.520
52	23	01.04555.525

#### **Anatomical Shoulder Glenoid**



Pegged Glenoid, cemented

Keeled Glenoid, cemented

Size	REF
S	01.04214.340
M	01.04214.370
L	01.04214.400

Size REF S 01.04214.345

M 01.04214.375 L 01.04214.405

#### **Instrument Overview**

#### **Humeral Preparation**



Sidus Stem-Free Shoulder Instrument Tray (complete) REF ZS01.04555.000

Sidus Stem-Free Shoulder Tray Lid

> REF 00-5900-099-00

Sidus Stem-Free Shoulder Tray (empty)

REF 01.04555.600



Alignment Guide "Supraspinatus" REF 01.04555.620



Resection Guide
REF
01.04555.630



Kirschner Wire 2x100 mm REF 01.04555.640



Kirschner Wire 2x70 mm REF 01.04555.650



Gentral pin 3.2 mm REF 01.04555.660



Central pin positioner

REF

01.04555.670



Impactor/Extractor
REF
01.04555.680



Irial	head	
Size		REF
38-1	13	01.04555.690
40-1	4	01.04555.700
42-1	15	01.04555.710
44-1	16	01.04555.720
46-1	16	01.04555.730
48-1	17	01.04555.740
50-1	.8	01.04555.760
52-1	9	01.04555.780



Trial head

Size

48-20 01.04555.750 50-21 01.04555.770 52-23 01.04555.790



Anchor sizer and countersink Countersink shaft

S 01.04555.800 Μ 01.04555.810 01.04555.820 L



01.04555.830



Humerus puncher

Size REF

01.04555.840 S 01.04555.850 Μ 01.04555.860 L



Anchor & head pusher

01.04555.870



Head distractor

01.04555.880



Fixation release

01.04555.890



Revision connector

01.04555.900



Pin retractor 2 mm

01.04555.910

#### **Instrument Overview**







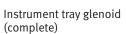


Drill guide for right glenoid

REF

72.09.10-21

Drill guide for left glenoid REF 72.09.10-11



REF ZS01.04230.030

Tray cover

REF 01.00029.029

Tray (empty)

KEF

01.04230.031



Glenoid trial prothesis S, convex

52.09.00-05



Glenoid trial prosthesis M, convex

REF 52.09.00-06



Glenoid trial prosthesis L, convex

REF 52.09.00-07



Countersink for glenoid REF 72.11.20-10



Centering peg for glenoid REF 72.11.20-11



Drill  $\varnothing$  6.2 mm for glenoid REF 72.11.20-13



Handle for cutter for glenoid REF 72.09.01-30



Handle for milling cutter glenoid

REF
72.11.20-01



Glenoid cement setting instrument

REF
01.04230.801



Holding forceps for glenoid 72.11.20-05



Rigid shaft REF 75.80.08



Guiding instrument for glenoid, 3D

\* Not included in ZS01.04230.030

01.0236.250\*



Humeral cut protection, Ø 40 mm

72.11.20-40



Kirschner Wire for glenoid 72.09.01-20



Flexible shaft 75.80.04



Guiding instrument for glenoid REF 72.09.01-10



Humeral cut protection, Ø 44 mm

REF 72.11.20-44



Impactor for glenoid 72.11.20-04



Screw for guiding instrument Glenoid reamer S for glenoid

01.04237.111



01.04236.534



Humeral cut protection, Ø 48 mm

72.11.20-48



Glenoid reamer M 01.04236.537



Glenoid reamer L 01.04236.540



Sclerotic milling cutter S 72.11.10-34



Instrument tray keeled glenoid (complete)

RFF

ZS01.04230.035

Small tray cover

REF

01.00029.032

Tray (empty)

REF

01.04230.038



Drill guide for keeled glenoid S & M

REF

01.04237.211



Drill guide for keeled glenoid

REF

01.04237.212



Glenoid trial prosthesis L, keeled

REF

01.04237.110



Glenoid trial prosthesis M, keeled

REF

01.04237.120



Glenoid trial prosthesis S, keeled

REF

01.04237.130



Glenoid cement setting instrument S & M

REF

01.04237.215



Glenoid cement setting instrument L

REF

01.04237.216



Handle glenoid cement setting instrument

REF

01.04237.217



Rasp for glenoid S & M

REF

01.04237.213



Rasp for glenoid L

REF

01.04237.214

#### **Zimmer Shoulder Solutions**

Early Stage Last Salvage

Durom® Shoulder Cemented Resurfacing Cup

Sidus Stem-free Shoulder

Variable head/stem angles; cemented 4-pegged Glenoid Multiple fixed head/ stem angles, 42°, 45° and 48°; compatible with a press-fit *Trabecular Metal* Glenoid Fixation

Anatomically fixed head/stem angle; anatomical repositioning of tuberosities The convertible Anatomical Shoulder Inverse/Reverse system allows the combination either with a convex, bone preserving design, or with a premium ingrowth Glenoid fixation

Bipolar fixed head/stem angle; "last salvage" shoulder solution



#### Disclaimer

This documentation is intended exclusively for physicians and is not intended for laypersons. Information on the products and procedures contained in this document is of a general nature and does not represent and does not constitute medical advice or recommendations. Because this information does not purport to constitute any diagnostic or therapeutic statement with regard to any individual medical case, each patient must be examined and advised individually, and this document does not replace the need for such examination and/or advice in whole or in part.

Please refer to the package inserts for important product information, including, but not limited to, contraindications, warnings, precautions, and adverse effects.

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