Galaxy Fixation System
Upper Extremities

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C. Grim, MD
INDICATIONS

The Galaxy Fixation System is intended to be used for bone stabilization in trauma and orthopedic procedures, both on adults and all pediatric subgroups except newborns as required. The indications for use include:

- open or closed fractures of the long bones;
- vertically stable pelvic fractures or as a treatment adjunct for vertically unstable pelvic fractures;
- infected and aseptic non-unions;
- joint pathologies/injuries of upper and lower limb, such as:
  - proximal humeral fractures;
  - intra-articular knee, ankle and wrist fractures;
  - delayed treatment of dislocated and stiff elbows;
  - chronic, persistent elbow joint instability;
  - acute elbow joint instability after complex ligament injuries;
  - unstable elbow fractures;
  - additional elbow stabilization of post-operative unstable internal fixation.

The Orthofix Galaxy Wrist external fixator is intended for the following indications:

- intra-articular or extra-articular fractures and dislocations of the wrist with or without soft tissue damage
- polytrauma
- carpal dislocations
- unreduced fractures following conservative treatment
- bone-loss or other reconstructive procedures
- infection

NOTE: The Shoulder Fixation System is intended to be used for proximal humeral fractures where two thirds of the metaphysis is intact.

For MRI Information see page 58.

The Rods and bone screws are strictly single patient use.
FEATURES AND BENEFITS

**Rods**
Strong radiolucent rods in three different diameters (12mm for Lower Limb, 9 and 6mm for Upper Limb) and various lengths.

### Code Description
- **Rods Diam. 12mm**
  - 932100: Rod 100mm long
  - 932150: Rod 150mm long
  - 932200: Rod 200mm long
  - 932250: Rod 250mm long
  - 932300: Rod 300mm long
  - 932350: Rod 350mm long
  - 932400: Rod 400mm long

- **Rods Diam. 9mm**
  - 939100: Rod 100mm long
  - 939150: Rod 150mm long
  - 939200: Rod 200mm long
  - 939250: Rod 250mm long
  - 939300: Rod 300mm long

- **Semi-Circular Rods Diam. 9mm**
  - 939010: Semi-Circular Rod small 115mm long
  - 939020: Semi-Circular Rod medium 140mm long
  - 939030: Semi-Circular Rod large 165mm long

- **Rods Diam. 6mm**
  - 936060: Rod 60mm long
  - 936080: Rod 80mm long
  - 936100: Rod 100mm long
  - 936120: Rod 120mm long
  - 936140: Rod 140mm long
  - 936160: Rod 160mm long
  - 936180: Rod 180mm long
  - 936200: Rod 200mm long

- **Rod Diam. 6mm**
  - 936010: 6mm L Rod

**Screws**

- **XCaliber Bone Screws** Shaft Ø 6mm - Thread Ø 6.0-5.6mm

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- **Drill bit Ø 4.8mm when the bone is hard
- **Drill bit Ø 3.2mm in poor quality bone or in the metaphyseal region**

- **Bone Screws** Shaft Ø 6mm - Thread Ø 4.5-3.5mm

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- **Self-drilling Bone Screws** Shaft Ø 4mm - Thread Ø 3.3-3.0mm

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- **Self-drilling Bone Screws** Shaft Ø 3mm - Thread Ø 3.0-2.5mm

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- **XCaliber Cylindrical Bone Screws** Shaft Ø 4mm - Thread Ø 3.0mm

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<td>948335</td>
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Galaxy Fixation System is compatible with all Orthofix Bone Screws with shaft and thread diameters as indicated above. Please refer to the Orthofix Products Catalogue.
**Clamps for Independent Screw Placement**

Allow easy and stable connection of either a rod and a bone screw or two rods.

**Large Clamp (93010)**
- To be used with 12mm Rod and 6mm shaft bone screws
-  
- Shaft Diameter Ø 6mm
- Thread Diameter Ø 6/5.6mm

**Small Clamp (93310)**
- To be used with 6mm Rods and 4mm shaft bone screws
-  
- Shaft Diameter Ø 4mm
- Thread Diameter Ø 3.3/3mm

**Medium Clamp (93110)**
- To be used with 9mm Rod and 6mm shaft bone screws
-  
- Shaft Diameter Ø 6mm
- Thread Diameter Ø 6/5.6mm

**Large-Medium Transition Clamp (99-93030)**
- (Sterile)
- To be used with 12mm Rod (Blue Part), 9mm Rod (Yellow Part) and 4mm Shaft Bone Screws
-  
- Ø 9mm Rod
- Ø 4mm Shaft Bone Screws
- Ø 12mm Rod
-  
**Galaxy Line Extension**

**Fast Closure**
- Metal Ring - pre closure by hand without the need of wrench

**Easy of use**:
- Internal Spring + Locking profile designed to provide “friction clutch on the rod” avoid sliding during surgery

**Fast Locking**:
- Cam closure in one step

**Fast Insertion**:
- Snap Lock

**Torsional Strength**
- Internal Spring + Locking profile designed to provide high torsional strength on rod

**Easy to use**:
- Internal Teeth + Spring to provide “friction clutch” between the 2 parts of the clamp avoiding sliding during surgery
Multiscrew Clamps

Multiscrew Clamp (93020)

- To be used with 12mm Rod and 6mm shaft bone screws.
- Allows parallel screw positioning either in a T- or a straight clamp configuration.

Note: the positions of the screw holes in the multiscrew clamp refer to the screw seats of the XCaliber fixator or the 1, 3, 5 screw seats of the LRS fixator T- or straight clamps.

Stability: Internal Spring + Locking profile designed to provide high torsional strength on rod

Flexibility of use: Rotation up to +/- 35°

Fast Insertion: Snap Lock

Fast Closure: Metal Ring - pre closure by hand without the need of wrench

Galaxy Line Extension

Medium Multiscrew Clamp (99-93120) (Sterile)

- To be used with 9mm Rod and 6mm Shaft Bone Screws.
- Allows parallel screw positioning (+/- 35°) in either a T-clamp or straight clamp configuration.

Note: the positions of the screw holes in the medium screw clamp refer to the screw seats of the Small Blue D.A.F. (31000) or the pediatric LRS system (series 55000)
Clamps Closure

1. **Start Position**
   Dot on cam in line with OPEN marking on metal ring

2. **Pre-Closure**
   Turn the metal ring fully by hand

3. **Final Closure**
   Tighten the cam with Wrench

1. **Pre-Closure**
   Turn the locking screw fully by hand

2. **Final Closure**
   Tighten the locking screw with Wrench

1. **Start Position**
   Dot on cam in line with OPEN marking on the base of the clamp

2. **Pre-Closure**
   Turn the knob fully by hand

3. **Final Closure**
   Tighten the cam with Wrench
Shoulder Components

Wire Locking Clamp (93620)
It consists of two disks which lock the 2.5mm Threaded Wire (93100) passing through it (NB: the clamp must not be removed but only slackened).

Wire Targeting Device (19975)
Allows positioning and fixation of the Wire Guides (19970) which can be fixed parallel, converging or diverging according to the type of fracture. The Wire Guides must be used to insert the 2.5mm Threaded Wires correctly.

Threaded Wire (93100)

WARNING: The Threaded Wires (93100) and the Wire Locking Clamps (93620) are not MR Conditional. Any construct/frame that is using Threaded Wires and Wire Locking Clamps must therefore be considered as MR Unsafe.
**Elbow Components**

**Elbow Hinge (93410)**
- To be used with 12mm Rod for the humerus and 9mm Rod for the Ulna
- Radiolucent hinge which allows easy location of the centre of rotation of the elbow, flexion-extension (up to 175°) and micrometric distraction (15mm) of the joint

**Elbow Distractor (932200 - 93431 - 93432)**
- To distract the joint intra-operatively in case of elbow stiffness (see page 34)

**Elbow Motion Unit (93420)**
- To be used with the Elbow Hinge for passive motion
- Allows controlled, limited flexion/extension of the joint
Wrist Components

Wrist Module (93350)

NOTE: Prepare the Proximal Rod half way distracted to allow Compression/Distraction maneuvers.
EQUIPMENT REQUIRED

INSTRUMENTS TRAY
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<th>Code</th>
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<td>11138</td>
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RODS & CLAMPS TRAY*
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* to order any of the Rods or Clamps, single-packaged and sterile, please add 99- prior to the part number, ex. 99-93010
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<td>Xwire Ø1,5x150mm</td>
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TRAY CONFIGURATIONS

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<td>Galaxy Upper + Lower Complete</td>
</tr>
<tr>
<td>93992C</td>
<td>Galaxy Instruments Complete</td>
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<tr>
<td>93993C</td>
<td>Galaxy Lower + Instruments Complete</td>
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HUMERAL APPLICATION

APPROACH TO THE HUMERUS
When dealing with the humerus, consideration should be given to the radial, axillary, musculocutaneous, ulnar and median nerves and brachial artery and vein. Proximally, screws should be inserted distal to the level of the axillary nerve. They can be placed from a lateral approach or ventro-lateral direction.

The middle segment of the humerus (shaded in red) should be avoided as the radial nerve has a variable course in this area.

Distally, a screw inserted from the lateral side between the triceps and brachioradialis muscles will avoid the radial nerve as long as it is just proximal to the upper border of the olecranon fossa. A more proximal screw can be inserted just medial to the lateral border of biceps, thereby avoiding the terminal branch of the musculocutaneous nerve. An alternative is a half screw inserted from the dorsal surface.
Screw Insertion
Screw positions should be planned with regard to zone of injury; often this may extend beyond the fracture lines visible on X-ray. Further thought into possible future surgeries, including plastic surgical and internal fixation procedures, should be given. X-rays of the fracture in two planes should be available. In general, screws should be placed anterolaterally in the femur; anteriorly (1 cm medial to the tibial crest in an anteroposterior direction) in the tibia; laterally in the proximal third of humerus and posterolaterally in the distal third of the humerus. Screws should be positioned for maximum mechanical stability in each bone segment, with bicortical purchase by the screw threads and with each pin as far apart in each segment as the fracture lines and neighbouring joints allow.

Insert two screws into each main fragment free-hand using the following technique:
1) Make a 15 mm incision through skin and deep fascia. Use blunt dissection to reach the underlying bone (Fig. 1).

2) Insert a screw guide perpendicular to the longitudinal axis of the bone. Use a trocar to locate the midline by palpation (Fig. 2).

3) Keeping the screw guide in contact with the cortex by gentle pressure, withdraw the trocar, and tap the screw guide lightly to anchor the pronged end against bone (Fig. 3).
4) Insert a screw through the screw guide into the bone using the Hand Drill (Fig. 4a). While drilling, the hand drill should be held steady so that the drilling direction is maintained throughout the procedure. Once the second cortex has been reached, reduce the drilling speed; four more turns are needed so that the tip just protrudes through the distal cortex. Diaphyseal bone screws should always be inserted across the diameter of the bone to avoid off axis placement. Off axis location of screws may result in screw threads lying entirely within the cortex and not traversing the medullary canal; this may weaken the bone. In all cases the surgeon should be mindful of the amount of torque required to insert the screw. In general, it is safer drill a hole with a 4.8mm drill bit prior to insertion of these screws in diaphyseal bone (Fig. 4b).

5) If a 6mm thread diameter screw is used, insert the 4.8mm drill guide into the screw guide and introduce 4.8mm drill bit (Fig. 5). Drill at 500-600 rpm through the first cortex, checking that the drill bit is at right angles to the bone. The force applied to the drill should be firm and the drilling time as short as possible to avoid thermal damage. Once the second cortex has been reached, reduce the drilling speed and continue through the bone. Ensure that the drill bit completely penetrates the second cortex.

6) Remove the drill bit and drill guide, keeping pressure on the handle of the screw guide. The screw is inserted with the T-Wrench until it reaches the second cortex. A further 4-6 turns are required to ensure that about 2mm of the screw protrudes beyond the second cortex (Fig. 6).

**Note:** The XCaliber self-drilling screws can be inserted by hand in cancellous bone. Pre-drilling is not often needed in this area. There is no need for the tip of the screw to protrude from the second cortex.

**Warning!** If thread is tapered, repositioning the screw by turning counter-clockwise more than two turns will loosen the bone-screw interface.
**XCaliber bone screw design**

The screws have a pointed tip and flute which allow them to be inserted as self-drilling implants in cancellous bone without the need for pre-drilling. Direct insertion with a hand drill is advised in most situations, irrespective of whether uncoated or HA coated screws are used. However, when insertion of these self-drilling screws is performed in diaphyseal bone, pre-drilling is recommended; use a 4.8mm drill bit through a drill guide when the bone is hard. If the bone quality is poor or, as in the metaphyseal region, where the cortex is thin, a 3.2mm drill bit should be used.

XCaliber bone screws should never be inserted with a power tool. This may result in high temperatures and cell necrosis from too high insertion speeds. Screw insertion, whether or not pre-drilling has been performed, should always be with the XCaliber Hand Drill (91120) or Rachet T Handle + Screw Shaft Connection (93150 + 93155). The screws have a round shank which is gripped securely by the XCaliber T-handle or Hand Drill. It is important that moderate force is applied initially for the screw to engage and gain entry into the first cortex.
7) Insert the remaining screws using the same technique (Fig. 7).

**Fixator Application**

8) The two screws in each bone segment are joined by rods of suitable length; each one mounted with two clamps positioned about 30mm from the skin. They are then locked manually by turning the knurled metal ring clockwise (Fig. 8).

9) A third rod is then used to join the first two rods together by 2 more clamps, which are not yet tightened. The surgeon now manipulates the fracture, if possible under X-ray control. When the position is satisfactory, the assistant locks all the clamps firmly by tightening the cams with the Universal T-Wrench or the 5mm Allen Wrench (Fig. 9).
10) The screw shafts are then cut with the bone screw cutter (Fig. 10). Although the screws can be cut before insertion, it is difficult to gauge the length accurately, and it is recommended that they are cut after the fixator has been applied. It is important that all of the screws are inserted first, and the fixator applied with the clamps locked firmly over the screws, about 30mm from the skin. The cutter can then be slid over the screw shanks in turn and the screws cut close to the fixator clamps. This will normally result in about 6mm of screw shank protruding from the clamp. The cutter is designed so that it can be used even when screws are in adjacent seats of the multiscrew clamp. The cut ends of the screws can then be protected with screw caps. When cutting the screws, the arms of the cutter should be extended for greater efficiency and the outer end of the screw held.

**MULTISCREW CLAMP**

Insert the first screw into one of the outer holes of the multiscrew clamp guide using the same technique as described above. Insert the second screw in the remaining outer seat and cut both screw shafts with the bone screw cutter. Lastly, insert the central screw if necessary.
SHOULDER APPLICATION

OPERATIVE TECHNIQUE

Positioning the patient in the operating room

Option 1: Percutaneous fixation. The patient must be positioned supine with the Image Intensifier on the contralateral side of the fracture and the X-Ray beam at right angles to the operating table.

NOTE: In order to allow the Image Intensifier to be handled correctly, we recommend using a modular table for shoulder surgery with removable proximal components.

Option 2: Fixation using an open procedure. The patient is placed in the beach chair position.
Assess the integrity of the external distal metaphyseal area (external 2/3 of the bone circumference), representing the entry point of the osteosynthesis means.

**NOTE:** A bone block or an excessively distal fracture level can contraindicate a percutaneous procedure, for both the technically difficult wire positioning and the final stability of implant. Alternatively, an open procedure must be performed, which facilitates the entry point of the wire in the cortex.

X-rays must always be carried out in AP, trans-thoracic or outlet view, and when possible, axillary view to define the configuration, position and size of the various bone fragments. A CT scan of the humeral head should also be performed.

Anterior, posterior and trans-thoracic X-rays.

Anterior, posterior and trans-thoracic CT-scans.
Reduction of the fracture

The reduction manoeuvres must be tested before preparing the surgical field and are performed following the usual procedures. For the radiological checks, the Image Intensifier must be positioned at the head end of the patient on the homolateral side of the injured limb with the C-arm able to move freely.

The sequence shows the reduction steps: forced abduction above 90°, firm retropulsion of the humeral diaphysis.

In parallel, sequential images of the fracture site were also taken for teaching purposes:
1. arm adducted in rest position
2. upper limb abducted to 90°: note that the scapulothoracic image falsifies the true humeroscapular ratios
3. upper limb abducted 120/130°: the proximal fragment starts to engage at subacromial level providing the fulcrum for the reduction manoeuvre
4. good position and start of diaphyseal retropulsion
5. retropulsion with arm normally abducted more than 90°
6. abduction of the arm which is kept abducted about 45° with a slight push to counteract the tension of the pectoralis minor

NOTE: If the reduction is not satisfactory or cannot be obtained with external manoeuvres, the surgery must be carried out with an open procedure. In this case the patient’s position must be changed from supine to the beach chair position.
Preparation of the surgical field

The area of the acromioclavicular joint must be visible: this is important for the percutaneous insertion of the wires. A poor surgical field will result in an excessively low insertion point. The upper limb must not impede the surgeon’s movement.

Positioning of the percutaneous wires

The system has proved to achieve good stability independently from the order in which the wires have been positioned. However, positioning of the first 2/3 wires depends upon the position in which the upper limb is placed in order to maintain reduction.

NOTE: It is extremely important that in maintaining the reduction the assistant keeps the injured arm parallel to the ground: in this position the humeral head is naturally offset more posteriorly than the diaphyseal plane, which corresponds to the horizontal reference plane. This will help to insert the first wire in the front plane, with an inclination of about 20° to the ground/humeral diaphysis, in order to target the apex of the humeral head. The entry point will be about 4/5cm proximal to the deltopectoral sulcus anterior to the line parallel to the humeral diaphysis which starts at the tip of the V insertion of the deltoid. The circumflex nerve anterior to this line is frayed and working anteriorly prevents iatrogenic neurological injuries. The diaphyseal cortical entry point must be as close as possible to the surgical neck fracture site: the condition of the area should have been carefully assessed pre-operatively with CT scan.

In 3 or 4 part fractures or fractures which show a certain instability after reduction, 2 wires with proximo-distal direction must be added to stabilise the greater tubercle to the head and to the humeral diaphysis as shown in the figure, both in case the procedure is carried out percutaneously or with an open access.

This operation requires further assembly to connect the distal osteosynthesis with the proximal osteosynthesis.
Inserting the wires with the aid of the Wire Targeting Device:

1) Insert the wires at slow speed. Position the first wire using the soft tissues protective guide (Fig. 1).

The correct position of the wires must be confirmed by X-rays.

2) Lock the Wire Targeting Device to the Wire Guide, turning the external knob in a clockwise direction (Fig. 2).

3) Insert the second Wire Guide into the Wire Targeting Device, place it in the most suitable position for reducing the fracture and lock it with the external knob (Fig. 3).

4) Insert the second wire into this Wire Guide. The wires have been marked to verify the correct insertion depth, reducing the use of image intensification (Fig. 4).
5) Repeat the procedure for the remaining wires. The implant must have at least 4 wires which do not overlap (Fig. 5a e Fig. 5b). If the reduction is not satisfactory, pull back the wires until the fracture is released, without removing them completely from the diaphysis. Improve the reduction with external manoeuvres and insert the wires back until gripping the humeral head fragment.

NOTE: In 3-part fractures with detachment of the greater tubercle, 1 or 2 extra wires should be applied to stabilise the fragment. The most successful insertion point is at the level of the greater tubercle-head junction. The direction can be targeted either towards the medial diaphyseal area or towards the humeral head itself. A further clamp and rod will be necessary to stabilise the wires with proximal-distal direction.

6) Once reduction has been achieved, bend the wires (93100) at about 90° with the Wire Bender (19980), leaving a distance of about 3cm from the skin: this will facilitate medication and removal at the end of treatment (Fig. 6a). The wires are oriented in pairs of 2 so that they run approximately parallel along the same plane. The flexibility of the system and the small rotational movements still possible with a single wire permit the appropriate wire direction (Fig. 6b).

Stabilisation of the wires

7) Holding the Wire Locking Clamp (93620) in place with the Open End Wrench 10mm (81031), tighten the upper disk of the clamp using the Universal T-Wrench (91150) (Fig. 7).
8) Repeat the same procedure for the remaining pairs of wires. Cut the wire distally close to the Wire Locking Clamp (Fig. 8).

9) Connect each clamp with a Galaxy Small Clamp (93310) and then connect them with a 6mm diameter rod (Fig. 9). Test the stability of the fixation under image intensification.

10) Cover the wires with the Wire Cover (80200) (Fig. 10).
POST-OPERATIVE MANAGEMENT

The wires are kept in place for an average of 6 weeks with the arm supported in a sling, but the period may be extended up to 8 weeks depending upon the fracture type. During the first 15 days the patient must keep the shoulder strictly at rest: the sling may be removed for personal hygiene, and mobilisation of the elbow and swinging movements may be allowed several times a day. Starting from the third week, passive motion can be commenced with a range of freedom proportional to the severity of the fracture. Passive mobilisation will continue until removal of the wires.

Removal of the wires

The wires guarantee good mechanical stability until the end of treatment. Cut the 2.5mm Threaded Wire leaving enough space to connect the reverse drill to it. Anaesthesia may not be necessary. The procedure can be carried out in the Out-Patient Clinic.
ELBOW APPLICATION

Patient Positioning
a) Positioning of the patient: the patient is in a supine position. The injured arm is positioned on the table so that radiographs of the humerus can be performed. A tourniquet generally should not be applied. If concomitant injuries make open osteosynthesis necessary (radial head fracture, condyle dislocation, etc.), appropriate bleeding stoppage will be necessary if the fixator is to be applied in the same procedure. As an alternative, osteosynthesis can be first performed separately in a bloodless field. After repeated disinfection and draping, the fixator can then be applied. In this case, it is important to ensure adequate bleeding stoppage to avoid hemorrhage in the operative field after removing the tourniquet. A single-step approach with minimal tissue trauma and use of bleeding stoppage is preferred.

Hint: It can sometimes be useful to raise the shoulder by placing a rolled-up towel below it.

b) Preparation of the patient: when performing disinfection, the entire upper limb and shoulder are washed. The arm can be held by the hand during the disinfection process. For this, the patient’s hand is wrapped in an adhesive drape. As an alternative, the hand can also be disinfected. The surgeon sits at the patient’s head with the assistant on the other side of the patient. The Image Intensifier is moved in from the side. It is important that the surgeon has adequate access to the elbow when the Image Intensifier is in place.

c) Use of Image Intensifier: the left figure shows a good position for the monitor. During surgery, the surgeon and assistant should have an unobstructed view of the monitor.
OPERATIVE TECHNIQUE

1) Expose the lateral aspect of the Humerus by blunt dissection in order to avoid damage to the radial nerve, taking into account that the first screw has to be inserted at the proximal level, placed not completely lateral but 10-15 degrees anterior. Use the multiscrew clamp as a template to insert screws perpendicular to the longitudinal axis of the bone. Insert the screw guides and position the trocar (19955), into one of the outer holes of the multiscrew clamp. Use the trocar to locate the midline by palpation (Fig. 1).

NOTE: The middle segment of the Humerus should be avoided as the radial nerve has a variable course in this area.

2) Keeping the screw guide in contact with the cortex by gentle pressure, withdraw the trocar, and tap the screw guide lightly to anchor its distal end. Make sure that there are no soft tissues between the bone and the screw guide. Insert the 4.8mm drill guide (11102) into the screw guide, and drill with a 4.8mm drill bit (11001). Use a sharp drill and make sure that the drill bit is at right angles to the bone, the force is applied to the drill is firm and the drilling time as short as possible to avoid thermal damage (Fig. 2). Once the second cortex has been reached, reduce the drilling speed and continue through the bone. Ensure that the drill bit completely penetrates the second cortex.

3) Remove the drill bit and drill guide, keeping pressure on the handle of the screw guide. Insert a screw through the screw guide into the bone using the T-wrench (Fig. 3) or hand drill. While drilling, the hand drill should be held steadily so that the drilling direction is maintained throughout the procedure. The screw should completely engage the second cortex for bicortical purchase. Use the same technique for the second screw.
If Xcaliber screws are used, cut both screw shafts with the bone screw cutter. Lastly, insert the central screw if necessary (Fig. 4).

**NOTE:** In all cases the surgeon should be mindful of the amount of torque required to insert the screw. If it seems tighter than usual, it is safer to remove the screw and clean it, and drill the hole again with a 4.8mm drill bit, even if it has already been used.

**Warning!** As the thread is tapered, repositioning the screw by turning counterclockwise more than two turns will loosen the bone-screw interface.

5) Remove the 3 screw guides and lock the screws in the clamp (Fig. 5).

6) Choose a 12mm rod of appropriate length and connect it to the Multiscrew Clamp and to the Elbow Hinge (93410). Lock the rod to the hinge (Fig. 6).
The Elbow Hinge needs to be aligned with the centre of rotation of the joint and in order to achieve this:

- With the rod parallel to the longitudinal axis of the humerus, ensure that the Hinge is vertically aligned with the center of rotation of the joint and lock the rod to the Multiscrew Clamp by turning the knurled metal ring by hand (Fig. 7).

- Move the rod antero-posteriorly to achieve horizontal alignment (Fig. 8).

- Rotate the Elbow Hinge until perfectly aligned with the center of the elbow joint in the lateral view (Fig. 9).
7) Lock the Multiscrew Clamp Central Nut and check Elbow Hinge alignment under image intensification. (Fig. 10). The radiolucent central unit of the hinge has an in-built targeting cross which can be used to achieve a correct alignment, however it is advisable to use an additional 2mm k-wire (length approx. 10cm). Insert the k-wire through the hole in the centre of the hinge unit and manipulate the central hinge unit until the k-wire projects as a dot in the center of the condyles.

If necessary, a third screw can be inserted distally in the humerus to increase stability (caution to the radial nerve). In this case the screw should be inserted from a dorsal-lateral direction into the distal humerus leaving the radial nerve ventrally. Ensure the safety position of the screw by a mini-open approach.

8) Choose a 9mm rod of appropriate length, lock it to the Elbow Hinge and attach a 9mm Clamp to it. With the forearm in neutral position or pronation, align the ulnar rod with the ulna shaft. The ulnar screws may reach the shaft from the lateral side or from a slightly latero-dorsal side. A minimum of 2 screws are necessary. The screws should be well spaced for mechanical stability and they are positioned using the Galaxy Medium Clamps. For predrilling, insert the 3.2mm drill guide (11116) directly into the screw seat of the clamp and drill with the 3.2mm drill bit (11003) (Fig. 11).

9) Insert a 120/20 4.5-3.5mm cortical screw (10137) (Fig. 11a). The distal screw should be inserted first taking care that the ulnar rod is parallel to the posterior border of the ulna. After insertion of the distal screw, the clamp has to be locked tightly before the second screw hole is prepared.

10) While predrilling the second screw hole—again using the drill guide—the clamp must be fully closed. Depending on the soft tissue conditions, the elbow hinge unit can be locked for a short post-operative time (Fig. 12) or can be left open for immediate mobilisation.
11) The in-built distraction unit is not necessarily used in an acute elbow trauma. Sometimes it may help to protect the joint surfaces but distraction should be limited to 3-4mm. (Fig. 13).

12) **Option:**
Alternatively to the non-invasive targeting technique described above, it sometimes might be helpful to insert a 2mm K-wire into the center of the condyles. The K-wire is percutaneously inserted from the lateral side and its tip centered into the radiologically visible centre of the condyles (Fig. 14).

13) With the of the K-wire at the entry point of the bone, the K-wire is then drilled approx. 4cm into the bone, along the joint axis both in the lateral and AP view (Fig. 15).
14) If the wire has not been inserted exactly along the joint axis, it is seen as a small line instead of a dot in the lateral view. In this case, under fluoroscopy bend the wire exiting from the skin until it is seen as a single dot (Fig. 16).

15) The elbow assembly is then slid over the K-wire and previously inserted the humeral screws - see above described technique (Fig. 17).

The Multiscrew Clamp is then closed fully and the application of the fixator is continued with the insertion of the ulnar screws as described above.
ELBOW MOVEMENT

Free Movement
With locking screw (A) loosened, the Elbow Hinge will allow free flexion-extension (Fig. 1).

Passive Movement
1) The elbow motion unit allows either a free movement of the elbow joint or a controlled movement towards flexion and extension by turning the worm-screw clockwise or anti-clockwise with the 5mm Allen Wrench (Fig. 2).

2) Passive flexion and extension is achieved by turning the worm-screw clockwise or anti-clockwise with the 5mm Allen Wrench (Fig. 3).

Limited Movement
3) If the screws are removed from the central part of the elbow motion unit, they can be used to limit the amount of flexion and extension (Fig. 4).
ELBOW DISTRACTER UNIT
POST-TRAUMATIC STIFFNESS

The Elbow Distractor is intended to be used to distract the joint intra-operatively in case of elbow stiffness.
1) It is mandatory to expose or free the ulnar nerve prior to distraction and arthrolysis (Fig. 1).

2) Cleaning of the joint might be necessary prior to the application of the Elbow Distractor. Expose the lateral aspect of the Humerus by blunt dissection in order to avoid damage to the radial nerve, taking into account that the proximal screws are inserted first, on the antero-lateral side, at an angle of 10-15° to the frontal plane (Fig. 2).

**NOTE:** The middle segment of the Humerus (shaded in red) should be avoided as the radial nerve has a variable course in this area.

3) Use the Humeral Distractor Clamp as a template for screw insertion. Insert the Screw Guides into the clamp, perpendicular to the longitudinal axis of the bone, and position the Trocar (19950) into one of the outer holes to locate the midline by palpation (Fig. 3).
4) Keeping the Screw Guide (11137) in contact with the cortex by gentle pressure, withdraw the Trocar (19955), and tap the Screw Guide lightly to anchor its distal end. Insert the 4.8mm Drill Guide (11138) into the Screw Guide, and introduce a 4.8mm Drill Bit (11001) (Fig. 4). Drill at 500-600 rpm through the first cortex, checking that the Drill Bit is at right angles to the bone. The force applied to the drill should be firm. Use a sharp drill and make sure that the drilling time is as short as possible to avoid thermal damage.

**NOTE:** The positions of the screw seats in the Humeral Distractor Clamp refer to the screw seats of the Galaxy Multiscrew Clamp or the 1, 3, 5 screw seats of the LRS ADV Straight Clamps.

5) Once the second cortex has been reached, reduce the drilling speed and continue through the bone. Ensure that the drill bit completely penetrates the second cortex. Remove the Drill Bit and Drill Guide, keeping pressure on the handle of the Screw Guide. Insert a 110/30 cortical screw (10110) or if necessary a longer screw through the Screw Guide into the bone using the Universal T Wrench (93150+93155) (Fig. 5).

While inserting the screw, the T Wrench should be held steady so that the direction of insertion is maintained throughout the procedure. Make sure that the tip of the screw protrudes through the distal cortex (Image Intensifier).

6) Insert the second screw in the outermost hole using the same technique. If XCaliber screws are used, cut both screw shafts with the Bone Screw Cutter (91101). Lastly, insert the middle screw if necessary. Remove the Screw Guides and tighten the clamp (Fig. 6).

**NOTE:** In all cases, the surgeon should be mindful of the amount of torque required to insert the screw. If it seems tighter than usual, it is safer to remove the screw and clean it, and drill the hole again with a 4.8mm Drill Bit, even if it has already been used.

**Warning!** As the thread is tapered, repositioning the screw by turning counterclockwise more than two turns will loosen the bone-screw interface.
7) With the Ulnar Micrometric Distraction Mechanism in Close position, adjust the distance of the Humeral Distractor Clamp, making sure that the Ulnar Distractor Clamp is aligned with the Ulna (Fig. 7).

8) To ensure that distraction between the Humerus and Ulna is carried out in a concentric way without any impingement, the axis of the Micrometric Distraction Mechanism should be perpendicular to the virtual line between the coronoid and olecranon (Fig. 8).

9) Tighten the ball-joints with the Allen Wrench (10017) (Fig. 9).
10) Insert now the temporary ulnar screws for distraction. Position the Trocar (19955) in one of the available holes of the Ulnar Distractor Clamp and locate the bone. The distal screw is usually inserted first, preferably opposite to the coronoid process. Remove the Trocar, insert a 3.2mm Drill Guide (19950) and drill with a 3.2mm Drill Bit (11003) (Fig. 10a). Insert a 4.5-3.5mm bone screw (10135 or 10137) (Fig. 10b).

11) If necessary, adjust the position of the Ulnar Distractor Clamp so that its distal border is aligned with the ulna (Fig. 11b). Insert a second ulnar screw in one of the remaining holes of the Ulnar Distractor Clamp using the same procedure. This second screw should enter the olecranon.

12) Tighten the screw into the clamp with the 5mm Allen Wrench (30017) and tighten the cams with the 6mm Torque Wrench (10025) (Fig. 12).

13) Apply joint distraction by turning the Micrometric Distraction Mechanism with 5mm Torque Wrench (93440) which indicates the distraction force (9 Nm correspond approximately to 100 Kg of distraction force) (Fig. 13a e 13b). Joint distraction is checked under image intensification and the appropriate amount of distraction should be decided by the surgeon, in accordance with clinical and radiological findings.

During the distraction process, the ulnar nerve should be monitored closely to make sure that there is no tension on the nerve. If necessary the ulnar nerve has to be transposed to the ventral side. The distraction process should be repeated 2-3 times and might take 5-10 minutes to relax the capsule and the collagen fibres in the ligaments. At the end, release the distraction, remove the temporary ulnar screws and the Elbow Distractor.
14) Leave the humeral screws in place for the application of the Elbow Hinge Fixator. After having centered the hinge fixator as described from page 26 onwards and inserted the ulnar screws, use the in-built distractor unit (central unit) and re-distract the elbow minimum twice the normal joint space. Do not exceed 10mm. Once the articular surfaces have been separated in this way, the elbow joint can be forced gently into flexion and extension. The resistance must be overcome by controlled continued manipulation. The ulnar nerve must be monitored. If a severe extension deficit is treated by forcing the elbow into extension, care has to be taken to the radial nerve as this manoeuvre might damage it (Fig. 14).

15) Lock the Elbow Hinge in the maximum flexion and leave the elbow in this position for 1-3 days. After this period allow elbow mobilisation (Fig. 15), advising the patient to use the elbow motion unit.

**ATTENTION:** Pain releasing drugs can be given as soon as the neurological situation (motor and sensory function) in the operated arm is fully intact and there is no severe swelling of the forearm which could be a cause and early sign of compartment syndrome. The external fixator should be kept in place for 6-8 weeks. During this time Indometacin or equivalent drugs may be administered to release pain and inflammation at the same time. A stomach protection could be advisable.
WRIST APPLICATIONS

APPROACH TO THE WRIST

Proximal screws are placed within the middle third of the radius. At this level, the radius is covered by the tendons of extensor carpi radialis longus (ECRL) and extensor carpi radialis brevis (ECRB) as well as the extensor digitorum communis (EDC). Screws can be inserted in the standard midlateral position by retracting the brachioradialis (BR) tendon and the superficial radial nerve (SRN), in the dorsoradial position between the ECRL and ECRB or dorsally between the ECRB and EDC. Screw placement is done through a limited open approach to insure identification and protection of the radial sensory and lateral antebrachial-cutaneous nerves.

In non-bridging wrist applications, the distal screws must be applied in the safety zones between the extensor compartments dorsally and dorsoradially.

In wrist bridging applications, the distal screws are applied into the second metacarpal bone, paying attention to the extensor tendon and the radiodorsal neuro-vascular bundle on the extensor and radiodorsal side. If the screws are placed too laterally, they will impede the function of the thumb. For this reason, an angle of 30-45° dorsally from the frontal plane is preferable.
INTRA-ARTICULAR APPLICATION

Surgical Area Preparation
- Regional or general anaesthesia may be used
- Tourniquet should be available if desired
- Use a hand table
- Make sure that X-ray equipment is available
- Reduce approximately the fracture before the fixator is applied
- Place the wrist in moderate (manual) traction, flexion and radial abduction (i.e. ulnar deviation) with a folded towel on the ulnar side to support it (Fig. 1)

Distal Screws Insertion
- Insert first the proximal metacarpal screw close to the base of the bone on the flare of the tubercle
- Make a longitudinal incision to the skin for each metacarpal screw
- Dissect the soft tissues down to the bone taking care to retract the interosseus muscle anteriorly and the extensor muscle dorsally
- The screw guide is positioned on the bone with the trocar (19965) (Fig. 2)
• Insert the screw following one of the screw insertion techniques described below:

**Pre-drilled Bone Screws (4mm shaft) Insertion**
• Remove the trocar, replace it with the drill guide and drill the bone over the drill guide with a 2.7mm drill bit (Fig. 3)

• Insert a bone screw with the T-Wrench 4mm Shaft (93175) or the T-Wrench (M210) over the screw guide (Fig. 4)

**Cylindrical Bone Screws (4mm shaft) and *Self-drilling Bone Screws (3 or 4mm shaft) Insertion**
• Remove the trocar and insert the screws directly through the screw guide without pre-drilling. In case of the 4mm cylindrical screws, they are inserted using either the Screw T-Wrench QC (93160) or power drill with moderate speed. In case of the 3mm or 4mm shaft conical screws, they are inserted using either the T-Wrench (M210) or power drill with moderate speed (Fig. 5)

**NOTES:**
• Take care to avoid damage to the carpo-metacarpal joint
• There is no need for the tip of the screw to protrude from the second cortex
• Great care must be taken to ensure that the screws are inserted in the central bone axis

**WARNING:** Due to their tapered design, screws become loose if they are backed out
Small Multiscrew Clamp-Short positioning

- For an easier application of the second screw, it is advisable to temporarily fix a rod to the clamp and use it as handle.
- Insert the two screw guides into the clamp (Fig. 6).
- Insert the Small Multiscrew Clamp-Short (93330) over the first metacarpal screw.

- Use the Small Multiscrew Clamp-Short (93330) as guide to insert the second distal screw, leveraging on the rod-handle to find the most appropriate position on the bone using the trocar (Fig. 7-8).

**NOTE:** it is important to identify the central axis of the second metacarpal bone for the correct insertion of the distal screw.
• Remove the temporary rod-handle and the screw guides
• Close the clamp cover by hand to secure the screws (Fig. 9)

• Set up the wrist module in neutral position with the two rods longitudinally aligned and tighten the Arch Nut with the Allen Wrench (Fig. 10)

Galaxy Wrist Fixator positioning
• Attach the Wrist Module to the Small Multiscrew Clamp-Short (93330) and close the clamp by hand (Fig. 11)

NOTE: At this stage, both Locking Screws of the Small Multiscrew Clamp-Short should be closed only by hand enabling movement in all planes to locate the centre of rotation
• If necessary a Ø 1.5mm K-wire can be used to assist in aligning the fixator with the centre of rotation of the wrist joint, which is located within the head of the capitate1 in both flexion and extension, and radial and ulnar deviation (Fig. 12)


• Under image intensification, identify the centre of rotation of the wrist in AP and Lateral view and ensure that the Wrist Module Central Unit is aligned with it (see figure 13)
• Once the centre of rotation has been identified, both Locking Screws of the Small Multiscrew Clamp-Short (93330) are locked into position using the Allen Wrench (Fig. 16)

NOTES:
The clamp must be tightened gently in order to preserve the correct alignment, taking care that it is maintained at all times

Correct positioning of the proximal rod in ML Plane
• Loosen the ML locking Screw of the central joint (Fig. 17)
• Move the proximal rod to find its best position in this plane

Correct positioning of the proximal rod in AP Plane
• Loosen the AP locking Screw of the central joint (Fig. 18)
• Move the proximal rod to find its best position in this plane
- Mount the second Small Multiscrew Clamp-Short (93330) over the proximal rod and lock it in place
- Make a longitudinal skin incision for each screw
- Dissect the soft tissues down to the bone taking care to retract the soft tissues
- The screw guide is positioned with the trocar over the Small Multiscrew Clamp-Short (93330)

- Insert the proximal screws following one of the below screw insertion techniques (Fig. 20):

*Pre-drilled Bone Screws (4mm shaft) Insertion
- Remove the trocar, replace it with the drill guide and drill the bone over the drill guide with a 2.7mm drill bit
- Insert a bone screw with the T-Wrench 4mm Shaft (93175) or the T-Wrench (M210) over the screw guide

Cylindrical Bone Screws (4mm shaft) and *Self-drilling Bone Screws (3 or 4mm shaft) Insertion
- Remove the trocar and insert the screws directly through the screw guide without pre-drilling. In case of the 4mm cylindrical screws, they are inserted using either the Screw T-Wrench QC (93160) or power drill with moderate speed. In case of the 3mm or 4mm shaft conical screws, they are inserted using either the T-Wrench (M210) or power drill with moderate speed

NOTES:
- Pay attention to the safe corridors during pin insertion, taking care to avoid tendon entrapment and radial nerve damage
- There is no need for the tip of the screw to protrude from the second cortex
- Great care must be taken to ensure that the screws are inserted in the central bone axis
- Once they have been placed, an X-Ray should be used to verify the position and penetration of the far cortex by all four screws
- Remove the screw guides and close the clamp cover by hand (Fig. 21).

* WARNING: Due to their tapered design, screws become loose if they are backed out
• Remove the K-wire (Fig. 22)

• Check the joint movement by loosening the nut of the arch, and, if the movement is correct, lock the Wrist Module in neutral position (Fig. 23)

• Check fracture reduction under X-Ray and if necessary restore the wrist anatomy with the fixator in place before locking all clamps

NOTE: Close tightly both the ML and the AP Locking Screw using the Allen Wrench. If the joint does not move freely, adjust the position of the Wrist Module before locking the Arch Nut and the AP/ML Locking Screws
CONTROLLED RANGE OF MOVEMENT
- The system allows for a ±20° and ±40° controlled flexion-extension movement of the wrist
- In order to achieve this, first loosen the Arch Nut (Fig. 24)

- Push the Motion Range Selector in its outer seat with the Allen Wrench to gain ±40° of freedom (Fig. 25)

- Push the Motion Range Selector in its inner seat with the Allen Wrench to gain ±20° freedom (Fig. 26)
COMPRESSON-DISTRACTION

• Loosen the Compression-Distraction Set Screw (Fig. 27)

• Using the Allen Wrench, turn the Compression-Distraction Nut clockwise or anti-clockwise to achieve distraction or compression respectively (Fig. 28)
If necessary, cut the screws shaft with the 4mm Cutter (94401 - not provided in the tray) (Fig. 29)
EXTRA-ARTICULAR APPLICATION

- If there is no extra-articular involvement of the fracture line and the epiphyseal fragment has a volar length of 10mm minimum, bridging of the joint is not required and the following technique is applicable

Surgical Area Preparation
- Regional or general anaesthesia may be used
- Tourniquet should be available if desired
- Use a hand table
- Make sure that X-ray equipment is available
- Sterilise the skin over the iliac crest in case a bone graft should be needed
- Reduce approximately the fracture before the fixator is applied
- Place the wrist in moderate (manual) traction, flexion and radial abduction (i.e. ulnar deviation) with a folded towel on the ulnar side to support it (Fig. 31)
- Identify the Lister tubercle and the safe corridors

Distal Screws Insertion
- Make a longitudinal incision to the skin for each screw, making sure to follow safe corridors
- Dissect the soft tissues down to the bone taking care to retract the muscles
- The screw guide is positioned over the bone with the trocar
• Insert first the lateral proximal screw following one of screw insertion techniques described below:

*Pre-drilled Bone Screws (4mm shaft) Insertion
• Remove the trocar, replace it with the drill guide and drill the bone over the drill guide with a 2.7mm drill bit
• Insert a bone screw with the T-Wrench 4mm Shaft (93175) or the T-Wrench (M210) over the screw guide

Cylindrical Bone Screws (4mm shaft) and *Self-drilling Bone Screws (3 or 4mm shaft) Insertion
• Remove the trocar and insert the screws directly through the screw guide without pre-drilling. In case of the 4mm cylindrical screws, they are inserted using either the Screw T-Wrench QC (93160) or power drill with moderate speed. In case of the 3mm or 4mm shaft conical screws, they are inserted using either the T-Wrench (M210) or power drill with moderate speed

NOTES:
• Take care to avoid damage to the radial nerve and the radio-ulnar joint
• After insertion, check by X-Ray the screw positions
• The screw should engage the volar cortex securely and penetrate by one thread
• Carefully evaluate the bone quality before positioning the screws
• There is no need for the tip of the screw to protrude from the second cortex
• Great care must be taken to ensure that the screws have the maximum bone purchase

* WARNING: Due to their tapered design, screws become loose if they are backed out

Small Multiscrew Clamp-long positioning
• Insert the Small Multiscrew Clamp-Long (93320) over the first distal screw
• For an easier application of the second screw, it is advisable to temporarily fix a rod to the clamp to be used as handle before positioning the Small Multiscrew Clamp-Long (93320) as described in Fig. 6, Page 43 for the Small Multiscrew Clamp-Short
• Insert the second distal screw following the same procedure described above (Fig. 34)
• Remove the screw guides and close the Locking Screw of the Clamp cover by hand
• Attach the L-Rod to the Small Multiscrew Clamp-Long
• Close the clamp by hand (Fig. 36)

• Insertion of the proximal screws is carried out at a distance of about 14cm from the distal screws depending on the fracture site
• Make a 25mm incision to the skin in order to avoid injury to the superficial branch of the radial nerve
• Screw insertion must follow one of the techniques described above, paying attention to safe corridors (Fig. 37)
• All clamps are tightened using the Allen Wrench avoiding loss of position (Fig. 38)

• If necessary cut the screw shafts with the 4mm Cutter (94401-not provided in the tray) (Fig. 39)
UPPER LIMB APPLICATIONS

- Shoulder
- Humerus
- Elbow
- Forearm
- Wrist
MRI INFORMATION

Galaxy System Fixator Components are labeled MR CONDITIONAL according to the terminology specified in ASTM F2503 Standard Practice for Marking Medical Devices and Other Items in the Magnetic Resonance Environment.

Non-clinical testing has demonstrated that the Galaxy System Fixator Components is MR Conditional according to the terminology specified in ASTM F2503 Standard Practice for Marking Medical Devices and Other Items in the Magnetic Resonance Environment. Non-clinical testing, done according to ASTM F2052-06, F2213-06, F2182–11, F2119-07, demonstrated that a patient with the Galaxy Fixation System can be safely scanned under the following conditions:

- Static magnetic field of 1.5 Tesla and 3.0Tesla
- Maximum spatial magnetic field gradient of 900-Gauss/cm (90mT/cm)
- Maximum whole-body-averaged specific absorption rate (SAR) of 4.0 W/kg in the First Level Controlled Mode for 15 minutes of scanning.
- No local transmit/receive coils can be used on the device.
- The Galaxy Fixation System must be entirely outside the MR scanner bore. No part of the Galaxy Fixation System must extend into the MR bore. Therefore MR scanning of body parts where the Galaxy Fixation System is located is Contraindicated.

**Note:** All components of Galaxy Fixation System frames must be identified as MR Conditional prior to being placed in or near an MR Environment.

The Threaded Wires (93100), the Wire Locking Clamps (93620), and the L-Rod (936010) and Semi-Circular Rods (939010, 939020, 939030) are not MR Conditional. Any construct/frame that is using Threaded Wires, the Wire Locking Clamps, the L-Rod and Semi-Circular Rods must therefore be considered as MR Unsafe.

HEATING INFORMATION

Comprehensive electromagnetic computer modeling and experimental testing was performed on the following systems:

- 1.5-Tesla/64-MHz: Magnetom, Siemens Medical Solutions, Malvern, PA. Software Numaris/4, Version Syngo MR 2002B DHHS Active-shielded, horizontal field scanner
- 3-Tesla/128-MHz: Excite, HDx, Software 14X.M5, General Electric Healthcare, Milwaukee, WI, Active-shielded, horizontal field scanner

To determine the worst heating in ten configurations of Orthofix Galaxy Fixation System. From these studies, it is concluded that once the entire external fixation frame is visible outside the MRI bore, the maximum heating is less than 1 degree Celsius. In non-clinical testing the worst scenarios produced the following temperature rises during MRI under the conditions reported above:

<table>
<thead>
<tr>
<th>Galaxy Fixation System</th>
<th>1.5 Tesla System</th>
<th>3.0 Tesla System</th>
</tr>
</thead>
<tbody>
<tr>
<td>Minutes of scanning</td>
<td>15</td>
<td>15</td>
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<tr>
<td>Calorimetry measured values, whole body averaged SAR (W/kg)</td>
<td>2.2 W/Kg</td>
<td>2.5 W/Kg</td>
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<tr>
<td>Highest temperature rise less than (°C)</td>
<td>1°C</td>
<td>1 °C</td>
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</table>

Please note that temperature changes reported apply to the designed MR systems and characteristics used. If a different MR system is used, temperature changes may vary but are expected to be low enough for safe scanning as long as all Galaxy System Fixator Components are placed outside the MR bore. Since different frame configurations and frame sizes might lead to higher temperature increases, Orthofix recommends that the SAR settings are minimised as much as possible.

DISPLACEMENT INFORMATION

The system will not present an additional risk or hazard to a patient in the 3-Tesla 1.5 and MR environment with regard to translational attraction or migration and torque.

ARTIFACT INFORMATION

MR image quality may be compromised if the area of interest is in the same vicinity or relatively close to the position of the device. For complete information on MR indications, please refer to PQ GAL.
MR PATIENT SAFETY

MRI in patients with Galaxy Fixation System can only be performed under these parameters. It is not allowed to scan the Galaxy Fixation System directly. Using other parameters, MRI could result in serious injury to the patient. When the Galaxy Fixation System is used in conjunction with other External Fixation Systems please be advised that this combination has not been tested in the MR environment and therefore higher heating and serious injury to the patient may occur. Because higher in vivo heating cannot be excluded, close patient monitoring and communication with the patient during the scan is required. Immediately abort the scan if the patient reports burning sensation or pain.

Galaxy Fixation System can only be guaranteed for MRI when using the following components to build a frame: (*the following components are listed in non-sterile configuration. Please consider that the same MRI information and performance are applicable to the same components in gamma-sterile configuration if available (code number preceeded by 99- e.g 99-93030))

**RODS**

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<td>Rod 150mm long, 12mm diameter</td>
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<td>932200</td>
<td>Rod 200mm long, 12mm diameter</td>
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**CLAMPS**

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**GALAXY WRIST**

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<td>93350</td>
<td>Wrist Module</td>
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**References**

Instructions for Use: See actual package insert for Instructions for Use.

Caution: Federal law (USA) restricts this device to sale by or on the order of a physician.

Proper surgical procedure is the responsibility of the medical professional. Operative techniques are furnished as an informative guideline. Each surgeon must evaluate the appropriateness of a technique based on his or her personal medical credentials and experience. Please refer to the "Instructions for Use" supplied with the product for specific information on indications for use, contraindications, warnings, precautions, adverse reactions and sterilization.