Zimmer® PSI Knee System
For Use with Persona®
The Personalized Knee System
Surgical Technique
# Zimmer® PSI Knee Surgical Technique

## Table of Contents

1. **Intra-Operative Guide**
2. **Femur Exposure**
3. **Position the Femoral PSI Jig**
4. **Pin the Distal Cutting Guide Pin Holes**
5. **Drill 4-in-1 Cutting Guide Pin Holes**
6. **Resect Distal Femur**
7. **Locate Anterior Cutting Guide Pins**
8. **Place 4-in-1 Femoral Finishing Guide**
9. **Position Tibial PSI Jig**
10. **Verify Tibial PSI Jig Alignment**
11. **Pin Tibia Cut Guide Pin Holes**
12. **Remove Tibial PSI Jig**
13. **Resect Proximal Tibia**
14. **Optional: Install PSI Tibia Rotation on Persona Sizing Plate Handle**
15. **Set Tibial Rotation**
16. **Verify Overall Alignment**
17. **Supported Zimmer Persona Systems**
18. **Cleaning/Sterilization Methods And Equipment Inventory**
   - **Cleaning**
   - **Sterilization Parameters**
   - **PSI Knee Reusable Instruments and Additional Specific Cleaning Instructions**
19. **Zimmer PSI Knee Disposable Kits**
20. **Reusable Instruments Ordering**
21. **Zimmer Contact Information**
Introduction

Overview

The Zimmer® Patient Specific Knee System consists of: disposable patient specific tibial and femoral instrument guides (also called PSI jigs), optional bone models, and an optional Tibial Rotational Guide (to set the axial rotation of the tibial component), per the available kits listed in the section titled “Zimmer® PSI Knee Disposable Kits”. A copy of the approved pre-operative surgical planning is also provided in the Zimmer® PSI Knee packaging to be referenced by the surgeon intra-operatively. The bone models are a reconstruction of the patient’s knee joint tibial and femur bones from the medical imaging data, aiding the surgeon in verifying the Zimmer PSI Jigs’ placement intra-operatively, they are required to hold a place in the sterile field prior and during the surgery.

The customized PSI instrument guides are to be used with the given Persona® implant families as described in the following section, “Indication for Use”. The Zimmer® PSI Knee instrument guides are placed on the distal femur and proximal tibia intra-operatively, and have pin holes to allow the surgeon to precisely insert reference pins, in accordance with the pre-operative surgical plan, that set the position of the cut guides.

The PSI Knee Reusable instruments, provided by Zimmer CAS, are listed in the section titled “Reusable Zimmer® PSI Knee Instruments”. All other reusable instruments that are part of the applicable standard instrumentation sets, described in the “Intra-Operative Guide” section, are listed with a Persona identifier.

The scope of this document is to provide information on the surgical technique, cleaning/sterilization methods, as well as the available Zimmer® PSI Knee kits. The pre-operative guide and instructions for use of the Zimmer® PSI Knee Planner application are provided in the Zimmer® PSI Knee Planner Software User Guide 97-5970-035-00.

Indication for use

The Zimmer® PSI Knee System is indicated as an orthopedic instrument system to assist in the positioning of knee replacement components. It involves surgical planning software used pre-operatively to plan the surgical placement of the components on the basis of provided patient radiological images with identifiable placement anatomical landmarks, and surgical instrument components that include patient specific or customized guides fabricated on the basis of the surgical plan to precisely reference the placement of the implant components intra-operatively per the surgical plan.

The Zimmer® PSI Knee System is to be used with the following fixed bearing knee replacement systems in accordance with their indications and contraindications: NexGen® CR, NexGen CR-Flex, NexGen CR-Flex Gender, NexGen LPS, NexGen LPS-Flex, NexGen LPS-Flex Gender, Persona® CR and Persona® PS.

The patient specific guide components are intended for single-use only.

Contraindications

The Zimmer PSI Knee system should not be used in any of the following situations: in cases with active infections of the knee joint, in cases with Hip-Knee-Ankle (HKA) alignment deformities larger than 15° varus or valgus, in cases where femoral anterior cut first surgical techniques will be used, in cases of knee replacement revision surgery, or in cases which are contraindicated for the implant as given by Zimmer.

Complications

Possible complications associated with the use of the system may include, but are not limited to: infection, complication due to misplacement of the implants that may potentially lead to dislocation, leg misalignment or knee ligament imbalance. The occurrence of one of these complications may affect the patient’s mobility.
Precautions
The following are general precautions and warnings related to the use of Zimmer PSI instrument guides:

- **Caution: Federal (U.S.) law restricts this device to sale by or on the order of a physician.**
- Zimmer strongly recommends formal Zimmer PSI Knee System training prior to use of the system. Contact your local Zimmer representative or the Zimmer Institute (1-855-ZSurgeon or 1-855-978-7436) for more information.
- The Zimmer PSI Knee System should not be used to perform surgical procedures other than those specified in this surgical technique.
- The Zimmer PSI Knee System should be used in conjunction with a femur first technique.
- The Disposable Zimmer PSI Knee Instruments, including instrument guides and bone models are patient specific and single use and should be discarded after surgery.
- The Disposable Zimmer PSI Knee Instruments and Reusable Zimmer PSI Knee Instruments are provided non-sterile and must be cleaned and sterilized before use per instructions provided in this surgical technique (in the section “Cleaning/Sterilization Methods and Equipment Inventory”). These instructions are also provided with the components, refer to Zimmer PSI Jigs & Bone Models Package Insert (20-8014-043-00).
- The Disposable Zimmer PSI Knee Instruments have a limited shelf life of 6 months after the manufacturing date, as indicated on the package label. Given the potential for patient morphological changes, the surgeon will need to reassess the patient to identify any potential changes prior to surgery. In case of any doubt the Zimmer PSI Knee guides and bone models must not be used.
- The Disposable Zimmer PSI Knee Instruments are to be used with the given implant system per the related pre-operative planning. The implant must be used in accordance with its respective package labeling. The user should refer to the surgical technique published by the implant manufacturer.
- The Disposable Zimmer PSI Knee Instruments can withstand two autoclave sterilizations. Re-sterilization is only permissible when they have not been in contact with the patient or otherwise contaminated.
- The Disposable Zimmer PSI Knee Instruments are designed to fit the patient anatomy as it was at the moment when the patient radiological images were acquired. If the anatomy or condition of the articular surface has changed since the radiological images were acquired, the patient specific instrument should not be used.
- If you experience difficulties with the Zimmer PSI Knee Jigs during surgery, stop using the Jigs and revert to the standard (non-PSI) surgical technique.

**Warning:** Ensure that the delivered Disposable Zimmer PSI Knee Instruments correspond to the intended patient. A copy of the approved surgical plan is provided in the Zimmer PSI Knee packaging. Only use the Disposable Zimmer PSI Knee Instruments if the PSI Case ID marking are both legible on the Zimmer PSI Knee instrument guides and bone models and match the PSI Case ID specific to the intended patient. If the two PSI Case ID markings do not match, DO NOT USE the Disposable Zimmer PSI Knee Instruments on the patient and notify your Zimmer representative.
The PSI Case ID can be either 8 or 15 characters, automatically assigned based on region. The nomenclature in the following table is based off of a fictitious patient with a First Initial: S, the First Two Letters of the Last Name: AM, and Operating Side: Left (L). The Marking on the Guides and Bone Models will be the whole case ID if it’s 8 characters and the first seven digits if it is 15 characters (Fig. 1).

### PSI Case ID with 8 Characters

**EXAMPLE: SAM1234L**

<table>
<thead>
<tr>
<th>S</th>
<th>AM</th>
<th>1234</th>
<th>L</th>
<th>-</th>
<th>-</th>
<th>-</th>
<th>-</th>
</tr>
</thead>
<tbody>
<tr>
<td>First letter of patient first name</td>
<td>First 2 letters of patient last name</td>
<td>Unique number assigned by Zimmer</td>
<td>Operated side (Left/Right)</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

**PSI Case ID with 15 Characters**

**EXAMPLE: SAM123L77DD13US**

<table>
<thead>
<tr>
<th>S</th>
<th>AM</th>
<th>123</th>
<th>L</th>
<th>77</th>
<th>DD</th>
<th>13</th>
<th>US</th>
</tr>
</thead>
<tbody>
<tr>
<td>First letter of patient first name</td>
<td>First 2 letters of patient last name</td>
<td>Unique number assigned by Zimmer</td>
<td>Operated side (Left/Right)</td>
<td>Year of patient birthday</td>
<td>Surgeon initial</td>
<td>Year when the case created</td>
<td>Region where the case ID created</td>
</tr>
</tbody>
</table>

*Fig. 1*

PSI Case Identifier

**Warning:** If the Case ID markings do not match the patient, do not use the PSI Knee Instrument Guides and Bone Models on the patient. Notify your Zimmer representative immediately.
Intra-Operative Guide

The *Zimmer* PSI Knee Jigs are designed for use with conventional incision as well as the MIS Sub-Vastus, the MIS Mid-Vastus, and the MIS Medial-Parapatellar approaches for the placement of given *Persona* implant families (defined in the “Indication for Use” section). These surgical approaches are described in the following *Zimmer* Surgical Techniques:

- *Zimmer Persona* Surgical Technique (97-5026-001-00) for the *Persona* implant family.
Femur Exposure

- Expose the femur and tibia according to the applicable surgical technique (Fig. 2).
- Remove soft tissues on the bone that could prevent a good contact with the PSI jigs, such as meniscus and fat tissue.
- Do not remove any osteophytes or cartilage from the femur.

Note: If the bone model significantly differs from the actual anatomy in those regions, it is indicated not to use the femoral PSI jig.
Position the Femoral PSI Jig

- Position the PSI jig on the distal femur by first locking on the anterior ridge of the femur and then applying pressure distally to secure the fit (Fig. 3). Avoid rotating the jig towards the posterior condyles, as this would cause excessive flexion.

- Use the visual cues on the jig indicating the mechanical axis entry point, Whiteside’s Line and the transepicondylar axis, to help position the PSI jig and decide if proper alignment is achieved.

Note: If the PSI jig does not have the appropriate snug fit, if there is any doubt on the jig position, or if the marking on the PSI jig does not match the anatomic landmarks, be sure that no soft tissue interferes between the PSI jig and the bone. The positioning of the PSI jig can be double checked on the optional bone model. If the above conditions remain, do not insert pins or drill holes and revert to standard surgical technique. At this point, intramedullary instrumentation should be used.
Pin the Distal Cutting Guide Pin Holes

- Hold the PSI jig in position by hand, and pin the medial and lateral distal cutting guide pin holes on the PSI jig using the standard instrument accessory 3.2mm x 75 mm Persona Trocar Tipped Drill Pins (2.5 hex) (00-5901-020-00) or use the 3.2 Headless Trocar Drill Pin (20-8000-00-16) (Fig. 4). Insert pins using the Persona Pin/Screw Inserter (00-5901-021-00).
Drill 4-in-1 Cutting Guide Pin Holes

- Using the 3.2mm drill bit, available through Zimmer Standard Instrumentation, drill the medial and lateral 4-in-1 pin holes of the PSI jig deep enough to ensure that after the distal cut, the holes are still visible (Fig. 5).

Note: If the drill contacts a trocar pin, DO NOT drill further and remove the pin.
Resect Distal Femur

- Remove the Femoral PSI jig by sliding it off the pins, leaving the distal cutting guide pins in place (Fig. 6).
- In case the Femoral PSI jig gets locked over the bone during its retrieval, it is recommended to disengage one pin at a time to ease the removal of the PSI jig. If the pins are removed in the process, re-insert them in the pin holes after having removed the jig.

- Secure the Persona 0° Captured/Uncaptured Cutting Head (42-5099-010-00) in the holes marked ‘0’ (Fig. 7).

- Check alignment, if desired, and make the distal cut (Fig. 8).
Locate the 4-in-1 Cut Guide Pin Holes

- Remove the medial and lateral distal 3.2mm x 75mm Headless Trocar Tipped Drill Pin with the Persona Multi Pin Puller (00-5901-022-00) (Fig. 9).
- Locate the 4-in-1 pinholes.
Place 4-in-1 Femoral Finishing Guide

- By hand, place the Persona 4-in-1 Cut Guide (42-5099-085-54 for Anterior Referencing or 42-5099-044-54 for Posterior Referencing), on the femur by aligning the two pins on the back of the guide with the previously drilled positioning holes. (Fig. 10).
- Impact the face of the guide until it is flush with the femur.
- Refer to the Zimmer Persona Surgical Technique (97-5026-001-00) for complete instruction.

Note: The instrumentation and related size per the planning can be found in the pre operative Surgical Planning Report.

Warning: The appropriate Anterior or Posterior Persona Cut Guide has to be used in accordance with the pre-operative Surgical Planning Report. In case of any doubt revert to the standard surgical technique.
Position Tibial PSI Jig

- Look at the mating surfaces of the Tibial PSI jig on the tibia bone model or on the pre-operative planning (Fig. 11).
- Remove soft tissues on the bone that could prevent good contact with the PSI Jig, such as meniscus and fatty tissue.
- Do not remove osteophytes or cartilage from the tibia.

To position the Tibial PSI jig, first ensure good medial contact between the jig and the bone, confirming that the medial side of the jig is properly wrapped around the bone. Then, press the two arms perpendicular on the plateau and then jig as a whole to maintain proper placement and full contact with the bone. Avoid rotating the jig by pressing too strongly anteriorly (Fig. 12).

Note: If the representation of the bone on the planning record or the optional Bone Models significantly differs from the actual anatomy in those regions, it is indicated not to use the Tibial PSI Jig.

Note: If the PSI jig does not mate appropriately, or if there is any doubt on the baseplate position or the marking on the PSI jig of the medial third of the tubercle does not match the anatomic landmarks, make sure that no soft tissue interferes between the PSI jig and the bone. The position of the PSI Jig can also be double checked on the optional Bone Model. If the above conditions persist, DO NOT insert pins or drill holes and revert to standard surgical technique. Remove the PSI jig from the assembly and set the baseplate orientation and rotation on the tibial cut as per standard surgical technique.
Verify Tibial PSI Jig Alignment

- Insert the *Persona* Drop Rod Adaptor (20-8014-015-00) to help position the PSI jig. Make sure the Drop Rod Adaptor pushed until it is flush with the PSI jig. Also make sure the Drop Rod Adaptor is inserted on the proper side by using the left or right laser marking *(Fig. 13)*.

- The Drop Rod Adaptor slot lines up with two landmarks, the PCL insertion point and the medial 1/3 of the tibial tubercle *(Fig 14)*.

- Insert Alignment Rod (00-5785-080-00) through *Persona* Drop Rod Adaptor to verify alignment of the PSI Guide *(Fig. 15)*. Alignment rod should point towards the center of the malleoli.
Pin Tibia Cut Guide Pin Holes

- Hold the PSI jig by and pin the medial and lateral tibia cut guide pin holes of the PSI jig using 3.2mm x 75mm Persona Trocar Tipped Drill Pins (2.5 hex) (00-5901-020-00) with the Persona Pin/Screw Inserter (00-5901-021-00) (Fig. 16).

Note: Avoid applying excessive force anteriorly to the PSI Tibial Jig to prevent adding anterior slope.

Note: The PSI drop rod adaptor can stay in place while pinning the PSI Tibia jig (Fig. 16).
Remove Tibial PSI Jig

- Remove the PSI Tibial jig gently to avoid pulling the pins out. Verify that both pins are still placed in the drilled holes (Fig 17).
- In case the PSI Tibial jig gets locked over the bone during its retrieval, it is recommended to disengage the medial pin first, and if the jig is still locked over the pins, remove the lateral pin. If pins have been removed, re-insert them in the holes after the jig is removed.

Note: Avoid pulling too hard on the jig, as this can damage the drilled pin hole, possibly causing misalignment.
Resect Proximal Tibia

- Following the standard technique, align the Persona Tibial Cut Guide Left/Right - 3° or 7° (42-5399-051-03, 42-5399-051-07, 42-5399-052-03, 42-5399-052-07) in place on the bone in the holes marked '0' (Fig. 18).

- Insert a 3.2mm Trocar Tipped Drill Pin in the oblique hole indicated by a lock pin symbol, to further secure the cut guide (Fig. 19).

- Verify the alignment of the Persona Tibial cut Guide by inserting the Persona Alignment Rod with coupler (00-5785-080-00) together with the Persona Drop Rod Adaptor (Fig. 20).

- Use a 1.27mm (.050-inch) oscillating saw blade through the slot of the Captured Cute Guide to resect the proximal surface of the tibia (Fig. 21).

Instruments

Persona Tibial Cut Guide Right - 3° 42-5399-052-03
Persona Tibial Cut Guide Right - 7° 42-5399-052-07
Persona Tibial Cut Guide Left - 3° 42-5399-051-03
Persona Tibial Cut Guide Left - 7° 42-5399-051-07
Persona Drop Rod Adapter 42-5399-006-00
Optional: Install PSI Tibia Rotation on Persona Sizing Plate Handle

- Insert the PSI Tibia Rotational Guide on the Persona Tibial Sizing Plate Handle (42-5399-017-00) by sliding the handle on the open side of the PSI Tibia Rotational Guide (Fig 22).

- Attach the proper tibial Sizing Plate, as defined in the pre-operative plan (Fig. 23).
Set Tibial Rotation

• Slide the tibial sizing and rotation assembly, described in the previous step, on the tibial cutting guide pins (Fig. 24).
• Mate the PSI Rotational Guide on the anterior surface of the tibia to assess the planned rotation and bone cut coverage.

Note: If the PSI Jig does not mate appropriately or if there is any doubt on the baseplate position or the marking on the PSI Jig of the medial third of the tubercle does not match the anatomic landmarks make sure that no soft tissue interferes between the PSI Jig and the bone. If the above conditions remain DO NOT insert pins or drill hole and revert to standard surgical technique. Remove the PSI Jig from the assembly and set the baseplate orientation and rotation on the tibial cut as per standard surgical technique.

• Insert 25mm x 3.2 mm (2.5mm female hex) screws or 25mm x 3.2mm Persona Shorthead Holding pins in the medial and lateral holes near the PCL cutout of the Persona Cemented Tibial Sizing Plate, to secure the tibial baseplate with the Persona Multi Pin Puller (00-5901-022-00) (Fig. 25).
• Remove the PSI Rotational Guide by pressing the lever and pulling off the handle.
Verify Overall Alignment

- Insert the drop rod, available from Zimmer Standard Instrumentation, into the Tibial Sizing Plate Handle (Fig. 26).
- When the alignment has been verified remove the PSI Rotational Guide by pushing on its clipping mechanism and pulling back the PSI Rotation jig. Remove the handle as per the standard surgical technique.
## Supported Zimmer Persona Systems

<table>
<thead>
<tr>
<th>Femur</th>
<th>Tibia</th>
</tr>
</thead>
<tbody>
<tr>
<td>CR Cemented Standard</td>
<td>Stemmed Cemented Tibia</td>
</tr>
<tr>
<td>CR Cemented Narrow</td>
<td>C, D, E, F, G, H, J</td>
</tr>
<tr>
<td>PS Cemented Standard</td>
<td>TM Porous</td>
</tr>
<tr>
<td>PS Cemented Narrow</td>
<td>C, D, E, F, G, H</td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td>3, 4, 5, 6, 7, 8, 9, 10, 11</td>
<td></td>
</tr>
</tbody>
</table>
Cleaning/Sterilization Methods and Equipment Inventory

- Disposable Zimmer PSI Knee Instruments are provided non-sterile and are single use. They must be cleaned and sterilized by the end user before the surgery. The Reusable Zimmer PSI Knee Instruments must be cleaned after use and prior to sterilization.
- The instruments should not be sterilized in the protective bag or packaging supplied with them. All sterilizations should be performed using standard and regularly maintained equipment.
- In the case a surgery is re-scheduled or in the case of another issue requiring the Zimmer Jigs to be re-cleaned and re-sterilized, the Disposable Zimmer PSI Knee Instruments can only be re-cleaned and re-sterilized once for a given patient, if they have not been otherwise contaminated. This is to avoid patient infection and contamination. Validated cleaning methods have not been established for such re-use conditions. Cleaning and Sterilization methods are described below.

Warning: Before every surgery, the user must verify that all jigs (including bone models) and instruments have been cleaned and sterilized.

Cleaning

For cleaning, both the single use Zimmer PSI Knee Jigs (including bone models) and the reusable instruments require manual cleaning steps as follows (additional component-specific cleaning instructions are provided in the next subsections):

1. Pre-soak components in an enzyme solution.
2. Scrub components with a soft bristle brush to remove all visible soil.
3. Use a water jet to flush difficult access areas and closely mated surfaces (see areas labeled “A” in the images in the following tables: “Reusable Zimmer PSI Knee Instruments” and “Zimmer PSI Knee Disposable Kits”).
4. Ultrasound clean (Sonication) all components in an enzyme solution with a minimum cycle time of 5 minutes.
5. Thoroughly rinse and dry all components.
Sterilization Parameters

- All components (disposable and reusable) require steam sterilization before use per the following methods (Fig. 27).

### Steam Sterilization (Autoclave)

<table>
<thead>
<tr>
<th>Cycle Type</th>
<th>Temperature $^1$</th>
<th>Exposure Time $^1$</th>
<th>Minimum Dry Time $^2$</th>
<th>Minimum Cool Time $^1$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pre-Vacuum</td>
<td>132°C (270°F)</td>
<td>4 minutes</td>
<td>30 minutes</td>
<td>30 minutes</td>
</tr>
</tbody>
</table>

1 Both the given cycle temperature and time can be increased to 134°C + 3°C (273.2°F + 5.4°F) and 18 minutes according to local requirements outside of the United States such as in the European Union.

2 Drying times vary according to load size and should be increased for larger loads.

3 Cooling times vary according to the type of sterilizer used, device design, temperature and humidity of ambient environment, and type of packaging used. Cooling process should comply with ANSI/AAMI ST79.

---

### Reusable Zimmer PSI Knee Instruments and Additional Specific Cleaning Instructions

- The table below (Fig. 28) shows the reusable instruments for Persona Zimmer PSI Jigs Kit. Additional specific cleaning instructions as applicable to each instrument are provided.

<table>
<thead>
<tr>
<th>Catalog No.</th>
<th>Instrument</th>
<th>Qty</th>
<th>Sterilization and specific cleaning instructions</th>
<th>Additional Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>20-8014-015-00</td>
<td><em>Persona</em> Drop Rod Adaptor</td>
<td>1</td>
<td>Autoclave Additional specific cleaning requirements: Use a water jet to flush difficult access areas (see areas labeled “A”)</td>
<td>Re-usable, Provided non-sterile</td>
</tr>
</tbody>
</table>
Zimmer PSI Knee Disposable Kits

The table below shows the available Disposable Zimmer PSI Knee Instruments. Additional specific cleaning instructions as applicable to each component are provided (Fig. 29).

<table>
<thead>
<tr>
<th>Catalog No.</th>
<th>Instrument</th>
<th>Qty</th>
<th>Sterilization and specific cleaning instructions</th>
<th>Additional Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>20-8070-001-01</td>
<td>Zimmer PSI Knee Persona Jigs Aref</td>
<td>1</td>
<td>Autoclave</td>
<td>Single use, Provided non-sterile</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Additional specific cleaning requirements : Use a water jet to flush difficult access areas (see areas labeled “A”)</td>
<td></td>
</tr>
<tr>
<td>20-8070-001-02</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>20-8070-002-01</td>
<td>Zimmer PSI Knee Persona Jigs Aref &amp; Tibia Rotation</td>
<td>1</td>
<td>Autoclave</td>
<td>Single use, Provided non-sterile</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Additional specific cleaning requirements : Use a water jet to flush difficult access areas (see areas labeled “A”)</td>
<td></td>
</tr>
<tr>
<td>20-8070-002-02</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Fig. 29
<table>
<thead>
<tr>
<th>Catalog No.</th>
<th>Instrument</th>
<th>Qty</th>
<th>Sterilization and specific cleaning instructions</th>
<th>Additional Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>20-8070-010-01</td>
<td><em>Zimmer PSI Knee Persona Jigs PREF</em></td>
<td>1</td>
<td>Autoclave</td>
<td>Single use, Provided non-sterile</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Additional specific cleaning requirements : Use a water jet to flush difficult access areas (see areas labeled “A”)</td>
<td></td>
</tr>
<tr>
<td>20-8070-010-02</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>20-8070-011-01</td>
<td><em>Zimmer PSI Knee Persona Jigs PREF &amp; Tibia Rotation</em></td>
<td>1</td>
<td>Autoclave</td>
<td>Single use, Provided non-sterile</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Additional specific cleaning requirements : Use a water jet to flush difficult access areas (see areas labeled “A”)</td>
<td></td>
</tr>
<tr>
<td>20-8070-011-02</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Fig. 29 (continued)
### Catalog No. Instrument Qty Sterilization and specific cleaning instructions Additional Notes

<p>| | | | | |</p>
<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>20-8070-009-00</td>
<td><em>Zimmer</em> PSI Knee Bone Models</td>
<td>1</td>
<td>Autoclave</td>
<td>Single use, Provided non-sterile</td>
</tr>
</tbody>
</table>

### Standard Instrumentation Pins:

<p>| | | | | |</p>
<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>00-5901-020-00*</td>
<td>Headless Trocar Drill Pin, 75mm</td>
<td>2</td>
<td>See package insert for re-sterilization instruction if permissible.</td>
<td>Single use, Provided sterile</td>
</tr>
</tbody>
</table>

- or -

<p>| | | | | |</p>
<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>20-8000-000-16</td>
<td>3.2mm Headless Trocar Drill Pin</td>
<td>2</td>
<td>Autoclave</td>
<td>Single use, Provided non-sterile</td>
</tr>
</tbody>
</table>

*The Headless trocar pin (00-5901-020-00) is manufactured by *Zimmer* (not *Zimmer CAS*). It should be ordered directly from *Zimmer*.

**Warning:** Do not use pins or any other fasteners than those recommended above.

**Fig. 29 (continued)**
### Reusable Instruments Ordering

- In order to perform a PSI case, some key reusable instruments have to be part of the kit. The Zimmer PSI is compatible with Persona Standard Instrumentation.
- A list of instruments that are required for each type of implant is listed in Figure 30, only one of these instruments is required per case for all types of implants. The Zimmer division responsible of supplying the instrument is written in the last row.
- To order a Zimmer instrument, please place your order through DCS.

<table>
<thead>
<tr>
<th>Implant/Instruments</th>
<th>20-8014-014-00 PRI Drop Rod Adaptor</th>
<th>20-8014-015-00 Persona Drop Rod Adaptor</th>
<th>00-5901-021-00 Trocar Screw Pin Driver</th>
<th>00-5901-075-00 PRI 0° Left Cut Guide or-00-5901-076-00 PRI 0° Right Cut Guide</th>
<th>00-5901-086-00 PRI alignment adapter</th>
<th>Key Remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td>Persona</td>
<td>X</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Do not use NexGen Tibial cut guide</td>
</tr>
<tr>
<td>Supplier</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Zimmer, Warsaw</td>
</tr>
</tbody>
</table>

*Fig. 30*
Zimmer Contact Information

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

Manufacturer
Zimmer CAS
75, Queen Street, Suite 3300
Montreal (Quebec) H3C 2N6
CANADA
Tel: 1 (514) 395-8883
Fax: 1 (514) 878-3801
Web site: www.zimmer.com
Email: ZimmerPSI@zimmercas.com

Customer Support
Tel: 1 (866) 336-7846

European Community (EC) Representative
Zimmer U.K. Ltd.
9 Lancaster Place
South Marston Park
Swindon, SN3 4FP, UK
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.