Zimmer Biomet Hip Preservation Portfolio
Zimmer Biomet Hip Preservation - a new opportunity to treat the hip joint.

Today, preserving the hip joint often equates to performing a hip arthroscopy or replacement. At Zimmer Biomet, we view all conditions leading up to osteoarthritis as an opportunity to preserve the native anatomy. From dysplasia and avascular necrosis, to femoroacetabular impingement and labral damage, we provide conservative treatment options to make it simple for surgeons to restore a healthy hip.
Patient Specific Analysis
3D Motion Hip Analysis Service, powered by Clinical Graphics

Hip Arthroscopy

1. JuggerKnot® Long Soft Anchor System
2. Quattro® GL Suture Anchor (US Only)
3. SureLock® All-Suture Anchor (US Only)
4. SpeedSnare™ Suture Passer System
Hip Subchondroplasty Procedure

The Subchondroplasty® Procedure targets and fills subchondral defects in the acetabulum and femur including cysts, bone marrow lesions and bone defects associated with early stage pre-collapse AVN.²

Surgical Head Dislocation (SHD)

DeNovo® NT (Natural Tissue) Graft

JuggerKnot® Soft Anchor-2.9 mm/2.9mm with Needles
Core Decompression

BioCUE® Blood and Bone Marrow Aspirate (BBMA) Concentration System³

PerFuse™ Percutaneous Decompression System⁴

Products are not available in all markets.
Please contact your local sales representative for product availability.
## Osteotomy

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<th>Product Description</th>
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<td>Large Titanium Cannulated Screw System (5.0 mm and larger screws)</td>
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<td><strong>Zimmer</strong>® <strong>Natural Nail</strong>® System (Antegrade and Cephalomedullary options)</td>
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1. Quattro & SureLock Suture Anchors have not been approved for Hip procedures in the European Union. Please refer to the IFU for indications and contraindications.

2. AccuFill® Bone Substitute material is indicated to fill bone voids or defects of the skeletal system that are not intrinsic to the stability of the bony structure, and has not been evaluated for any clinical indications.

3. The BioCUE System is indicated to prepare autologous platelet-rich plasma (PRP) to mix with autograft and/or allograft bone prior to application to an orthopedic site. The PRP has not been evaluated for any clinical indications.

4. Core decompression is an avascular necrosis treatment option. The PerFuse System is designed to access the femoral head for core decompression; and to facilitate mixing/pre-mixing of bone graft material with I.V. fluids, blood, platelet-rich plasma, bone marrow, or other specified blood components.

5. Screws 5 mm and larger are intended for use in the femur; but only screws that are 6.5 mm or larger in diameter are indicated for both the pelvis and femur.

6. The DHS and BP (sterile) are intended for temporary fixation of fractures or osteotomies in the proximal or distal femur, and are not available in the US.

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