References

- Zimmer ZTR_BI_0245_18.
- Zhang, Y., et al. Interfacial Frictional Behavior: Cancellous Bone, Cortical Bone, and a Novel Porous Tantalum Biomaterial. Journal of Musculoskeletal Research. 3(4):, 245-251, 1999.
- 3. Bobyn, J.D., et al. Characteristics of Bone In-growth and Interface Mechanics of a New Porous Tantalum Biomaterial. Journal of Bone and Joint Surgery (British), 81-B(5): 907, 1999.
- Shirazi-Adl, A., et al. Experimental Determination of Friction Characteristics at the Trabecular Bone / Porous-coated Metal Interface in Cementless Implants. The Journal of Biomedical Research. 27: 167-175, 1993.
- Zimmer ZRR_WA_2409_11. Zimmer ZRR_WA_2403_11. 5.
- 7. Zimmer ZRR_WA_2537_12.
- Zimmer ZTR_BI_0249_18.
- Zimmer ZRR_WA_2551_12. 9.
- 10 Zimmer ZTR BI 0133 18.
- 11. NexGen Legacy Constrained Condylar Knee release to market date 1997.
- Natural Knee II Revision Knee System release to market date 2003. 12.
- 13. Vanguard SSK Revision Knee System release to market date 2005.
- Vanguard 360 Revision Knee System release to market date 2012. 14.
- 15. Peters, C.L., et al. Clinical and Radiographic Results of 184 Consecutive Revision Total Knee Arthroplasties Placed with Modular Cementless Stems. The Journal of Arthroplasty. 24(6): 48-53, 2009.
- Maynard, L. M., et al. Survival of Primary Condylar-Constrained Total Knee Arthroplasty at a Minimum of 7 Years. The Journal of Arthroplasty. 29(6): 1197-1202, 2014.
- Kim, J.H., et al. Revision Total Knee Arthroplasty with Use of a Constrained Condylar Knee Prosthesis. Journal of Bone Joint Surgery (America). 91(6): 1440-1447, 2009.
- Kim, Y.H., et al. Long-Term Clinical Outcomes and Survivorship of Revision Total Knee Arthroplasty with Use of a Constrained Condylar Knee Prosthesis. The Journal of Arthroplasty. 30(10): 1804-1809, 2015.
- Dai, Y., et al. Anatomical Tibial Component Design Can Increase Tibial Coverage and Rotational Alignment Accuracy: A Comparison of Six Contemporary Designs. Knee Surgery Sports Traumatology Arthroscopy. 22: 2911-2923; KSSTA. 2014.
- 20. Zimmer ZRM_WI_5438_18.
- Morgan-Jones, J. et al. Zonal Fixation in Revision Total Knee Arthroplasty. Bone 21. and Joint Journal, 97-B: 147-9, 2015.
- 22. Persona Revision Femoral and Tibial Cone and Sleeve Size Range and Shape. Memorandum. KN000253. 2018.
- 23. Zimmer ZTR_BI_0262_18.
- 24. FRA 160.40 Stem Extension Transition Length and FRA 160.50 Stem Extension Lead In Length Memorandum. KN000253. 2018.
- Dalury, D., et al. Why Are Total Knee Arthroplasties Being Revised? The Journal 25. of Arthroplasty. 28(8): 120-121, 2013.
- Sharkey, P.F., et al. Why Are Total Knee Arthroplasties Failing Today-Has 26. Anything Changed After 10 Years? The Journal of Arthroplasty. 29(7): 1774-1778, 2014,
- Chang, M.J., et al. Diagnosis, Causes and Treatments of Instability Following Total Knee Arthroplasty. Knee Surgery and Related Research. 26(2): 61-67,
- Zimmer ZTR_BI_0223_18. 28.

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- ZIMMER BIOMET Instrumentation Industrial Design Style Guideline. Revised 2015.09.02.
- 30. Levine, B. et al. Experimental and Clinical Performance of Porous Metal Tantalum in Orthopedic Surgery. Biomaterials. 27: 4671-81, 2006.
- 31. Bobyn, J.D., et al. Characterization of New Porous Tantalum Biomaterial for Reconstructive Orthopaedics. Scientific Exhibition. 66th Annual Meeting of the American Academy of Orthopaedic Surgeons. Anaheim, CA. 1999.
- 32. Karageorgiou V, Kaplan D. Porosity of biomaterial scaffolds and osteogenesis Biomaterials. 26: 5474-5491, 2005.
- Springer, B. CORR Insights®: The Alpha-defensin Test for Periprosthetic Joint Infection Responds to a Wide Spectrum of Organisms. Clinical Orthopaedics and Related Research. 473: 2236-7, 2015.
- 34. Deirmengian, C., et al. Diagnosing Periprosthetic Joint Infection. Has the Era of the Biomarker Arrived? Clinical Orthopaedics and Related Research.
- CDD-CLI-001: Clinical Validation of CD Diagnostics Synovasure PJI ELISA Test and Synovasure PJI Lateral Flow Test for Detection of Periprosthetic Joint Infection (PJI) in Synovial Fluid.
- Malchau, H., et al. Prognosis of Total Hip Replacement. The National Hip Arthroplasty Register. 9-11, 1996.
- 37. Malchau, H., et al. The Swedish Total Hip Replacement Register. Journal of Bone and Joint Surgery. 84A: 2-20, 2002.
- Swedish Hip Arthroplasty Register. Annual Report 1998.
- Kavanaugh A, et al. Factors Influencing the Initial Strength of the Tibial Tray-Cement Interface Bond. Bone Joint Journal. 95-B(34): 98, 2013.
- Refsum, A. M., et al. Cementing Technique for Primary Knee Arthroplasty: A Scoping Review. Acta Orthopaedica. Published online: Aug 27, 2019. DOI: 10.1080/17453674.2019.1657333
- 41. Wang, J.S., et al. Bone Cement Pososity in Vacuum Mixing Systems. Bone Cements and Cementing Technique. Walenkamp, Murray (Eds) Springer Verlag, 2001.
- Dunne, N.J., et al. Influence of the Mixing Techniques on the Physical Properties of Acrylic Bone Cement. Biomaterials. 22: 1819-26, 2001.
- Parvizi, J., et al. Efficacy of Antibiotic-impregnated Cement in Total Hip Replacement. A Meta-Analysis. Acta Orthopaedica (Scandinavica). 79(3): 335-341, 2008.
- Kühn, K.D. Bone Cements, Up-to-Date Comparison of Physical and Chemical Properties of Commercial Materials. 254-258. Springer Verlag: Berlin, 2000.
- Data on File at Zimmer Biomet. Verification Report, Release of antibiotics. Project Verification Number CD0278. June 20, 2017.
 - *Laboratory studies are not necessarily indicative of clinical performance.

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Biomet France SARL Plateau de Lautagne 26000 Valence, France Persona Revision Knee System



Legal Manufacturer

Zimmer, Inc. 1800 W. Center Street Warsaw, Indiana 46580 USA







THE POWER TO PERSONALIZE™

PERSONALIZED. PRECISE. PROVEN.

Personalized implants, precise instrumentation and proven technologies enable surgeons with the ability to personalize their surgical experience to best meet the needs of each patient.

Persona Revision offers anatomic implant geometry to provide patients an individualized fit while achieving the desired bone coverage, fixation¹ and stability. The intuitive instrumentation platform empowers surgeons with the flexibility to tailor their surgical approach intraoperatively, while the streamlined delivery system's modularity is designed to help increase OR efficiencies and workflow. Combined with the clinically successful Trabecular Metal™ Material²-⁴ and Vivacit-E® Highly Crosslinked Polyethylene,⁵-¹0 Persona Revision truly offers personalization for revision knee arthroplasty.



THE PERSONA REVISION KNEE SYSTEM ENABLES SURGEONS WITH

THE ABILITY TO PERSONALIZE

REVISION KNEE ARTHROPLASTY.



WHEN SOMETHING WORKS WELL, WHY CHANGE IT?

Built on a legacy of clinical heritage, components of the Persona Revision Knee System are evolutionary refinements of their respective predecessors: Persona the Personalized Knee, NexGen® Legacy Constrained Condylar Knee, Natural-Knee® II, and Vanguard SSK/ 360 Revision Knee System. These design and material properties have a combined clinical history of over 55 years on the market. 11-14 By integrating the successful design elements from its predecessors, Persona Revision was born with an updated implant design, modernized instrumentation platform and advanced technologies.





TRABECULAR METAL TECHNOLOGY

- Over 20 years of clinical history²⁻⁴
- Well documented in over 350+ peer reviewed journal articles, poster exhibits and abstracts

VANGUARD®SSK/360 REVISION KNEE SYSTEMS

• 93% survivorship at 4 years¹⁵



VIVACIT E® VITAMIN-E HIGHLY CROSSLINKED POLYETHYLENE

Actively stabilized with Vitamin-E to help protect against oxidation and maintain wear resistance and strength throughout the life of the implant. Exceptional oxidative stability,^{5,6} ultra-low wear^{7,8} and enhanced strength.^{9,10}

NEXGEN® LEGACY® CONSTRAINED CONDYLAR KNEE (LCCK)

- 97.6% survivorship at 10+ years in a primary knee arthroplasty¹⁶
- 96% survivorship at 10+ years in a revision knee arthroplasty¹⁷
- 92% survivorship at 16 years in a revision knee arthroplasty¹⁸



PERSONALIZED IMPLANTS

ACHIEVING AN INDIVIDUALIZED FIT.

When designing the Persona Revision Implants, the ultimate goal was to create the most anatomic revision system on the market. During the design process, extensive research was performed using the ZiBRATM Anatomical Modeling System which studied the morphology of native tibias of various ethnicities, genders and sizes.

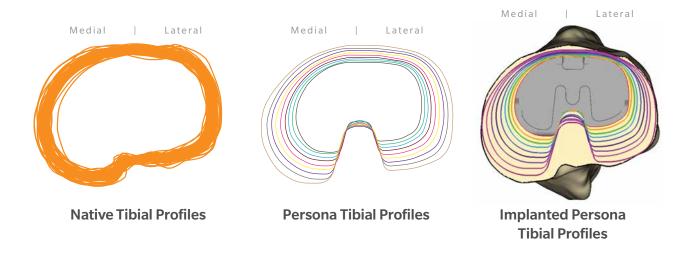
Data showed previous asymmetric and symmetric tibial designs once served a purpose, but surgeons had to choose between rotation, coverage and position.¹⁹ The anatomic implant geometry of the Persona Revision Tibia should help achieve proper rotation, position and optimal bone coverage.

Persona Revision offers surgeons an array of anatomic components designed with patient anatomy in mind. The system provides tibial and femoral cones and numerous stem choices to address fixation needs while the femoral components and multiple bearing constraint options aim to restore soft tissue balance and stability. With these advancements, Persona Revision can offer patients a truly individualized implant fit.



Anatomic Implants

The Persona Primary and Revision Tibial Implants were designed by studying the morphology of native tibias of various ethnicities, genders and sizes. Hundreds of virtual tibial resections were performed using the ZiBRA Anatomic Modeling System with varying surgical parameters. This thorough research helped to better understand that the variation of the tibial shape was only subtle between ethnicities and gender. It was determined that the ideal size and shape of the tibial implant should be anatomic.



Tibial Component

The shape of the Persona Primary Tibial Implant was incorporated into the anatomic design of the Persona Revision Tibial Implant. This consistent design provides a unified system approach giving surgeons the confidence to address both difficult primary and complex revision knee arthroplasties. In vitro, the Persona Anatomic Tibia has demonstrated:

- 92 percent bone coverage with proper rotation¹⁹
- Less compromise of coverage (0.5 percent anatomic vs 5 percent non-anatomic)19
- Six percent average improvement in coverage compared to non-anatomic designs¹⁹



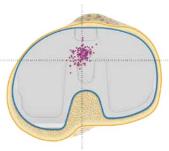
Medialized Tibial Keel

The ZiBRA Anatomic Modeling System database was also utilized to determine proper stem housing location. By observing the location of the native diaphysis compared to the outer tibial implant shape, an anterior medialized keel was determined to be the most anatomically accurate location.²⁰

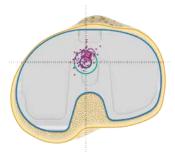
ZiBRA data was also used to determine that the 3 mm and 6 mm offset stems would help achieve optimal coverage and rotation for almost all ZiBRA model data resulting in a desired fit as shown in the graphic below.²⁰



Average Tibial Canal Locations*



Tibial Baseplate Location in Relation to Canal Data Points



3 and 6 mm Offset Stem Coverage in Relation to Tibial **Canal Data Points**



Keel Location on the Underside of the Persona Revision Tibial **Baseplate**

Persona Revision Tibial Implant Specifications

- Nine left and right anatomic micro, macro and core sizes (A-J)
- Anatomic disproportional M/L growth
- Compatible with all Persona Stems (tapered, smooth and splined designs)
- Triple wedge bearing locking mechanism
- No through holes to prevent osteolytic pathways
- Made of Tivanium® Alloy (titanium)
- Compatible with anatomic Trabecular Metal and Tivanium Tibial Augments to minimize bone overhang



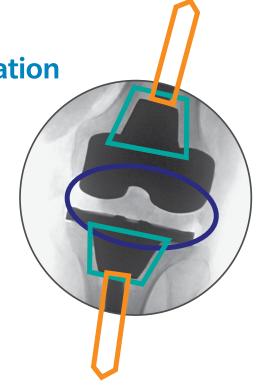
Splined Stem



*Based on ZiBRA Anatomical Modeling System Data

Achieving Zonal Fixation

One study shows that in order to achieve robust implant fixation, most successful knee revision procedures require a combined approach to fixation with a multi-zone strategy. There are three anatomical zones; the joint surface (epiphysis), metaphysis and diaphysis. To provide a stable construct, two of the three zones must achieve adequate fixation. Persona Revision offers numerous anatomic tibial and femoral cones, augments and an array of stem options to address fixation needs.



Zone 3 – Diaphysis

Zone 2 - Metaphysis

Zone 1 - Joint Surface (Epiphysis)

Zone 2 – Metaphysis

Zone 3 – Diaphysis

Anatomic Tibial and Femoral Trabecular Metal Cones

At times while performing revision TKAs, surgeons encounter bone defects and poor bone quality that make it challenging to restore the patient's natural knee. "The achievement of solid fixation of revision implants is essential to allow early post-operative mobilization and rehabilitation, and improves the longevity of the construct". 21 Persona Revision Trabecular Metal Cones provide customers a solution to effectively fill bone defects to closely replicate patients normal anatomy. The proven 2-4 Trabecular Metal material offers increased biological fixation and distributes loads closer to the joint line.

- Anatomic shapes designed to fill bone voids without cortical impingement²²
- Designed to fill cavitary bone defects and provide a stable platform for femoral and/or tibial articulating components²³
- Cones are designed to facilitate biologic fixation and promote a load sharing construct



Tibial Central Cone



Tibial Perimeter Cone



Femoral Central Cone



Femoral Metaphyseal Cone



Enhanced Stem Design

Boasting one of the most robust stem portfolios on the market, the Persona Revision Knee System presents customers an array of stem configurations in multiple lengths and sizes to address varying patient anatomy.

- 135 mm and 175 mm lengths designed to achieve the desired diaphyseal engagement
- Offset splined stem design promotes precise femoral and tibial component placement
 - o Monolithic offset splined stem design reduces the potential for fretting and corrosion
 - Shortened offset stem transition closer to the joint line results in a better fitting implant with less bone removal²⁴ and area to fill with cement
- Tapered connection for ease of implant removal over threaded type of stem connections

Stem Implant Specifications

- **Tapered Stems**
 - +30 mm* and +75 mm lengths
 - 11 mm and 14 mm diameters
 - *Additional compatibility with Persona Primary 5 Degree Tibial Component
- **Smooth Stems**
 - +135 mm and +175 mm lengths 10-24 mm diameters in 2 mm increments
- **Splined Stems**
 - 0 mm (straight), 3 mm and 6 mm offset configurations
 - +135 mm and +175 mm lengths
 - 10–18 mm diameters in 1 mm increments
 - 20-24 mm diameters in 2 mm increments

Note: + in addition to stem housing



Restoring Soft Tissue Balance

One of the most common causes of implant failure is joint instability. ²⁵⁻²⁷ In an effort to address joint instability, Persona Revision's multiple femoral components in varying sizes, along with a vast continuum of bearing constraints provide surgeons the tools to help achieve a stable knee.

Femoral Components

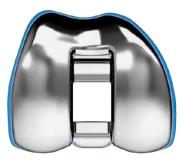
Persona Revision offers both standard and plus femoral component options. The plus femoral components feature a Flexion Fill[™] design, with an additional +3 mm of posterior condylar offset to provide additional flexion stability compared to the corresponding standard size. This addresses flexion laxity without the need to upsize, offset or augment the femur to accommodate soft tissue needs while not affecting implant overhang.





The plus femoral components can also be used to achieve a narrow femoral option. To achieve a narrow femoral option, the size smaller plus femoral component can be used to fill the same flexion space with a smaller M/L dimension.

Standard and Plus Femurs as a Narrow Option



M/L Femoral Size Comparison Blue Outline: Size 7 Standard Femur

Grav: Size 5 Plus Femur





A/P Femoral Size Comparison

Femoral Implant Specifications

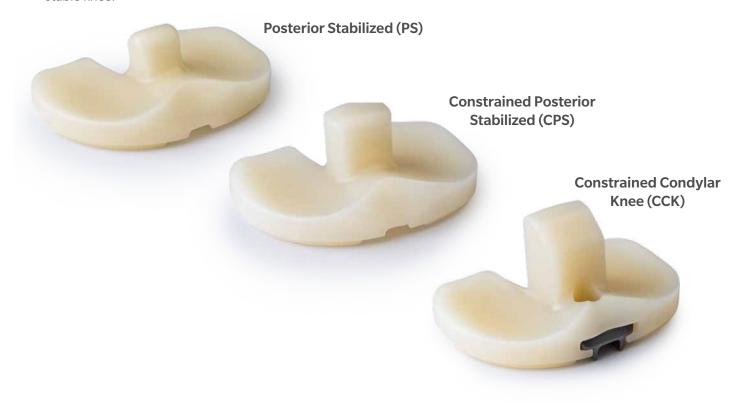
- 13 left and right sizes available in two distinct profiles, standard and plus*
- Designed to accommodate up to 135° of flexion
- Compatible with PS, CPS and CCK Bearings
- Compatible with the Persona Primary 5 Degree Stemmed Tibial Component with a 14mm x +30 mm stem extension
- Compatible with Trabecular Metal and Tivanium distal and posterior augments available in sizes 5, 10 and 15 mm**

**15 mm augments are not available for femoral sizes 1 and 3 All augment sizes can be used together in any configuration

*Plus femur not available in size 13

Continuum of Constraint

The Persona Revision Knee System includes a full continuum of constraint offering Posterior Stabilized (PS), Constrained Posterior Stabilized (CPS) and Constrained Condylar Knee (CCK) Bearings available in Vitamin-E and Conventional polyethylene*. All bearings are compatible with the Persona Primary 5 Degree Stemmed Tibia and Persona Revision Tibial Components. From no constraint to full constraint, the system helps achieve a balanced, stable knee.



Bearing Implant Specifications

- PS Bearing offered in 1 mm increments**
- CPS and CCK Bearings offered in 2 mm increments
- CCK Bearings available in thicknesses of 10-26 mm
- CCK Bearing Lock Down Screw is designed to prevent torque loss²⁸ to improve screw backout resistance
- CCK Bearing offers a highly constrained option
- Patella friendly chamfer designed to help reduce impingement in high flexion
 - * Conventional polyethylene available in PS Only ** PS 1 mm 10-14 mm and 2 mm 16-20 mm

PRECISE INSTRUMENTATION

WITH A PERSONALIZED APPROACH.

Patient outcomes can be driven by the precision and accuracy of each step within the surgical procedure. In order to streamline workflow and provide a more personalized approach to instrumentation selection, the surgical techniques allow surgeons to individualize the surgical experience based on the patient's bone and joint condition.

The instrumentation allows surgeons to intraoperatively transition from a primary to a revision component using a condensed revision conversion set. This conversion instrumentation provides the flexibility to make efficient surgical decisions based on the specific needs of the patient.

These advancements along with updated instrumentation allow for simplified decisions tailored to surgical preference, philosophy and approach to achieve a personalized procedure.



Comprehensive Instrument Platform

Persona Revision Instrumentation was designed to be versatile in its capabilities and philosophies, precise in its measurements, comprehensive and comfortable with repetitive use. These options allow for more control, a smooth surgical flow and reproducible outcomes.

Instrumentation Design

- Multipurpose instrumentation design reduces excess tray weight and redundant instrumentation
- Anatomic contouring and ergonomic mechanical grip feature of the instruments designed to maximize comfort and control over repetitive use²⁹
- Weight balanced design helps minimize strain and allow for improved reproducibility and accuracy²⁹
- Proprietary surface finish reduces intraoperative glare and provides an enhanced grip in the surgical environment²⁹
- Enables postponement of bone cuts until flexion and extension gaps have been determined for trial early philosophy
- Lockable connections for cutting instruments designed to eliminate pin use and improve accuracy of cuts



TASP System

The Tibial Articular Surface Provisional (TASP) adds sophistication to the trialing system.

- Seamless soft tissue balancing with no need to remove the provisional
- Effortlessly make 2 mm adjustments designed to find the optimal bearing thickness every time
- Available in PS, CPS and CCK constraints



Personalized Surgical Techniques

The adaptable instrumentation and personalized surgical workflow accommodates surgeon's preferred philosophy or patient anatomy. The surgical techniques are capable of supporting various surgeon defined workflows such as:

- Standard Method
- Zonal Fixation (cones)
- Trial Early Method
- Primary Conversion
- Short Cemented Stem



Streamlined Delivery System

Revision systems have historically been bulky, containing redundant instrumentation that takes up tray space and clutters the OR and central sterile department leading to inefficiencies. Persona Revision's efficient kitting strategy with modular brackets aims to eliminate nonessential instrumentation and allows for adaptable tray configurations based on surgeon defined workflow and hospital preference to help reduce the burden of inventory management.

 Allows for personalization of trays based on procedure or surgical philosophy

 Simplified back-table set-up saving space and clutter in the sterile field

 Configured to reduce instrument footprint while providing the full array of implant offerings





Condensed Femoral Revision Set

Persona Revision provides surgeons a three tray condensed femoral set to address difficult primary arthroplasty. The primary to revision conversion instrumentation allows for a seamless transition from a primary to a stemmed revision femoral component with a CCK bearing intraoperatively based on patient soft tissue and fixation needs. This concise set offers an efficient solution by reducing the number of typical revision trays required in the OR to back up a primary case.





PROVEN TECHNOLOGY

BUILT ON CLINICAL PERFORMANCE.

Persona Revision boasts the most clinically proven²⁻⁴ biological fixation portfolio on the market with over 20 years of successful clinical use. Combined with our premium bearing technology, Persona Revision Knee System provides customers with solutions to improve patient outcomes.





Trabecular Metal Technology

For more than 20 years, Trabecular Metal Technology has clinically demonstrated excellent initial stability and biologic in-growth not only in knees, but in many orthopedic applications. ^{2-4, 30} This highly porous material is designed to replicate the structure, function and physiological properties of cancellous bone.³¹

Trabecular Metal Technology is available in multiple sized femoral and tibial cones, femoral and tibial augments and patellar implants giving surgeons the power to select the right implant to best meet the needs of each patient.

- Made from commercially pure elemental Tantalum
- Cancellous architecture up to 80% porosity with a 100% openinterconnected cell structure with an average pore size of 440 microns designed to support bony in-growth and vascularization^{31,32}
- Published 0.98 coefficient of friction* against cancellous bone for initial stability³²







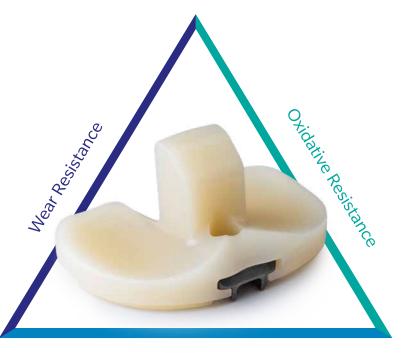
 $[\]hbox{``For net-shaped Trabecular Metal Material.}$

Vivacit-E Vitamin-E Highly Crosslinked Polyethylene (HXPE)

Patients are expecting more out of their implants, and with that in mind, Persona Revision Bearings were created exclusively out of Vivacit-E Vitamin-E Highly Crosslinked Polyethylene. Vivacit-E Polyethylene is actively stabilized with Vitamin-E to help protect against oxidation and maintain wear resistance and strength throughout the life of the implant.

- Stresses in the CCK Bearing Spine are lower than that of the PS Bearing Spine which demonstrated a 10% improvement in spine fatigue strength over conventional polyethylene^{14, 15*}
- Exceptional oxidative stability with delamination resistance and retention of mechanical properties 12 times longer than industry standards^{10, 11*}
- Vivacit-E Highly Crosslinked Polyethylene (HXPE) results in a technologically advanced material that provides mechanical strength, superior oxidative stability and wear resistance over other polyethylenes^{10-15*}
- Ultra-low wear with 96 percent wear reduction compared to conventional polyethylene and 73 percent wear reduction compared to re-melted HXPE polyethylene^{12, 13*}

Vivacit-E Vitamin-E Highly Crosslined Polyethylene



Strength

 $^{{}^*\}text{Laboratory testing not necessarily indicative of clinical performance}.$

BREAKING THE REVISION CYCLE

It is time to break the revision cycle and focus on the entire patient journey. From diagnosis to patient specific re-implantation, we unite customizable, interconnected and interdependent services and solutions to address each unique episode of care.







Care

Uniting Innovative Solutions



Therapy



Patient Specific Solutions



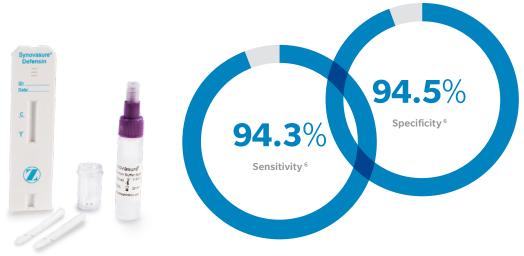
Limb Salvage



Re-implantation

Infection Diagnostic Solution

Diagnosis of Periprosthetic Joint Infection (PJI) continues to be a major challenge for orthopedic surgeons, especially in late and delayed infections.^{33, 34} Synovasure[®] Alpha Defensin Lateral Flow Test is the only in-vitro diagnostic device specifically designed to aid in the diagnosis of PJI by rapidly and accurately* detecting alpha defensin biomarkers in synovial fluid.



^{*}Synovasure Alpha Defensin Lateral Flow Test is 94.3% sensitive and 94.5% specific against the standard of care criteria, excluding samples diluted with >20% blood (RBC>1,000,000)³⁵

Microorganism Lavage Solution

Bacteria can produce Extracellular Polymeric Substance (EPS) to shield themselves from both mechanical and chemical attack. Bactisure™ Wound Lavage* was specifically designed to remove these structurally resistant forms of bacteria by physically deconstructing the EPS matrix, making bacteria more susceptible to traditional antibiotics and the body's normal defense mechanism.





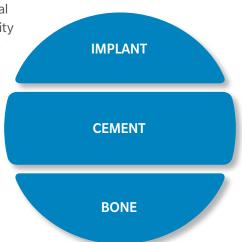
Bactisure Wound Lavage is intended to be used with jet lavage (Zimmer Pulsavac* Plus or Pulsavac Plus AC) for cleansing and removal of debris, including microorganisms, from wounds.

^{*}Bactisure Wound Lavage is not CE marked.

Modern Cementing Technique (MCT) - improved clinical outcome³⁶⁻³⁸

Modern Cementing Technique Knee (MCT Knee) addresses implant loosening with the objective to provide long term implant stability in knee arthroplasty and is incorporated into the Persona Revision Surgical Technique. The objective of MCT in facilitating long term implant stability is to optimize cement quality and the interfaces between both Implant-Cement and Cement-Bone.^{39*,40}

Zimmer Biomet offers solutions for standardized mixing procedures resulting in homogeneous cement with very low porosity. The mixing and collection under vacuum reduces the cement's porosity, which can lead to improvements in cement strength and fatigue life. 41*, 42*



Antibiotic-loaded Bone Cement - may reduce risk of re-infection⁴³

Gentamicin has shown to be the antibiotic of choice for bone cement, as its broad therapeutic spectrum covers gram-positive and gram-negative bacteria. Gentamicin is bactericidal and its release from the bone cement is superior to that of other antibiotics.⁴⁴

Zimmer Biomet bone cements including gentamicin, where available, have shown to provide antibiotic release over several days.^{45*}





* * Refobacin® Revision is not available in the United States.

* Laboratory testing is not necessarily indicative of clinical performance.

** Refobacin® Revision is not available in the United States.

StageOne™ Knee Cement Spacer Molds