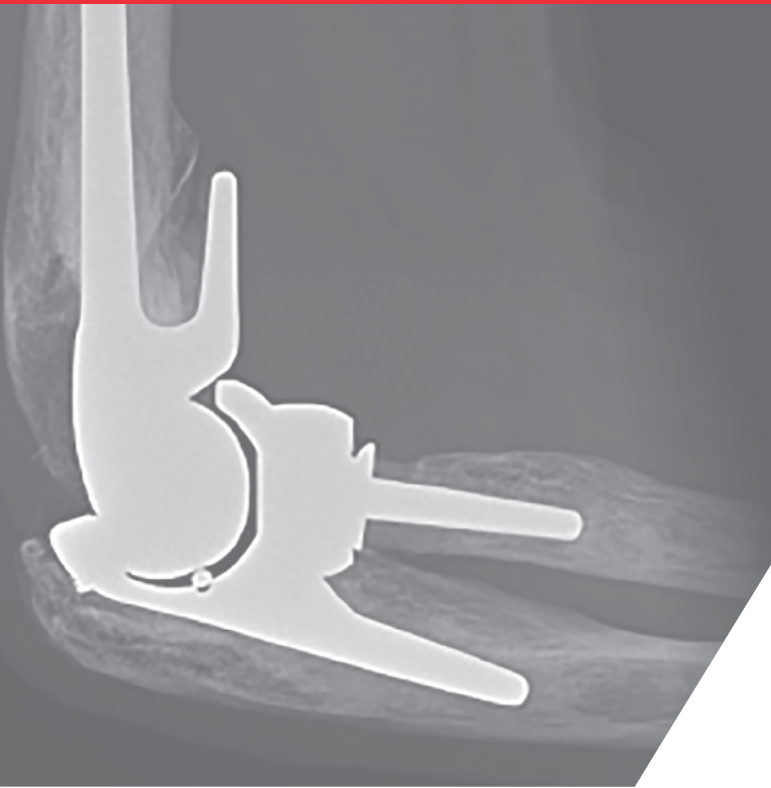


TORNIER
LATITUDE EV™
Total Elbow Arthroplasty



Welcome to the **EVOLUTION** of elbow replacement



The LATITUDE EV™ total elbow arthroplasty system is designed to replicate the natural anatomy of the elbow. The innovative LATITUDE™ gives surgeons the ability to reproduce the natural flexion/extension axis and restore natural kinematics of the elbow with its anatomic design. The LATITUDE EV implant design is founded on the original LATITUDE implant which has been in clinical use since 2001.

Reproduce the anatomy...

LATITUDE EV™ is designed to model the patient's anatomy allowing the surgeon to restore the natural kinematics of the elbow, while offering maximum flexibility in elbow arthroplasty.

| Design Feature | Advantage |
|--|---|
| Intraoperative flexibility to link or unlink the implant (a change in linkage can be performed at any time) | Flexible options to address a wide range of indications and patient needs |
| The LATITUDE EV provides multiple sizes of radial head implants for articulation with the anatomic humeral spool | Allows for optimal mechanical balance of radiohumeral articulation |
| Ti Plasma Spray | Enhances long term fixation |

Anatomic design to restore normal anatomy for optimal range of motion

Modeled after the human anatomy, LATITUDE EV reproduces natural elbow kinematics with an anatomic design



...simply restore the natural kinematics of the elbow.



Humeral Spool

Concave barrel shaped trochlea preserves linear contact through 7° varus/valgus movement with ulnar component

Ulnar Stem

Shape replicates natural bow of the ulna to aid in ease of insertion and reduce stresses on the cortical wall

Radial Head

Radial Head

60% of comprehensive loads are transferred across the radiohumeral joint. Morrey et al, JBS 70-A 1988. LATITUDE™ is the only total elbow prosthesis that enables surgeons to maintain the radiohumeral joint when alignment is adequate.

Humeral Stem

Square shape for rotational stability

Anterior Flange

For bone graft

Medial/Lateral Fins

Prevent rotation

Ulnar Cap Option

To enable use as a linked or unlinked implant





User Friendly, Intuitive Instrumentation

The LATITUDE EV™ system offers new, completely redesigned instrumentation. The instrumentation system allows for accurate component positioning with humeral, ulnar, and radial preparations that are separate but codependent.



LATITUDE EV™ Total Elbow Product Specifications

| Humeral Spools | | Ulnar Stems | | Ulnar Caps | |
|----------------------|-----------------------|-------------|-----------------------|-------------------------------|---|
| DKY211 | Small Right | 0030010 | Small Short Right | DKY067 | Small |
| DKY212 | Small Left | 0030011 | Small Short Left | DKY068 | Medium |
| DKY213 | Medium Right | 0030110 | Medium Short Right | DKY069 | Large |
| DKY214 | Medium Left | 0030111 | Medium Short Left | Radial Heads and Stems | |
| DKY215 | Large Right | 0030210 | Large Short Right | DKY056 | Small Head |
| DKY216 | Large Left | 0030211 | Large Short Left | DKY057 | Medium Head |
| DKY217 | Large+ Right | 0030020 | Small Standard Right | DKY058 | Large Head |
| DKY218 | Large+ Left | 0030021 | Small Standard Left | DKY059 | Large+ Head |
| Humeral Stems | | 0030120 | Medium Standard Right | DKY061 | 5.0 mm Stem |
| 0030302 | Small Standard Right | 0030121 | Medium Standard Left | DKY062 | 6.5 mm Stem |
| 0030303 | Small Standard Left | 0030220 | Large Standard Right | Cement Restrictors | |
| 0030402 | Medium Standard Right | 0030221 | Large Standard Left | EBO101 | Cement Restrictor (Diameter Range 8-15 mm) |
| 0030403 | Medium Standard Left | 0030030 | Small 125 mm Right | EBO102 | Cement Restrictor (Diameter Range 5-8 mm) |
| 0030502 | Large Standard Right | 0030031 | Small 125 mm Left | Single Use Items | |
| 0030503 | Large Standard Left | 0030130 | Medium 125 mm Right | DKY090 | Single Use Suture Passer |
| 0030312 | Small 150 mm Right | 0030131 | Medium 125 mm Left | DWD060 | 3 mm Drill Bit |
| 0030313 | Small 150 mm Left | 0030230 | Large 125 mm Right | | |
| 0030412 | Medium 150 mm Right | 0030231 | Large 125 mm Left | | |
| 0030413 | Medium 150 mm Left | 0030040 | Small 150 mm Right | | |
| 0030512 | Large 150 mm Right | 0030041 | Small 150 mm Left | | |
| 0030513 | Large 150 mm Left | 0030140 | Medium 150 mm Right | | |
| 0030322 | Small 200 mm Right | 0030141 | Medium 150 mm Left | | |
| 0030323 | Small 200 mm Left | 0030240 | Large 150 mm Right | | |
| 0030422 | Medium 200 mm Right | 0030241 | Large 150 mm Left | | |
| 0030423 | Medium 200 mm Left | | | | |
| 0030522 | Large 200 mm Right | | | | |
| 0030523 | Large 200 mm Left | | | | |

Indications:

LATITUDE EV is intended for total elbow arthroplasty. Prosthetic replacement with this device may be indicated to relieve severe pain or significant disability following the effects of primary or secondary osteoarthritis and rheumatoid arthritis; correction of functional deformities; revision procedures where other treatments or devices have failed; treatment of fractures that are unmanageable using other techniques. LATITUDE EV is intended for cemented use only.

Contraindications:

Systemic infection is an absolute contraindication. Every effort should be made to rule out the possibility of preoperative sepsis in patients who have one or more of the following abnormalities: fever and/or local inflammation; rapid joint destruction or bone resorption apparent on roentgenograms; elevation of sedimentation rate unexplained by other disease; elevation of WBC count; distant foci of infection from genitourinary, pulmonary, skin and other sites, dental focus infection which may cause hematogenous spread to the implant site; skeletally immature patients; cases where there is inadequate neuromuscular status, poor bone stock, or poor skin coverage around the elbow joint that would make the procedure unjustifiable; neuromuscular or psychiatric disorders which might jeopardize fixation and postoperative care; known allergy to one of the materials; pregnancy.

LATITUDE EV™ has been designed in conjunction with:

Graham King, MD (University of Western Ontario); Shawn O'Driscoll, MD, PhD (Mayo Foundation); Ken Yamaguchi, MD (Washington University)

Proper surgical procedures and techniques are the responsibility of the medical professional. This material is furnished for information purposes only. Each surgeon must evaluate the appropriateness of the material based on his or her personal medical training and experience. Prior to use of any Tornier implant system, the surgeon should refer to the product package insert for complete warnings, precautions, indications, contraindications and adverse effects. Package inserts are also available by contacting Wright. Contact information can be found in this document and the package insert.



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