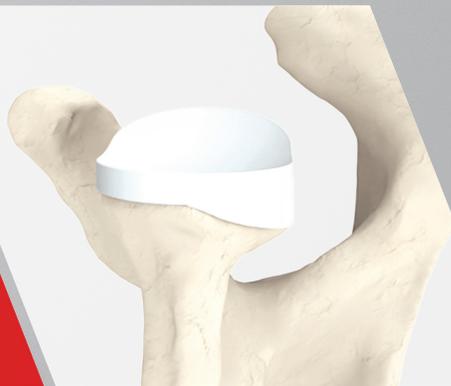




TORNIER
BLUEPRINT™
3D Planning + PSI



TORNIER
AEQUALIS™ PERFORM™ +
Shoulder System



PLAN, POSITION,
PREPARE & PRESERVE

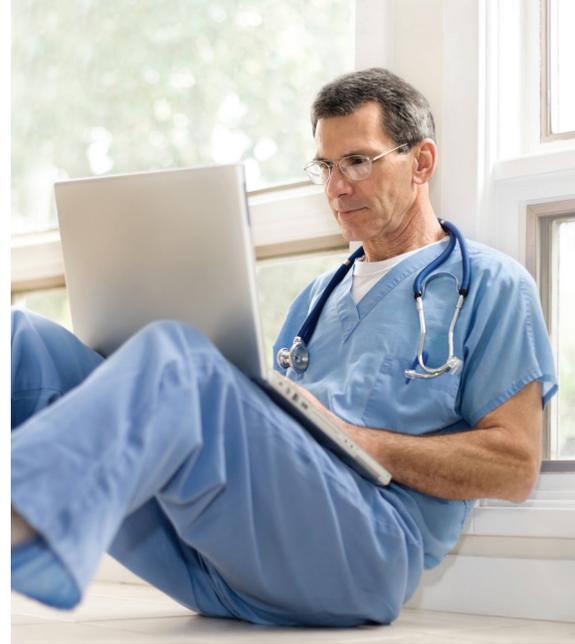
PLAN

Clarity from Complexity

3D Evaluation of a 3D Problem

Several studies^{1,2,3} have shown that 2D evaluations regularly underestimate retroversion by up to 15° in A1 and B2 glenoids compared to 3D measurements.

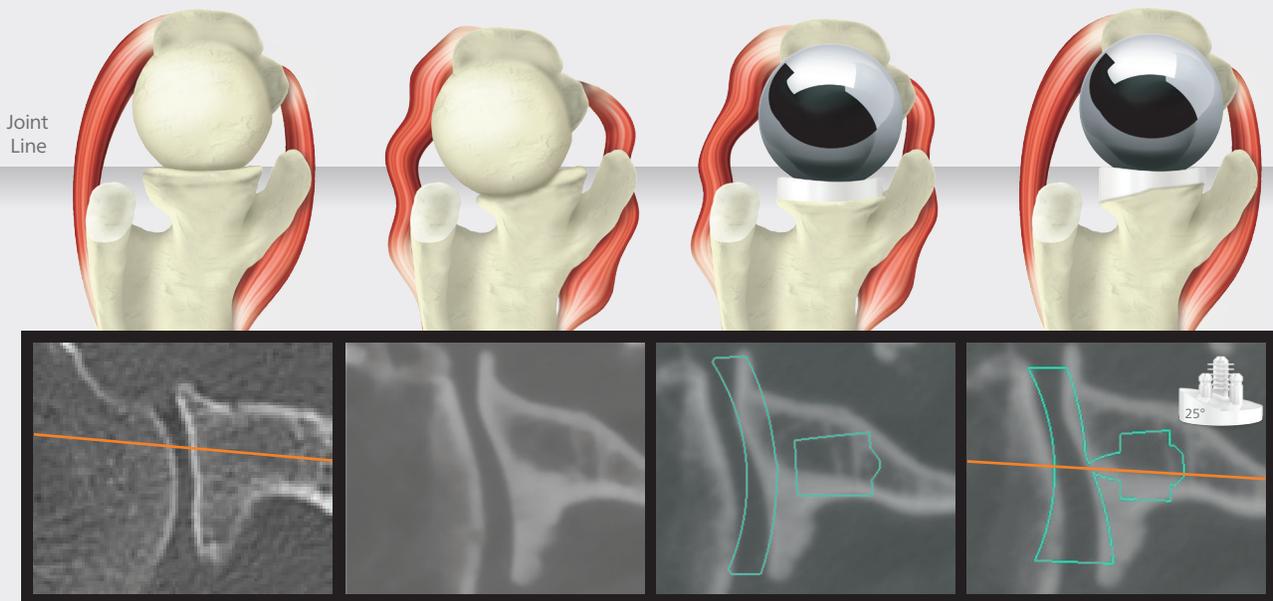
BLUEPRINT™ 3D Planning software automatically segments CT scan series in real time, on your own computer to generate a precise 3D reconstruction with associated 3D version and inclination calculations.



POSITION

Restores Joint Line, Corrects Version & Re-Centers⁴

Precise Positioning with BLUEPRINT 3D Planning



Normal Shoulder

Joint Line Compromised
Posterior wear resulting in 13° of pre-operative retroversion and posterior subluxation.

Standard Glenoid with Eccentric Reaming
Extensive bone removal and joint line medialization.

Joint Line Restored with Aequalis™ Perform™+
Minimal reaming, bone preservation, and version correction.



PREPARE

Accurate Execution of the Ideal Plan

Instrument Technology that Facilitates Accurate Bone Preparation

BLUEPRINT patient-specific instrumentation (PSI) **precisely** transfer the pre-operative plan to the OR.



The MARKSMAN reamer provides **axial control** for preparation of the "Paleo" surface.

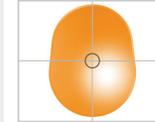


The angled NEO REAMER preserves critical subchondral bone by independently preparing the "Neo" surface on-axis **with ease**.

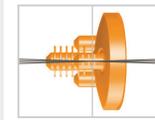


Proven Accuracy with BLUEPRINT PSI

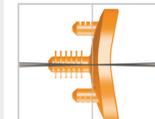
BLUEPRINT PSI



1.05 mm entry point



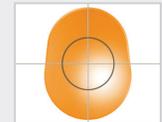
1.64° version



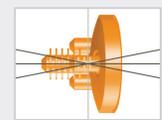
1.42° inclination

G. Walch et al. JSES, Vol. 24, Issue 2, p302-309

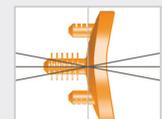
Standard Instrumentation



2.9 mm entry point



11.1° version



10.7° inclination

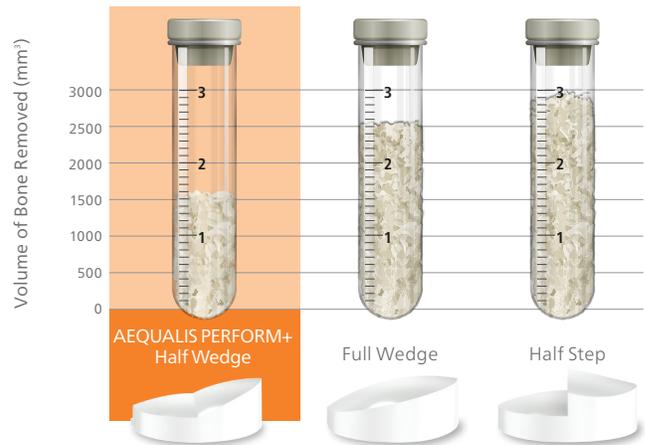
J. Iannotti et al JBJS, Vol 96-A, Number 9, May 7, 2014.

PRESERVE

More Bone, More Support

Defect Mimicking Implant Preserves Subchondral Bone⁵

The AEQUALIS PERFORM+ Shoulder System was developed to address posterior glenoid deficiencies, that when treated with traditional implants have demonstrated an increased risk for glenoid loosening via finite element analysis.⁶ The "defect mimicking" augment shape was developed to preserve subchondral bone which has been demonstrated to be a critical factor in long-term survivorship.⁷ In an independent head-to-head comparison conducted via virtual implantation in CAD, the posterior wedge shape removed substantially less bone than the other designs, with the remaining bone being of better quality.⁵



References

- 1 Hoenecke H et al. Accuracy of CT-based measurements of glenoid version for total shoulder arthroplasty. J Shoulder Elbow Surg. 2010 Mar;19(2):166-71.
- 2 Armstrong A et al. Comparison of standard two-dimensional and three-dimensional corrected glenoid version measurements. J Shoulder Elbow Surg. 2011 Jun;20(4):577-83
- 3 Farron A et al. Measurements of three-dimensional glenoid erosion when planning the prosthetic replacement of osteoarthritic shoulders. Bone Joint J. 2014 Apr;96-B(4):513-8.
- 4 Iannotti J et al. Correction of acquired glenoid bone loss in osteoarthritis with a standard versus an augmented glenoid component. J. Shoulder Elbow Surg (2013)
- 5 Athwal G et al. Augmented glenoid component designs for type B2 erosions: a computational comparison by volume of bone removal and quality of remaining bone. J Shoulder Elbow Surg (2015)
- 6 Juan C. Hermida, MD; Cesar Flores-Hernandez, BS; Heinz R. Hoenecke, MD; Darryl D. D'Lima, MD, PhD. Augmented wedge-shaped glenoid component for the correction of glenoid retroversion: a finite element analysis. J Shoulder Elbow Surg (2014) 23, 347-354
- 7 R. Sean Churchill, MD, Edwin E. Spencer Jr, MD, Edward V. Fehring, MD. Quantification of B2 glenoid morphology in total shoulder arthroplasty. J Shoulder Elbow Surgery. 2015; 24(8)



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