

**K120044 ARTHREX UNIVERSII XL GLENOID-PEGGED,
ARTHREX UNIVERS II XL GLENOID-KEELED**Mar 8, 2012
63 days to decisionK120044 · Product code: **KWS** · Orthopedic
Source: <https://www.510kdatabase.net/k120044/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Prosthesis, Shoulder, Semi-constrained, Metal/polymer Cemented (KWS)
Date received	Jan 5, 2012
Decision date	Mar 8, 2012
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Arthrex, Inc.
Location	Naples, FL, US
Contact	COURTNEY SMITH
Website	https://www.arthrex.com
510(k) history	346 submissions · 342 cleared · 1992-2026

Arthrex, Inc. is a medical device manufacturer based in Naples, US. The company specializes in surgical implants and instruments for orthopedic procedures. Arthrex has received FDA 510(k) clearances from total submissions since its first clearance in 1992. The company's portfolio is dominated by Orthopedic devices, which represent 89% of its regulatory submissions. Recent cleared devices include suture anchors, plating systems, nails, and specialized fixation devices for shoulder, ankle, and lower extremity procedures. The latest FDA 510(k) clearance was received in 2026,...

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Device record: <https://www.510kdatabase.net/k120044/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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