

K040500 ARTHROTEK LACTONAILMay 25, 2004
89 days to decisionK040500 · Product code: **HTY** · Orthopedic
Source: <https://www.510kdatabase.net/k040500/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pin, Fixation, Smooth (HTY)
Date received	Feb 26, 2004
Decision date	May 25, 2004
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Arthrotek, Inc.
Location	West Hartford, CT, US
Contact	TRACY J BICKEL
510(k) history	17 submissions · 17 cleared · 1987-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k040500/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026