

**K051827 JOIMAX ENDOSCOPE/MULTISCOPE, THESSYS
FORAMINOSCOPE/MULTISCOPE, THESSYS
LAMINOSCOPE/MULTISCOPE**Aug 12, 2005
37 days to decisionK051827 · Product code: **HRX** · Orthopedic
Source: <https://www.510kdatabase.net/k051827/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Jul 6, 2005
Decision date	Aug 12, 2005
Days to decision	37 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Joimax GmbH
Location	Los Gatos, CA, US
Contact	KARIN GASTINEAU
510(k) history	7 submissions · 7 cleared · 2005-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k051827/>. 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 26, 2026