

**K990345 RABEA DEVICE, MODEL PXXXXXX**Jul 30, 1999  
176 days to decisionK990345 · Product code: **JDK** · Orthopedic  
Source: <https://www.510kdatabase.net/k990345/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - U
Submission type	Traditional
Device classification	Prosthesis, Hip, Cement Restrictor (JDK)
Date received	Feb 4, 1999
Decision date	Jul 30, 1999
Days to decision	176 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Signus Medizintechnik GmbH</b>
Location	Minneapolis, MN, US
Contact	THOMAS HOGHAUG
510(k) history	23 submissions · 22 cleared · 1999-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k990345/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 21, 2026