FDA 510(k) Premarket Notification

K990345 RABEA DEVICE, MODEL PXXXXXX

CLEARED

Jul 30, 1999
176 days to decision

K990345 · Product code: **JDK** · Orthopedic
Source: https://www.510kdatabase.net/k990345/

SUBMISSION DETAILS

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

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