FDA 510(k) Premarket Notification

K982006 MICOMED - HALM ZIELKE INSTRUMENTATION

CLEARED

Jan 20, 1999
226 days to decision

K982006 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k982006/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Jun 8, 1998
Decision date	Jan 20, 1999
Days to decision	226 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Micomed GmbH**
Location	Fullerton, CA, US
Contact	CHARMAINE HENDERSON
510(k) history	2 submissions · 2 cleared · 1999-2000