FDA 510(k) Premarket Notification

K991882 NEXFLEX TOTAL KNEE SYSTEM

CLEARED

Dec 8, 1999
189 days to decision

K991882 · Product code: **JWH** · Orthopedic
Source: https://www.510kdatabase.net/k991882/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Jun 2, 1999
Decision date	Dec 8, 1999
Days to decision	189 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Nexmed, Inc.**
Location	Palm Desert, CA, US
Contact	JASON BLAIN
510(k) history	6 submissions · 5 cleared · 1995-2004

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k991882/ 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 20, 2026