FDA 510(k) Premarket Notification

K952629 NEXFLEX TOTAL HIP SYSTEM

CLEARED

Sep 8, 1995
92 days to decision

K952629 · Product code: **JDI** · Orthopedic
Source: https://www.510kdatabase.net/k952629/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Jun 8, 1995
Decision date	Sep 8, 1995
Days to decision	92 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k952629/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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