FDA 510(k) Premarket Notification

K011832 VANGUARD REPROCESSED FEMORAL COMPRESSION DEVICE

CLEARED

Dec 21, 2001
192 days to decision

K011832 · Product code: **NMF** · Cardiovascular
Source: https://www.510kdatabase.net/k011832/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clamp, Vascular, Reprocessed (NMF)
Date received	Jun 12, 2001
Decision date	Dec 21, 2001
Days to decision	192 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vanguard Medical Concepts, Inc.**
Location	Plant City, FL, US
Contact	MIKE SAMMON
510(k) history	33 submissions · 33 cleared · 1991-2005