FDA 510(k) Premarket Notification

K992053 AESCULAP VASCULAR INSTRUMENTS

CLEARED

Jan 13, 2000
209 days to decision

K992053 · Product code: **DXC** · Cardiovascular
Source: https://www.510kdatabase.net/k992053/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clamp, Vascular (DXC)
Date received	Jun 18, 1999
Decision date	Jan 13, 2000
Days to decision	209 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Aesculap, Inc.**
Location	Burlingame, CA, US
Contact	MARY ELLEN HOLDEN
510(k) history	207 submissions · 201 cleared · 1991-2025