FDA 510(k) Premarket Notification

K930758 DIAGNOSTIC INTRAVASCULAR CATHETER

CLEARED

Sep 24, 1993
224 days to decision

K930758 · Product code: **LYK** · Cardiovascular
Source: https://www.510kdatabase.net/k930758/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Angioscope (LYK)
Date received	Feb 12, 1993
Decision date	Sep 24, 1993
Days to decision	224 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Applied Medical Resources**
Location	Launa Hills, CA, US
Contact	JO STEGWELL
510(k) history	58 submissions · 58 cleared · 1992-2021