FDA 510(k) Premarket Notification

K931421 DISPOSABLE STEERABLE ANGIOSCOPE

CLEARED

Jun 16, 1993
86 days to decision

K931421 · Product code: **LYK** · Cardiovascular
Source: https://www.510kdatabase.net/k931421/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Angioscope (LYK)
Date received	Mar 22, 1993
Decision date	Jun 16, 1993
Days to decision	86 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Endovascular, Inc.**
Location	Costa Mesa, CA, US
Contact	KAREN U SALINAS
510(k) history	3 submissions · 3 cleared · 1993-1993