

K930758 DIAGNOSTIC INTRAVASCULAR CATHETERSep 24, 1993
224 days to decisionK930758 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k930758/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026