

K960917 PERCUGLIDEJul 2, 1996
118 days to decisionK960917 · Product code: **DRE** · Cardiovascular
Source: <https://www.510kdatabase.net/k960917/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, For Percutaneous Catheterization (DRE)
Date received	Mar 6, 1996
Decision date	Jul 2, 1996
Days to decision	118 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Scientific Device Manufacturer, LLC
Location	San Rafael, CA, US
Contact	RICHARD C BALL
510(k) history	6 submissions · 6 cleared · 1995-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k960917/> 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 June 29, 2026