FDA 510(k) Premarket Notification

K001694 GUIDANT RETRIEVER DEVICE

CLEARED

May 15, 2001
347 days to decision

K001694 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k001694/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jun 2, 2000
Decision date	May 15, 2001
Days to decision	347 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Guidant Cardiac and Vascular Surgery**
Location	Menlo Park, CA, US
Contact	BERNICE JURS
510(k) history	7 submissions · 7 cleared · 1997-2002