FDA 510(k) Premarket Notification

K831636 PERCUTANEOUS INTRODUCER KIT

CLEARED

Aug 24, 1983
93 days to decision

K831636 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k831636/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	May 23, 1983
Decision date	Aug 24, 1983
Days to decision	93 days
Third-party review	No

APPLICANT

Company	**Accu-Line**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1983-1983