FDA 510(k) Premarket Notification

K895932 PSG(TM) EDM(TM) INFUSION CATHETER

CLEARED

Jan 4, 1990
86 days to decision

K895932 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k895932/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Oct 10, 1989
Decision date	Jan 4, 1990
Days to decision	86 days
Third-party review	No

APPLICANT

Company	**Peripheral Systems Group**
Location	Mountain View, CA, US
Contact	JESSICA AYRES
510(k) history	17 submissions · 17 cleared · 1990-1994