FDA 510(k) Premarket Notification

K834352 ADULT VENT CATHETER

CLEARED

Jan 27, 1984
45 days to decision

K834352 · Product code: **DRA** · Cardiovascular
Source: https://www.510kdatabase.net/k834352/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Steerable (DRA)
Date received	Dec 13, 1983
Decision date	Jan 27, 1984
Days to decision	45 days
Third-party review	No

APPLICANT

Company	**Dlp, Inc.**
Location	Mchenry, IL, US
510(k) history	56 submissions · 56 cleared · 1979-1997