FDA 510(k) Premarket Notification

K813143 DESILET

CLEARED

Nov 27, 1981
22 days to decision

K813143 · Product code: **DYB** · Cardiovascular
Source: https://www.510kdatabase.net/k813143/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Nov 5, 1981
Decision date	Nov 27, 1981
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Exco, Inc.**
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1981-1985