FDA 510(k) Premarket Notification

K812143 INTRODUCER, PERCUTANEOUS

CLEARED

Aug 18, 1981
21 days to decision

K812143 · Product code: **DYB** · Cardiovascular
Source: https://www.510kdatabase.net/k812143/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Jul 28, 1981
Decision date	Aug 18, 1981
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**Willson**
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1980-1981