FDA 510(k) Premarket Notification

K914128 CONNECTING SET

CLEARED

Nov 27, 1991
75 days to decision

K914128 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k914128/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Sep 13, 1991
Decision date	Nov 27, 1991
Days to decision	75 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Procedure Products, Inc.**
Location	Mchenry, IL, US
Contact	ROBERT B EVERETT
510(k) history	16 submissions · 16 cleared · 1981-2017