FDA 510(k) Premarket Notification

K920157 SCHNEIDER 8 FR. STAMINA(TM) GUIDING CATHETER

CLEARED

Mar 16, 1992
62 days to decision

K920157 · Product code: **DXY** · Cardiovascular
Source: https://www.510kdatabase.net/k920157/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Jan 14, 1992
Decision date	Mar 16, 1992
Days to decision	62 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Schneider Intl., Ltd.**
Location	Minneapolis, MN, US
Contact	ROBERT L ULLEN
510(k) history	22 submissions · 22 cleared · 1989-1995