FDA 510(k) Premarket Notification

K921856 TOTAL CROSS .021 PERCUT. TRANSLUMINAL ANGIO. CATH.

CLEARED

Jul 21, 1992
95 days to decision

K921856 · Product code: **LIT** · Cardiovascular
Source: https://www.510kdatabase.net/k921856/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Angioplasty, Peripheral, Transluminal (LIT)
Date received	Apr 17, 1992
Decision date	Jul 21, 1992
Days to decision	95 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