FDA 510(k) Premarket Notification

K882238 VESSEL DILATOR AND INTRODUCER SHEATH

CLEARED

Aug 25, 1988
90 days to decision

K882238 · Product code: **DRE** · Cardiovascular
Source: https://www.510kdatabase.net/k882238/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, For Percutaneous Catheterization (DRE)
Date received	May 27, 1988
Decision date	Aug 25, 1988
Days to decision	90 days
Third-party review	No

APPLICANT

Company	**Ideal Medical, Inc.**
Location	Mchenry, IL, US
Contact	LORNA K LINVILLE
510(k) history	16 submissions · 14 cleared · 1981-1994