FDA 510(k) Premarket Notification

K881088 ADDITION OF A TIP GUARD TO THE ATHEROCATH(TM)

CLEARED

May 31, 1988
77 days to decision

K881088 · Product code: **LIT** · Cardiovascular
Source: https://www.510kdatabase.net/k881088/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Angioplasty, Peripheral, Transluminal (LIT)
Date received	Mar 15, 1988
Decision date	May 31, 1988
Days to decision	77 days
Third-party review	No

APPLICANT

Company	**Devices For Vascular Intervention, Inc.**
Location	Redwood City, CA, US
Contact	DIANE RUPPERT
510(k) history	14 submissions · 14 cleared · 1987-1994