FDA 510(k) Premarket Notification

K871546 SAFESIDE(TM) LUMED

CLEARED

Jun 17, 1987
58 days to decision

K871546 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k871546/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Apr 20, 1987
Decision date	Jun 17, 1987
Days to decision	58 days
Third-party review	No

APPLICANT

Company	**Luther Medical Products, Inc.**
Location	Walker, MI, US
Contact	RONALD B LUTHER
510(k) history	17 submissions · 16 cleared · 1980-1998