FDA 510(k) Premarket Notification

K842133 EYRTHROCATH DOUBLE-LUMEN CENTRAL VENOUS

CLEARED

Feb 19, 1985
265 days to decision

K842133 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k842133/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	May 30, 1984
Decision date	Feb 19, 1985
Days to decision	265 days
Third-party review	No

APPLICANT

Company	**Cardio Search, Inc.**
Location	Walker, MI, US
Contact	LARRY H BRIM
510(k) history	4 submissions · 4 cleared · 1983-1985