FDA 510(k) Premarket Notification

K892410 FOGARTY(R) THRU-LUMEN EMBOLECTOMY CATHETER

CLEARED

Jul 20, 1989
104 days to decision

K892410 · Product code: **DXE** · Cardiovascular
Source: https://www.510kdatabase.net/k892410/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Apr 7, 1989
Decision date	Jul 20, 1989
Days to decision	104 days
Third-party review	No

APPLICANT

Company	**Baxter Healthcare Corp**
Location	Mchenry, IL, US
Contact	J KRATTENMAKER
510(k) history	505 submissions · 496 cleared · 1977-2019