FDA 510(k) Premarket Notification

K911367 BAXTER MARATHON(TM) INTRAVASCULAR RELAY GUID CATH

CLEARED

Jun 10, 1991
74 days to decision

K911367 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k911367/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Mar 28, 1991
Decision date	Jun 10, 1991
Days to decision	74 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k911367/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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