FDA 510(k) Premarket Notification

K910208 BAXTER MARATHON(R) INTRAVASCULAR GUIDING CATHETERS

CLEARED

Apr 1, 1991
75 days to decision

K910208 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k910208/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jan 16, 1991
Decision date	Apr 1, 1991
Days to decision	75 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/k910208/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 17, 2026