FDA 510(k) Premarket Notification

K910429 SWAN-GANZ FLOW DIRECTED PRESSURE TD CATHETER

CLEARED

Jul 11, 1991
160 days to decision

K910429 · Product code: **DXO** · Cardiovascular
Source: https://www.510kdatabase.net/k910429/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Pressure, Catheter Tip (DXO)
Date received	Feb 1, 1991
Decision date	Jul 11, 1991
Days to decision	160 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k910429/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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