FDA 510(k) Premarket Notification

K901736 OXIMETRY PROBE CATHETERS

CLEARED

Sep 11, 1990
147 days to decision

K901736 · Product code: **DQE** · Cardiovascular
Source: https://www.510kdatabase.net/k901736/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Oximeter, Fiber-optic (DQE)
Date received	Apr 17, 1990
Decision date	Sep 11, 1990
Days to decision	147 days
Third-party review	No

APPLICANT

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