FDA 510(k) Premarket Notification

K900879 SOFT-WEDGE(TM) SYRINGE

CLEARED

May 10, 1990
73 days to decision

K900879 · Product code: **DYG** · Cardiovascular
Source: https://www.510kdatabase.net/k900879/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Flow Directed (DYG)
Date received	Feb 26, 1990
Decision date	May 10, 1990
Days to decision	73 days
Third-party review	No

APPLICANT

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