FDA 510(k) Premarket Notification

K882680 ALTERNATE BLOOD PORT DESIGN FOR ADDIT. MEMB. OF CA

CLEARED

Jul 22, 1988
23 days to decision

K882680 · Product code: **FJI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k882680/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, Capillary, Hollow Fiber (FJI)
Date received	Jun 29, 1988
Decision date	Jul 22, 1988
Days to decision	23 days
Third-party review	No

APPLICANT

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