FDA 510(k) Premarket Notification

K961040 BLOOD PORT CAP, DIALYSATE PORT CAP, RED HARD PLASTIC

CLEARED

Jun 11, 1996
89 days to decision

K961040 · Product code: **FJI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k961040/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, Capillary, Hollow Fiber (FJI)
Date received	Mar 14, 1996
Decision date	Jun 11, 1996
Days to decision	89 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Dayspring Medical, Inc.**
Location	Boulder, CO, US
Contact	NEIL RASMUSSEN
510(k) history	5 submissions · 5 cleared · 1996-1999