FDA 510(k) Premarket Notification

K863532 MODIFIED LABELING TO MODEL 2000 AMBULATORY INFUSER

CLEARED

Jan 14, 1987
127 days to decision

K863532 · Product code: **FRN** · General Hospital
Source: https://www.510kdatabase.net/k863532/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Sep 9, 1986
Decision date	Jan 14, 1987
Days to decision	127 days
Third-party review	No

APPLICANT

Company	**Parker Hannifin Corp.**
Location	Walker, MI, US
Contact	STEVE A WIRTZ
Website	http://www.parker.com/
510(k) history	8 submissions · 8 cleared · 1983-2017