

K863532 MODIFIED LABELING TO MODEL 2000 AMBULATORY INFUSERJan 14, 1987
127 days to decisionK863532 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k863532/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 1, 2026