FDA 510(k) Premarket Notification

K831761 SIMPLICITY PLUS

CLEARED

Aug 12, 1983
72 days to decision

K831761 · Product code: **FRN** · General Hospital
Source: https://www.510kdatabase.net/k831761/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Jun 1, 1983
Decision date	Aug 12, 1983
Days to decision	72 days
Third-party review	No

APPLICANT

Company	**Critikon Company, LLC**
Location	Mchenry, IL, US
510(k) history	51 submissions · 51 cleared · 1979-2000