FDA 510(k) Premarket Notification

K823035 COMPUTER INTERFACE MODULE CIM

CLEARED

Nov 24, 1982
41 days to decision

K823035 · Product code: **FRN** · General Hospital
Source: https://www.510kdatabase.net/k823035/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Oct 14, 1982
Decision date	Nov 24, 1982
Days to decision	41 days
Third-party review	No

APPLICANT

Company	**Y**
Location	Mchenry, IL, US
510(k) history	78 submissions · 78 cleared · 1978-1996

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k823035/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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