FDA 510(k) Premarket Notification

K834106 INTRAVENOUS INFUSION CONTROLLER

CLEARED

May 30, 1984
183 days to decision

K834106 · Product code: **LDR** · General Hospital
Source: https://www.510kdatabase.net/k834106/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Nov 29, 1983
Decision date	May 30, 1984
Days to decision	183 days
Third-party review	No

APPLICANT

Company	**Cutter Laboratories, Inc.**
Location	Mchenry, IL, US
Website	https://www.bayer.com
510(k) history	39 submissions · 39 cleared · 1976-1986