FDA 510(k) Premarket Notification

K802101 I.V. SENTRY ALARM DEVICE

CLEARED

Sep 16, 1980
13 days to decision

K802101 · Product code: **FLN** · General Hospital
Source: https://www.510kdatabase.net/k802101/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Electric For Gravity Flow Infusion Systems (FLN)
Date received	Sep 3, 1980
Decision date	Sep 16, 1980
Days to decision	13 days
Third-party review	No

APPLICANT

Company	**Pike Medical Co.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980