

K802101 I.V. SENTRY ALARM DEVICESep 16, 1980
13 days to decisionK802101 · Product code: **FLN** · General HospitalSource: <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

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Device record: <https://www.510kdatabase.net/k802101/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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