

K822891 FLO CONTROLNov 24, 1982
57 days to decisionK822891 · Product code: **LDR** · General Hospital
Source: <https://www.510kdatabase.net/k822891/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Sep 28, 1982
Decision date	Nov 24, 1982
Days to decision	57 days
Third-party review	No

APPLICANT

Company	Dna Medical, Inc.
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1982-1983

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Device record: <https://www.510kdatabase.net/k822891/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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