

K833659 IV EXTENSION SETNov 29, 1983
42 days to decisionK833659 · Product code: **FPK** · General HospitalSource: <https://www.510kdatabase.net/k833659/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubing, Fluid Delivery (FPK)
Date received	Oct 18, 1983
Decision date	Nov 29, 1983
Days to decision	42 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k833659/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 27, 2026