

K780673 CENTRAL CATHETER KITMay 16, 1978
22 days to decisionK780673 · Product code: **FPK** · General HospitalSource: <https://www.510kdatabase.net/k780673/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubing, Fluid Delivery (FPK)
Date received	Apr 24, 1978
Decision date	May 16, 1978
Days to decision	22 days
Third-party review	No

APPLICANT

Company	Argon Medical Corp.
Location	Mchenry, IL, US
510(k) history	27 submissions · 27 cleared · 1976-1991

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Device record: <https://www.510kdatabase.net/k780673/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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