

K802602 L-3 PURGE LINENov 12, 1980
22 days to decisionK802602 · Product code: **DTM** · CardiovascularSource: <https://www.510kdatabase.net/k802602/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Blood, Cardiopulmonary Bypass, Arterial Line (DTM)
Date received	Oct 21, 1980
Decision date	Nov 12, 1980
Days to decision	22 days
Third-party review	No

APPLICANT

Company	Delta Medical Industries
Location	Mchenry, IL, US
510(k) history	27 submissions · 27 cleared · 1976-1984

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Device record: <https://www.510kdatabase.net/k802602/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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