FDA 510(k) Premarket Notification

K802602 L-3 PURGE LINE

CLEARED

Nov 12, 1980
22 days to decision

K802602 · Product code: **DTM** · Cardiovascular
Source: 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