FDA 510(k) Premarket Notification

K831315 TRILEX

CLEARED

Aug 12, 1983
112 days to decision

K831315 · Product code: **FPB** · General Hospital
Source: https://www.510kdatabase.net/k831315/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Infusion Line (FPB)
Date received	Apr 22, 1983
Decision date	Aug 12, 1983
Days to decision	112 days
Third-party review	No

APPLICANT

Company	**Tri-Med, Inc.**
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1977-2004