FDA 510(k) Premarket Notification

K945160 PARACENTESIS NEEDLES/CANNULAS

CLEARED

Nov 14, 1994
24 days to decision

K945160 · Product code: **LRO** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k945160/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	General Surgery Tray (LRO)
Date received	Oct 21, 1994
Decision date	Nov 14, 1994
Days to decision	24 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Tri-Med Specialties, Inc.**
Location	Overland Park, KS, US
Contact	MIKE KNOTH
510(k) history	14 submissions · 12 cleared · 1985-1997