FDA 510(k) Premarket Notification

K981619 VACUETTE MULTIPLE SAMPLE LUER ADAPTER

CLEARED

Jul 15, 1998
70 days to decision

K981619 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k981619/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	May 6, 1998
Decision date	Jul 15, 1998
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Greiner Meditech, Inc.**
Location	Bel Air, MD, US
Contact	DOUGLAS L HARRIS
510(k) history	10 submissions · 10 cleared · 1997-1999