FDA 510(k) Premarket Notification

K943137 B-D/BFI MAIL DISPOSAL SERVICE

CLEARED

Nov 7, 1994
131 days to decision

K943137 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k943137/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jun 29, 1994
Decision date	Nov 7, 1994
Days to decision	131 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Med-Safe Systems, Inc.**
Location	Oceanside, CA, US
Contact	VIRGINIA SHILLINGTON
510(k) history	8 submissions · 8 cleared · 1994-1994