FDA 510(k) Premarket Notification

K980003 M500-100 SHARPS CONTAINER

CLEARED

Mar 13, 1998
70 days to decision

K980003 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k980003/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jan 2, 1998
Decision date	Mar 13, 1998
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Imagination Medical, Inc.**
Location	Jacksonville, FL, US
Contact	TIM WEIST
510(k) history	1 submissions · 1 cleared · 1998-1998