FDA 510(k) Premarket Notification

K992404 THE WAND

CLEARED

Jun 6, 2000
323 days to decision

K992404 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k992404/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jul 19, 1999
Decision date	Jun 6, 2000
Days to decision	323 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Spintech, Inc.**
Location	Minneapolis, MN, US
Contact	DANIEL J MANELLI
510(k) history	10 submissions · 10 cleared · 1992-2022