FDA 510(k) Premarket Notification

K001690 PROTECTIV SAFETY BLOOD COLLECTION NEEDLE

CLEARED

Jul 31, 2000
59 days to decision

K001690 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k001690/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jun 2, 2000
Decision date	Jul 31, 2000
Days to decision	59 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Johnson & Johnson Medical, Inc.**
Location	Arlington, TX, US
Contact	DORENE MARKWIESE
510(k) history	53 submissions · 50 cleared · 1990-2000