FDA 510(k) Premarket Notification

K041941 SAFE T RETRACTABLE BLOOD COLLECTION DEVICE

CLEARED

Sep 13, 2004
56 days to decision

K041941 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k041941/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jul 19, 2004
Decision date	Sep 13, 2004
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Safe T Medical Devices, Ltd.**
Location	Conway, NH, US
Contact	DEBBIE IAMPIETRO
510(k) history	1 submissions · 1 cleared · 2004-2004

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k041941/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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