FDA 510(k) Premarket Notification

K052445 DUOPROSS NEEDLE

CLEARED

Dec 5, 2005
89 days to decision

K052445 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k052445/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Sep 7, 2005
Decision date	Dec 5, 2005
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Duopross Meditech Corporation**
Location	Albuquerque, NM, US
Contact	FRANK FERGUSON
510(k) history	2 submissions · 2 cleared · 2005-2005