FDA 510(k) Premarket Notification

K861560 MEDIPOINT MULTI-SAMPLE BLOOD COLLECTING NEEDLE

CLEARED

May 12, 1986
17 days to decision

K861560 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k861560/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Apr 25, 1986
Decision date	May 12, 1986
Days to decision	17 days
Third-party review	No

APPLICANT

Company	**Medi Point, Inc.**
Location	Mineola, NY, US
Contact	PETER GOLLOBIN
510(k) history	1 submissions · 1 cleared · 1986-1986