

**K861560 MEDIPOINT MULTI-SAMPLE BLOOD COLLECTING NEEDLE**May 12, 1986  
17 days to decisionK861560 · 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	<b>Medi Point, Inc.</b>
Location	Mineola, NY, US
Contact	PETER GOLLOBIN
510(k) history	1 submissions · 1 cleared · 1986-1986

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k861560/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026