

**K890473 NEEDLE GUARD(TM)**Apr 3, 1989  
62 days to decisionK890473 · Product code: **FMI** · General Hospital  
Source: <https://www.510kdatabase.net/k890473/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jan 31, 1989
Decision date	Apr 3, 1989
Days to decision	62 days
Third-party review	No

**APPLICANT**

---

Company	<b>Martech Medical Products, Inc.</b>
Location	Riverside, CA, US
Contact	MICHAEL D MARTELL
510(k) history	7 submissions · 6 cleared · 1989-1996

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890473/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 26, 2026