

**K943485 MONOJECT HORIZONTAL ENTRY SHARPS
CONTAINER**Sep 9, 1994
52 days to decisionK943485 · Product code: **FMI** · General Hospital
Source: <https://www.510kdatabase.net/k943485/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jul 19, 1994
Decision date	Sep 9, 1994
Days to decision	52 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Sherwood Medical Co.
Location	Mchenry, IL, US
Contact	MICHAEL KROEHNKE
510(k) history	191 submissions · 177 cleared · 1976-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k943485/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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