

K905305 SAFE-SITE NEEDLE COVERFeb 12, 1991
76 days to decisionK905305 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k905305/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Nov 28, 1990
Decision date	Feb 12, 1991
Days to decision	76 days
Third-party review	No

APPLICANT

Company	North American Medical Products, Inc.
Location	Walker, MI, US
Contact	ARTHUR GIANAKOS
510(k) history	6 submissions · 6 cleared · 1984-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k905305/>; 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 May 27, 2026