

K960248 GENERAL PROCEDURE KITApr 12, 1996
87 days to decisionK960248 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k960248/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jan 16, 1996
Decision date	Apr 12, 1996
Days to decision	87 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Physician Industries, Inc.
Location	North Salt Lake, UT, US
Contact	BRIAN BAKER
510(k) history	4 submissions · 3 cleared · 1995-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k960248/> 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