

K882975 RE-VENT ARTERIAL BLOOD SAMPLER AND/OR BIOSAMPLERJan 25, 1989
194 days to decisionK882975 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k882975/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jul 15, 1988
Decision date	Jan 25, 1989
Days to decision	194 days
Third-party review	No

APPLICANT

Company	Richmann Laboratories
Location	Lake Park, FL, US
Contact	REED
510(k) history	1 submissions · 1 cleared · 1989-1989

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k882975/> 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