

K925210 90DX MULTITEST METER, MODIFICATIONJan 25, 1994
475 days to decisionK925210 · Product code: **FLB** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k925210/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Meter, Conductivity, Induction, Remote Type (FLB)
Date received	Oct 7, 1992
Decision date	Jan 25, 1994
Days to decision	475 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Mesa Medical, Inc.
Location	Mchenry, IL, US
Contact	JEFF ZEPP
510(k) history	7 submissions · 7 cleared · 1982-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k925210/> 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 July 2, 2026