

**K920410 EMILITH ELECTROMECHANICALIMPACTOR
LITHOTRIPSY PROB**Feb 14, 1994
745 days to decision

K920410 · Product code: FFK · Gastroenterology & Urology

Source: <https://www.510kdatabase.net/k920410/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lithotripter, Electro-hydraulic (FFK)
Date received	Jan 31, 1992
Decision date	Feb 14, 1994
Days to decision	745 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Psi Medical Products, Inc.
Location	Norwalk, CT, US
Contact	JOHN SOKOLOWSKI
510(k) history	7 submissions · 7 cleared · 1992-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k920410/>. 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