

K921612 INTERTRON 4500, INTERTRON 5000, MASTERSTIM I-2Aug 18, 1992
137 days to decisionK921612 · Product code: **LIH** · Neurology
Source: <https://www.510kdatabase.net/k921612/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Interferential Current Therapy (LIH)
Date received	Apr 3, 1992
Decision date	Aug 18, 1992
Days to decision	137 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Medstar, Inc.
Location	Glenview, IL, US
Contact	CRAIG SEYL
510(k) history	2 submissions · 2 cleared · 1988-1992

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k921612/> 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 1, 2026