

K935760 TEC 2100 (MODEL MT-LPLGD)Jan 25, 1994
60 days to decisionK935760 · Product code: **IWE** · Radiology
Source: <https://www.510kdatabase.net/k935760/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Patient Position, Light-beam (IWE)
Date received	Nov 26, 1993
Decision date	Jan 25, 1994
Days to decision	60 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Medtec, Inc.
Location	Orange City, IA, US
Contact	DONALD RIIBE
510(k) history	41 submissions · 41 cleared · 1993-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k935760/> 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 May 17, 2026