FDA 510(k) Premarket Notification

K070931 MODEL PMT850

CLEARED

May 24, 2007
51 days to decision

K070931 · Product code: **ILX** · Physical Medicine
Source: https://www.510kdatabase.net/k070931/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Diathermy, Shortwave, For Use Other Than Applying Therapeutic Deep Heat (ILX)
Date received	Apr 3, 2007
Decision date	May 24, 2007
Days to decision	51 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Promedtek, Inc.**
Location	Washington, DC, US
Contact	DIANE MANDELL
510(k) history	2 submissions · 2 cleared · 2007-2016

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k070931/ 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 2, 2026