

**K072256 HF54 COMBINATION ULTRASOUND INTERFERENTIAL
AND PREMODULATED STIMULATION SYSTEM WITH
OPTIONAL HANDS-FREE OPERATION, MODEL**Mar 12, 2008
212 days to decisionK072256 · Product code: **PFW** · Physical Medicine
Source: <https://www.510kdatabase.net/k072256/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stationary Ultrasonic Diathermy Device For Use In Applying Therapeutic Deep Heat (PFW)
Date received	Aug 13, 2007
Decision date	Mar 12, 2008
Days to decision	212 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Hill Laboratories Co.
Location	Malvern, PA, US
Contact	BRADY ALLER
510(k) history	17 submissions · 17 cleared · 1993-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k072256/> 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 21, 2026