FDA 510(k) Premarket Notification

K072551 LD-I 75 AND LD-I 200

CLEARED

Apr 2, 2008
205 days to decision

K072551 · Product code: **NHN** · Physical Medicine
Source: https://www.510kdatabase.net/k072551/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Light Based Laser Non-thermal Instrument With Non-heating Effect For Adjunctive Use In Pain Therapy (NHN)
Date received	Sep 10, 2007
Decision date	Apr 2, 2008
Days to decision	205 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Meditech International, Inc.**
Location	Stamford, CT, US
Contact	RICHARD KEEN
510(k) history	5 submissions · 5 cleared · 2003-2008