

**K072551 LD-I 75 AND LD-I 200**Apr 2, 2008  
205 days to decisionK072551 · Product code: **NHN** · Physical MedicineSource: <https://www.510kdatabase.net/k072551/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072551/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 24, 2026