

**K063162 GOLDEN LITEWAY**Nov 2, 2006  
15 days to decisionK063162 · Product code: **INI** · Physical Medicine  
Source: <https://www.510kdatabase.net/k063162/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Vehicle, Motorized 3-wheeled (INI)
Date received	Oct 18, 2006
Decision date	Nov 2, 2006
Days to decision	15 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Golden Technologies, Inc.</b>
Location	Wyoming, PA, US
Contact	GENE R KULON
510(k) history	8 submissions · 8 cleared · 1988-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k063162/> 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 29, 2026