FDA 510(k) Premarket Notification

K063162 GOLDEN LITEWAY

CLEARED

Nov 2, 2006
15 days to decision

K063162 · Product code: **INI** · Physical Medicine
Source: https://www.510kdatabase.net/k063162/

SUBMISSION DETAILS

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

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