FDA 510(k) Premarket Notification

K131909 DALTON LIFT CHAIR

CLEARED

Feb 24, 2014
243 days to decision

K131909 · Product code: **INO** · Physical Medicine
Source: https://www.510kdatabase.net/k131909/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chair, Positioning, Electric (INO)
Date received	Jun 26, 2013
Decision date	Feb 24, 2014
Days to decision	243 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Dalton Instrument Corp.**
Location	Richardson, TX, US
Contact	MEI LEIN
510(k) history	5 submissions · 5 cleared · 1985-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k131909/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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