FDA 510(k) Premarket Notification

K090632 COMPEX REHAB

CLEARED

Jun 1, 2009
84 days to decision

K090632 · Product code: **IPF** · Physical Medicine
Source: https://www.510kdatabase.net/k090632/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Mar 9, 2009
Decision date	Jun 1, 2009
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Chattanooga Group**
Location	Hixson, TN, US
Contact	NORA C.R. YORK
510(k) history	11 submissions · 11 cleared · 2002-2009