

**K090632 COMPEX REHAB**Jun 1, 2009  
84 days to decisionK090632 · Product code: **IPF** · Physical MedicineSource: <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	<b>Chattanooga Group</b>
Location	Hixson, TN, US
Contact	NORA C.R. YORK
510(k) history	11 submissions · 11 cleared · 2002-2009

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090632/> 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 23, 2026