

**K090532 T.E.A.R. TECH3**Nov 13, 2009  
259 days to decisionK090532 · Product code: **IPF** · Physical Medicine  
Source: <https://www.510kdatabase.net/k090532/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Feb 27, 2009
Decision date	Nov 13, 2009
Days to decision	259 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vision Quest Industries, Inc.</b>
Location	Irvine, CA, US
Contact	WALLACE FISCHER
510(k) history	6 submissions · 6 cleared · 1998-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090532/> 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 5, 2026