FDA 510(k) Premarket Notification

K090532 T.E.A.R. TECH3

CLEARED

Nov 13, 2009
259 days to decision

K090532 · Product code: **IPF** · Physical Medicine
Source: https://www.510kdatabase.net/k090532/

SUBMISSION DETAILS

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

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