FDA 510(k) Premarket Notification

K123354 REVITIVE IX

CLEARED

Jun 28, 2013
240 days to decision

K123354 · Product code: **IPF** · Physical Medicine
Source: https://www.510kdatabase.net/k123354/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Oct 31, 2012
Decision date	Jun 28, 2013
Days to decision	240 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Actegy , Ltd.**
Location	Washington, DC, US
Contact	JOHN J SMITH
510(k) history	7 submissions · 7 cleared · 2013-2021