FDA 510(k) Premarket Notification

K123746 AMES THERAPY DEVICE

CLEARED

May 24, 2013
169 days to decision

K123746 · Product code: **IKK** · Physical Medicine
Source: https://www.510kdatabase.net/k123746/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Isokinetic Testing And Evaluation (IKK)
Date received	Dec 6, 2012
Decision date	May 24, 2013
Days to decision	169 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Ames Technology, Inc.**
Location	Eugene, OR, US
Contact	SHEILA RAMERMAN, RAC
510(k) history	1 submissions · 1 cleared · 2013-2013