FDA 510(k) Premarket Notification

K093553 POSTURETEK

CLEARED

Aug 13, 2010
269 days to decision

K093553 · Product code: **LZW** · Physical Medicine
Source: https://www.510kdatabase.net/k093553/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Spine Curvature (LZW)
Date received	Nov 17, 2009
Decision date	Aug 13, 2010
Days to decision	269 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Perseus Althetics,Llc**
Location	Washington, DC, US
Contact	KEITH BARRITT
510(k) history	1 submissions · 1 cleared · 2010-2010