

**K093553 POSTURETEK**Aug 13, 2010  
269 days to decisionK093553 · Product code: **LZW** · Physical MedicineSource: <https://www.510kdatabase.net/k093553/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093553/> 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 20, 2026