

**K131253 PERIFLUX 6000**Oct 22, 2013  
173 days to decisionK131253 · Product code: **LKD** · Anesthesiology  
Source: <https://www.510kdatabase.net/k131253/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Monitor, Carbon-dioxide, Cutaneous (LKD)
Date received	May 2, 2013
Decision date	Oct 22, 2013
Days to decision	173 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Perimed AB</b>
Location	Jarfalla, SE
Contact	MARIA LILJEVRET
510(k) history	4 submissions · 4 cleared · 2001-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k131253/> 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 May 24, 2026