FDA 510(k) Premarket Notification

K131253 PERIFLUX 6000

CLEARED

Oct 22, 2013
173 days to decision

K131253 · Product code: **LKD** · Anesthesiology
Source: https://www.510kdatabase.net/k131253/

SUBMISSION DETAILS

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

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