

**K990960 PERIMED TRANSCUTANEOUS PO2 AND PCO2
MONITOR (PF5040)**Sep 23, 1999
185 days to decisionK990960 · Product code: **LKD** · Anesthesiology
Source: <https://www.510kdatabase.net/k990960/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Carbon-dioxide, Cutaneous (LKD)
Date received	Mar 22, 1999
Decision date	Sep 23, 1999
Days to decision	185 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Perimed, Inc.
Location	Piscataway, NJ, US
Contact	JAMES HOWARD
510(k) history	6 submissions · 6 cleared · 1989-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k990960/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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