

**K900333 E5280 COMBINED TRANSCUTANEOUS
TCP02/TCPCO2 ELECT.**Apr 16, 1990
83 days to decisionK900333 · Product code: **LKD** · Anesthesiology
Source: <https://www.510kdatabase.net/k900333/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Carbon-dioxide, Cutaneous (LKD)
Date received	Jan 23, 1990
Decision date	Apr 16, 1990
Days to decision	83 days
Third-party review	No

APPLICANT

Company	Radiometer America, Inc.
Location	Mchenry, IL, US
Contact	DONALD BAKER
510(k) history	50 submissions · 50 cleared · 1983-2016

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k900333/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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