

K860124 HORIZON 2000 TRANSCUTANEOUS GAS MONITORINGMay 9, 1986
115 days to decisionK860124 · Product code: **KLK** · Anesthesiology
Source: <https://www.510kdatabase.net/k860124/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Oxygen, Cutaneous, For Infant Not Under Gas Anesthesia (KLK)
Date received	Jan 14, 1986
Decision date	May 9, 1986
Days to decision	115 days
Third-party review	No

APPLICANT

Company	Mennen Medical, Inc.
Location	Clarence, NY, US
Contact	RONALD WIDMAN
510(k) history	34 submissions · 34 cleared · 1985-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k860124/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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