

K802846 TCM202 TCPO2 & TCM204 TCPO2/PO2 REC. SY.Dec 17, 1980
35 days to decisionK802846 · Product code: **KLK** · Anesthesiology
Source: <https://www.510kdatabase.net/k802846/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Oxygen, Cutaneous, For Infant Not Under Gas Anesthesia (KLK)
Date received	Nov 12, 1980
Decision date	Dec 17, 1980
Days to decision	35 days
Third-party review	No

APPLICANT

Company	The London Co.
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1976-1980

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k802846/> Data sourced from FDA 510(k) public records (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. - Generated May 28, 2026