FDA 510(k) Premarket Notification

CLEARED

K802846 TCM202 TCPO2 & TCM204 TCPO2/PO2 REC. SY.

Dec 17, 1980
35 days to decision

K802846 · 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