

**K860951 KONTRON CUTANEOUS PO2 MONITOR 820**Apr 9, 1986  
28 days to decisionK860951 · Product code: **KLK** · AnesthesiologySource: <https://www.510kdatabase.net/k860951/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Oxygen, Cutaneous, For Infant Not Under Gas Anesthesia (KLK)
Date received	Mar 12, 1986
Decision date	Apr 9, 1986
Days to decision	28 days
Third-party review	No

**APPLICANT**

---

Company	<b>Kontron Instruments, Inc.</b>
Location	Mchenry, IL, US
Contact	GEORGE CHO
510(k) history	57 submissions · 57 cleared · 1981-1993

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860951/>, Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 21, 2026