

**K844731 KONTRON SUPERMON 7210 PATIENT MONITOR**May 16, 1985  
163 days to decisionK844731 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k844731/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiometer & Rate Alarm) (DRT)
Date received	Dec 4, 1984
Decision date	May 16, 1985
Days to decision	163 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/k844731/>; 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