

**K895425 MINIMON 7137B PATIENT MONITOR**Sep 26, 1989  
19 days to decisionK895425 · Product code: **DQA** · AnesthesiologySource: <https://www.510kdatabase.net/k895425/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Sep 7, 1989
Decision date	Sep 26, 1989
Days to decision	19 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895425/>; 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