

**K812144 CENTRAL MONITOR 360**Sep 25, 1981  
59 days to decisionK812144 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k812144/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jul 28, 1981
Decision date	Sep 25, 1981
Days to decision	59 days
Third-party review	No

**APPLICANT**

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Company	<b>Life Science Instrumentation, Inc.</b>
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1981-1985

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k812144/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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