

**K792081 SPECTRA CENTRAL PT MONITORING SYS**Nov 13, 1979  
28 days to decisionK792081 · Product code: **DSI** · CardiovascularSource: <https://www.510kdatabase.net/k792081/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Oct 16, 1979
Decision date	Nov 13, 1979
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>American Optical Corp.</b>
Location	Mchenry, IL, US
510(k) history	35 submissions · 35 cleared · 1976-1995

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Device record: <https://www.510kdatabase.net/k792081/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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