

K781845 CAMSCAN SYSTEMNov 20, 1978
19 days to decisionK781845 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k781845/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Nov 1, 1978
Decision date	Nov 20, 1978
Days to decision	19 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k781845/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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