

K780043 ESIS MODULEFeb 13, 1978
35 days to decisionK780043 · Product code: **DRR** · CardiovascularSource: <https://www.510kdatabase.net/k780043/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Amplifier And Signal Conditioner, Biopotential (DRR)
Date received	Jan 9, 1978
Decision date	Feb 13, 1978
Days to decision	35 days
Third-party review	No

APPLICANT

Company	Datascope Corp.
Location	Mchenry, IL, US
510(k) history	136 submissions · 135 cleared · 1976-2019

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Device record: <https://www.510kdatabase.net/k780043/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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