

K802179 CARDIOSCOPE J200Nov 12, 1980
64 days to decisionK802179 · Product code: **DRS** · CardiovascularSource: <https://www.510kdatabase.net/k802179/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Sep 9, 1980
Decision date	Nov 12, 1980
Days to decision	64 days
Third-party review	No

APPLICANT

Company	Telectronics, Inc.
Location	Mchenry, IL, US
510(k) history	107 submissions · 107 cleared · 1977-1990

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Device record: <https://www.510kdatabase.net/k802179/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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