

K791710 CARDIOVASCULAR INSTRUMENTS-VARIOUSSep 12, 1979
16 days to decisionK791710 · Product code: **DWS** · Cardiovascular
Source: <https://www.510kdatabase.net/k791710/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instruments, Surgical, Cardiovascular (DWS)
Date received	Aug 27, 1979
Decision date	Sep 12, 1979
Days to decision	16 days
Third-party review	No

APPLICANT

Company	Edward Weck, Inc.
Location	Mchenry, IL, US
510(k) history	140 submissions · 140 cleared · 1976-1992

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Device record: <https://www.510kdatabase.net/k791710/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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