

K792088 AORTIC OCCLUDER #000390Oct 26, 1979
9 days to decisionK792088 · Product code: **DWS** · CardiovascularSource: <https://www.510kdatabase.net/k792088/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instruments, Surgical, Cardiovascular (DWS)
Date received	Oct 17, 1979
Decision date	Oct 26, 1979
Days to decision	9 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/k792088/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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