FDA 510(k) Premarket Notification

# K792088 AORTIC OCCLUDER #000390

**CLEARED**

Oct 26, 1979
9 days to decision

K792088 · Product code: **DWS** · Cardiovascular
Source: 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

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