FDA 510(k) Premarket Notification

K791710 CARDIOVASCULAR INSTRUMENTS-VARIOUS

CLEARED

Sep 12, 1979
16 days to decision

K791710 · 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