FDA 510(k) Premarket Notification

K791703 TEW BYPASS INSTRUMENTS & DISSECTING KIT

CLEARED

Oct 4, 1979
37 days to decision

K791703 · Product code: **GDI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k791703/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dissector, Surgical, General & Plastic Surgery (GDI)
Date received	Aug 28, 1979
Decision date	Oct 4, 1979
Days to decision	37 days
Third-party review	No

APPLICANT

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