FDA 510(k) Premarket Notification

K810324 AUTOMATIC HEMOCLIP APPLIER, 527700

CLEARED

Mar 20, 1981
42 days to decision

K810324 · Product code: **GDW** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k810324/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Implantable (GDW)
Date received	Feb 6, 1981
Decision date	Mar 20, 1981
Days to decision	42 days
Third-party review	No

APPLICANT

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