

K810324 AUTOMATIC HEMOCLIP APPLIER, 527700Mar 20, 1981
42 days to decisionK810324 · Product code: **GDW** · General & Plastic SurgerySource: <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

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Device record: <https://www.510kdatabase.net/k810324/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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