

K791741 FLOOR LINE INSTRUMENTSOct 4, 1979
24 days to decisionK791741 · Product code: **FZT** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k791741/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cutter, Surgical (FZT)
Date received	Sep 10, 1979
Decision date	Oct 4, 1979
Days to decision	24 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/k791741/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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