

**K791703 TEW BYPASS INSTRUMENTS & DISSECTING KIT**Oct 4, 1979  
37 days to decisionK791703 · Product code: **GDI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k791703/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k791703/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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