

**K771332 PATIENT GROUNDING CABLES**Sep 6, 1977  
50 days to decisionK771332 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k771332/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 18, 1977
Decision date	Sep 6, 1977
Days to decision	50 days
Third-party review	No

**APPLICANT**

---

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

---

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

Device record: <https://www.510kdatabase.net/k771332/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 23, 2026