

K771330 DISP. GROUND PLATESep 6, 1977
50 days to decisionK771330 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k771330/>**SUBMISSION DETAILS**

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k771330/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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