

**K823923 MF 360B ELECTROSURGICAL UNIT**Feb 15, 1983  
49 days to decisionK823923 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k823923/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 28, 1982
Decision date	Feb 15, 1983
Days to decision	49 days
Third-party review	No

**APPLICANT**

---

Company	<b>Aspen Laboratories, Inc.</b>
Location	Mchenry, IL, US
510(k) history	55 submissions · 55 cleared · 1976-1998

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k823923/> 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 22, 2026