

K852355 SINGLE USE ARTHROSCOPY ELECTRODESep 3, 1985
92 days to decisionK852355 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k852355/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 3, 1985
Decision date	Sep 3, 1985
Days to decision	92 days
Third-party review	No

APPLICANT

Company	Aspen Laboratories, Inc.
Location	Mchenry, IL, US
Contact	JOHNSON
510(k) history	55 submissions · 55 cleared · 1976-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k852355/>; Data sourced from FDA 510(k) public records (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. - Generated May 22, 2026