

K822092 CAMERON-MILLER #40-1516Sep 21, 1982
67 days to decisionK822092 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k822092/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 16, 1982
Decision date	Sep 21, 1982
Days to decision	67 days
Third-party review	No

APPLICANT

Company	Cameron-Miller, Inc.
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1977-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k822092/> 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 June 30, 2026