

K820494 SYBRON BOVIE CSV HAND CONTR. ELECTROSURGApr 9, 1982
45 days to decisionK820494 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k820494/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 23, 1982
Decision date	Apr 9, 1982
Days to decision	45 days
Third-party review	No

APPLICANT

Company	Sybron Corp.
Location	Mchenry, IL, US
510(k) history	37 submissions · 37 cleared · 1977-1986

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k820494/> 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 21, 2026