

**K914883 CIRCON ACMI GEN SURG INSTRU FOR MINI
INVASIVE SURG**Jan 15, 1993
443 days to decisionK914883 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k914883/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 30, 1991
Decision date	Jan 15, 1993
Days to decision	443 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Circon Video
Location	Stanford, CT, US
Contact	ERVIN F TAYLOR
510(k) history	14 submissions · 14 cleared · 1990-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k914883/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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