

**K960695 MODEL 200C ELECTROSURGICAL PROBE**Apr 30, 1996  
70 days to decisionK960695 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k960695/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 20, 1996
Decision date	Apr 30, 1996
Days to decision	70 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Zomed Intl.</b>
Location	Mountain View, CA, US
Contact	THOMAS C WEHMAN, PH.D.
510(k) history	7 submissions · 7 cleared · 1995-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960695/> 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 July 5, 2026