

**K021299 CONMED HEATWAVE HAND-CONTROLLED  
ELETRODE**Jun 26, 2002  
63 days to decisionK021299 · Product code: **GEI** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k021299/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Apr 24, 2002
Decision date	Jun 26, 2002
Days to decision	63 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Conmedcorp</b>
Location	Dayton, OH, US
Contact	IRA D DUESLER
510(k) history	92 submissions · 92 cleared · 1981-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k021299/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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