

**K982130 RETURN ELECTRODE CLAMP SHELL CORD MODEL
RC 201**Aug 7, 1998
51 days to decisionK982130 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k982130/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 17, 1998
Decision date	Aug 7, 1998
Days to decision	51 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	New Deantronics, Ltd.
Location	Boulder, CO, US
Contact	LEWIS WARD
510(k) history	2 submissions · 2 cleared · 1998-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k982130/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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