FDA 510(k) Premarket Notification

CLEARED

K982130 RETURN ELECTRODE CLAMP SHELL CORD MODEL RC 201

Aug 7, 1998
51 days to decision

K982130 · Product code: **GEI** · General & Plastic Surgery
Source: 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