FDA 510(k) Premarket Notification

K033880 ELECTRO-LUBE

CLEARED

Mar 10, 2004
86 days to decision

K033880 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k033880/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 15, 2003
Decision date	Mar 10, 2004
Days to decision	86 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Mectra Labs, Inc.**
Location	Bloomfield, IN, US
Contact	CHARLES E ALLGOOD JR.
510(k) history	9 submissions · 9 cleared · 1992-2012