FDA 510(k) Premarket Notification

K081431 MEG LAPAROSCOPIC ELECTRODES

CLEARED

Jul 24, 2008
64 days to decision

K081431 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k081431/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 21, 2008
Decision date	Jul 24, 2008
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Bovie Medical**
Location	St. Petersburg, FL, US
Contact	RICHARD KOZLOFF
510(k) history	8 submissions · 8 cleared · 2006-2010