FDA 510(k) Premarket Notification

K080741 MEGA SOFT REUSABLE PATIENT RETURN ELECTRODE

CLEARED

Dec 16, 2008
274 days to decision

K080741 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k080741/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 17, 2008
Decision date	Dec 16, 2008
Days to decision	274 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Megadyne Medical Products, Inc.**
Location	Murray, UT, US
Contact	RONDA K MAGNESON
510(k) history	50 submissions · 48 cleared · 1990-2024