FDA 510(k) Premarket Notification

K020711 WALLACH LOOP ELECTRODE

CLEARED

Jun 3, 2002
90 days to decision

K020711 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k020711/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 5, 2002
Decision date	Jun 3, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Wallach Surgical Devices, Inc.**
Location	Walker, MI, US
Contact	MICHAEL MALIS
510(k) history	27 submissions · 27 cleared · 1981-2002