FDA 510(k) Premarket Notification

K971711 SOMNUS MODEL 215 ELECTROSURGICAL GENERATOR

CLEARED

Jun 24, 1997
47 days to decision

K971711 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k971711/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 8, 1997
Decision date	Jun 24, 1997
Days to decision	47 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Somnus Medical Technologies, Inc.**
Location	Sunnyvale, CA, US
Contact	EVE CONNER
510(k) history	16 submissions · 16 cleared · 1996-2000