FDA 510(k) Premarket Notification

K970837 SOMNUS MODEL 6000 DISPOSABLE TISSUE COAGULATING ELECTRODE

CLEARED

Jan 9, 1998
308 days to decision

K970837 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k970837/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 7, 1997
Decision date	Jan 9, 1998
Days to decision	308 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