FDA 510(k) Premarket Notification

CLEARED

K000936 VAPR 2.3MM WEDGE ELECTRODE FOR USE WITH VAPR SYSTEM

Apr 19, 2000
27 days to decision

K000936 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k000936/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 23, 2000
Decision date	Apr 19, 2000
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Mitek Products**
Location	Westwood, MA, US
Contact	MARY P LEGRAW
510(k) history	30 submissions · 30 cleared · 1996-2002