FDA 510(k) Premarket Notification

K023924 AUTOCON II 200 ELECTROSURGICAL GENERATOR

CLEARED

Feb 21, 2003
88 days to decision

K023924 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k023924/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 25, 2002
Decision date	Feb 21, 2003
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Karl Storz Endoscopy**
Location	Culver City, CA, US
Contact	JAMES A LEE
510(k) history	35 submissions · 35 cleared · 1995-2005