FDA 510(k) Premarket Notification

K001382 AARON 2100 HIGH FREQUENCY ELECTROSURGICAL GENERATOR

CLEARED

Jun 27, 2000
56 days to decision

K001382 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k001382/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 2, 2000
Decision date	Jun 27, 2000
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Aaron Medical Industries**
Location	St. Petersburg, FL, US
Contact	RICK KOZLOFF
510(k) history	46 submissions · 46 cleared · 1990-2006