FDA 510(k) Premarket Notification

K151696 Caiman Seal and Cut Technology

CLEARED

Jul 20, 2015
27 days to decision

K151696 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k151696/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 23, 2015
Decision date	Jul 20, 2015
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Aesculap, Inc.**
Location	Burlingame, CA, US
Contact	Denise Adams
510(k) history	207 submissions · 201 cleared · 1991-2025