FDA 510(k) Premarket Notification

K070004 ACCENT

CLEARED

Apr 23, 2007
110 days to decision

K070004 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k070004/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 3, 2007
Decision date	Apr 23, 2007
Days to decision	110 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Alma Lasers , Ltd.**
Location	Pleasanton, CA, US
Contact	ANNE WORDEN
510(k) history	14 submissions · 14 cleared · 2007-2022

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k070004/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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