FDA 510(k) Premarket Notification

K171986 Sterling Supreme Diode Laser

CLEARED

Sep 22, 2017
81 days to decision

K171986 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k171986/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jul 3, 2017
Decision date	Sep 22, 2017
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**CAO Group, Inc.**
Location	West Jordan, UT, US
Contact	Robert K. Larsen
Website	https://www.caogroup.com
510(k) history	31 submissions · 31 cleared · 2001-2026

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k171986/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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