FDA 510(k) Premarket Notification

K123443 PRECISE SHP DIODE LASER

CLEARED

Aug 13, 2013
278 days to decision

K123443 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k123443/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Nov 8, 2012
Decision date	Aug 13, 2013
Days to decision	278 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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