FDA 510(k) Premarket Notification

K070466 REVOLIX DUO LASER SYSTEM

CLEARED

Apr 17, 2007
60 days to decision

K070466 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k070466/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Feb 16, 2007
Decision date	Apr 17, 2007
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Allmed Systems, Inc.**
Location	Pleasanton, CA, US
Contact	PETER ALLEN
510(k) history	8 submissions · 8 cleared · 2004-2014