FDA 510(k) Premarket Notification

K061720 COSMELIGHT

CLEARED

Jan 31, 2007
226 days to decision

K061720 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k061720/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jun 19, 2006
Decision date	Jan 31, 2007
Days to decision	226 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Penntack Enterprises, Inc.**
Location	Miami, FL, US
Contact	PABLO PENA
510(k) history	1 submissions · 1 cleared · 2007-2007