FDA 510(k) Premarket Notification

K951117 LIHTAN 532 LASER

CLEARED

Oct 18, 1995
222 days to decision

K951117 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k951117/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Mar 10, 1995
Decision date	Oct 18, 1995
Days to decision	222 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Buckman Co., Inc.**
Location	Mchenry, IL, US
Contact	DAVID W SCHLERF
510(k) history	111 submissions · 104 cleared · 1983-1998