FDA 510(k) Premarket Notification

K920004 PBI MULTILASE D COPPER VAPOR LASER

CLEARED

May 14, 1992
133 days to decision

K920004 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k920004/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jan 2, 1992
Decision date	May 14, 1992
Days to decision	133 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Wm.I.Kern & Co., Inc.**
Location	Smithtown, NY, US
Contact	WILLIAM I.KERN
510(k) history	1 submissions · 1 cleared · 1992-1992