FDA 510(k) Premarket Notification

K782030 LASER SYSTEM, SHARPLAN 791/HE, CO2

CLEARED

Jan 22, 1979
48 days to decision

K782030 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k782030/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Dec 5, 1978
Decision date	Jan 22, 1979
Days to decision	48 days
Third-party review	No

APPLICANT

Company	**Laser Industries , Ltd.**
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1978-1994

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k782030/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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