FDA 510(k) Premarket Notification

K853900 ENDO-LASE CD40 FOR NEUROSURGERY

CLEARED

Jun 18, 1986
271 days to decision

K853900 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k853900/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Sep 20, 1985
Decision date	Jun 18, 1986
Days to decision	271 days
Third-party review	No

APPLICANT

Company	**Endo Lase, Inc.**
Location	Walker, MI, US
Contact	& HARTSON
510(k) history	24 submissions · 24 cleared · 1984-1986