FDA 510(k) Premarket Notification

K913109 LATERALASE(TM) CATHETER WITH THERMAL FEEDBACK

CLEARED

Oct 8, 1991
88 days to decision

K913109 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k913109/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jul 12, 1991
Decision date	Oct 8, 1991
Days to decision	88 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Trimedyne, Inc.**
Location	Walker, MI, US
Contact	MERRITT M GIRGIS
Website	http://www.trimedyne.com/
510(k) history	58 submissions · 58 cleared · 1981-2005

Trimedyne, Inc. is a manufacturer of Holmium:YAG lasers and surgical peripherals. The company specializes in laser-based surgical solutions for minimally invasive procedures across multiple specialties including urology, orthopedics, spine surgery, and general surgery. Trimedyne has received FDA 510(k) clearances from total submissions since its first clearance in 1981. The company's regulatory focus centers on General & Plastic Surgery devices, which represent 83% of its submission history. The latest clearance on record dates to 2005, reflecting the company's historical...