

**K914952 MODEL 1210 OMNIPULSE HOLMIUM LASER SYSTEM**Dec 20, 1991  
45 days to decisionK914952 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k914952/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Nov 5, 1991
Decision date	Dec 20, 1991
Days to decision	45 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Trimedynne, Inc.</b>
Location	Walker, MI, US
Contact	PAUL S.KRAMSKY
Website	<a href="http://www.trimedynne.com/">http://www.trimedynne.com/</a>
510(k) history	58 submissions · 58 cleared · 1981-2005

Trimedynne, 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. Trimedynne 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...

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