FDA 510(k) Premarket Notification

K830084 ANGIOSCOPE

CLEARED

Jun 2, 1983
142 days to decision

K830084 · Product code: **LYK** · Cardiovascular
Source: https://www.510kdatabase.net/k830084/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Angioscope (LYK)
Date received	Jan 11, 1983
Decision date	Jun 2, 1983
Days to decision	142 days
Third-party review	No

APPLICANT

Company	**Trimedyne, Inc.**
Location	Walker, MI, US
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...