FDA 510(k) Premarket Notification

K142310 Stryker Infrared Fluorescence (IRF) Imaging System

CLEARED

Dec 2, 2014
105 days to decision

K142310 · Product code: **OWN** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k142310/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Confocal Optical Imaging (OWN)
Date received	Aug 19, 2014
Decision date	Dec 2, 2014
Days to decision	105 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stryker Endoscopy**
Location	San Jose, CA, US
Contact	Golnaz Moeini
Website	https://www.stryker.com
510(k) history	100 submissions · 100 cleared · 1993-2026

Stryker Endoscopy is a medical device manufacturer based in San Jose, US. The company specializes in endoscopic and surgical imaging systems for minimally invasive procedures. Stryker Endoscopy has received FDA 510(k) clearances from total submissions since its first clearance in 1993. The company maintains active regulatory status, with its latest clearance in 2026. Its portfolio spans General & Plastic Surgery devices, orthopedic surgical tools, and obstetric and gynecologic instruments. Recent cleared devices include advanced imaging systems such as 4K camera platforms...