

**K022248 STRYKER L3 HYDROLERT**Aug 6, 2002  
25 days to decisionK022248 · Product code: **FLN** · General Hospital  
Source: <https://www.510kdatabase.net/k022248/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Electric For Gravity Flow Infusion Systems (FLN)
Date received	Jul 12, 2002
Decision date	Aug 6, 2002
Days to decision	25 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Endoscopy</b>
Location	San Jose, CA, US
Contact	MICHAEL HILLDOERFER
Website	<a href="https://www.stryker.com">https://www.stryker.com</a>
510(k) history	101 submissions · 101 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...

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