

# K222079 Connected OR Hub with Device and Voice Control, SDC4K Information Management System with Device and Voice Control

Oct 11, 2022  
88 days to decisionK222079 · Product code: **GCJ** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k222079/>

## SUBMISSION DETAILS

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, General & Plastic Surgery (GCJ)
Date received	Jul 15, 2022
Decision date	Oct 11, 2022
Days to decision	88 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

## APPLICANT

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Company	<b>Stryker Endoscopy</b>
Location	San Jose, CA, US
Contact	Divya Sekar
Website	<a href="https://www.stryker.com">https://www.stryker.com</a>
510(k) history	99 submissions · 99 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...