

**K952681 STRYKER HUMMER II MICRODEBRIDER SYSTEM**Jul 20, 1995  
38 days to decisionK952681 · Product code: **ERL** · Ear, Nose, Throat  
Source: <https://www.510kdatabase.net/k952681/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drill, Surgical, Ent (electric Or Pneumatic) Including Handpiece (ERL)
Date received	Jun 12, 1995
Decision date	Jul 20, 1995
Days to decision	38 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Endoscopy</b>
Location	San Jose, CA, US
Contact	JEFF KABLIK
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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