

**K874690 GASTROPLASTY CALIBRATING TUBE & GASTROSTENOMETER**Feb 8, 1988  
84 days to decisionK874690 · Product code: FFX · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k874690/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Gastrointestinal Motility (electrical) (FFX)
Date received	Nov 16, 1987
Decision date	Feb 8, 1988
Days to decision	84 days
Third-party review	No

**APPLICANT**

---

Company	<b>Mentor Corp.</b>
Location	Mchenry, IL, US
Contact	LYNN R BRECKENRIDGE
510(k) history	61 submissions · 61 cleared · 1977-2013

Mentor Corp. is a surgical aesthetics and medical device company based in McHenry, US. Now part of Johnson & Johnson MedTech, the brand supplies products to plastic surgeons and specialists worldwide. Mentor has received FDA 510(k) clearances from total submissions since its first clearance in 1977. The company's regulatory record spans General & Plastic Surgery, Gastroenterology & Urology, Obstetrics & Gynecology, and Radiology device categories. The latest clearance was recorded in 2013, reflecting the company's historical significance in surgical device innovation. Men...

---