

**K080253 KIMBERLY-CLARK INTRODUCER KITS**Feb 26, 2008  
26 days to decisionK080253 · Product code: **KGC** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k080253/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent - K
Submission type	Special
Device classification	Tube, Gastro-enterostomy (KGC)
Date received	Jan 31, 2008
Decision date	Feb 26, 2008
Days to decision	26 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Kimberly-Clark Corp.</b>
Location	Mchenry, IL, US
Contact	SHERRY SAURINI
Website	<a href="http://www.Kimberly-Clark.com">http://www.Kimberly-Clark.com</a>
510(k) history	104 submissions · 103 cleared · 1978-2014

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k080253/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 24, 2026