

**K090183 HERCULES 3 STAGE WIRE GUIDED DILATION BALLOON**

Sep 25, 2009  
242 days to decision

K090183 · Product code: **KNQ** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k090183/>

**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Dilator, Esophageal (KNQ)
Date received	Jan 26, 2009
Decision date	Sep 25, 2009
Days to decision	242 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Cook Endoscopy</b>
Location	Winstrom-Salem, NC, US
Contact	TIFFANNY THOMAS
510(k) history	4 submissions · 4 cleared · 2006-2020

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k090183/> Data sourced from FDA 510(k) public records (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. - Generated June 22, 2026