

**K040388 NEXT GENERATION SALEM SUMP**May 17, 2004  
90 days to decisionK040388 · Product code: **FEG** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k040388/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Double Lumen For Intestinal Decompression And/or Intubation (FEG)
Date received	Feb 17, 2004
Decision date	May 17, 2004
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Kendall</b>
Location	Mansfield, MA, US
Contact	JIM WELSH
510(k) history	12 submissions · 10 cleared · 2004-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k040388/> 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 June 22, 2026