

**K032138 MEDTRONIC SINGLE USE ESOPHAGEAL  
MANOMETRIC CATHETER, MEDTRONIC SINGLE USE  
ANORECTAL MANOMETRIC CATHETER**Oct 9, 2003  
90 days to decisionK032138 · Product code: **KLA** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k032138/>**SUBMISSION DETAILS**

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|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)                               |
| Submission type       | Traditional  |
| Device classification | Monitor, Esophageal Motility, Anorectal Motility, And Tube (KLA) |
| Date received         | Jul 11, 2003   |
| Decision date         | Oct 9, 2003  |
| Days to decision      | 90 days  |
| Third-party review    | No   |
| Summary / Statement   | Summary  |

**APPLICANT**

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|----------------|---|
| Company        | <b>Medtronic Vascular</b>                 |
| Location       | Walker, MI, US                            |
| Contact        | DEBORAH KIDDER                            |
| 510(k) history | 475 submissions · 453 cleared · 1977-2023 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k032138/> 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 18, 2026