

**K012147 DEVICE MODIFICATION OF ENTERAL FEEDING SETS  
FOR GRAVITY AND PUMP USE**Nov 2, 2001  
115 days to decisionK012147 · Product code: **KNT** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k012147/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Jul 10, 2001
Decision date	Nov 2, 2001
Days to decision	115 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Zevex, Inc.</b>
Location	Salt Lake City, UT, US
Contact	SUSAN P SCHMIDT
510(k) history	5 submissions · 5 cleared · 2001-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k012147/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 26, 2026