

**K943250 LIBERTY**Feb 9, 1995  
218 days to decisionK943250 · Product code: **KPI** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k943250/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Non-implantable, For Incontinence (KPI)
Date received	Jul 6, 1994
Decision date	Feb 9, 1995
Days to decision	218 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Utah Medical Products, Inc.</b>
Location	Mchenry, IL, US
Contact	KEVIN L CORNWELL
510(k) history	38 submissions · 38 cleared · 1979-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k943250/> 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 23, 2026