

**K141643 YARLAP**Apr 24, 2015  
309 days to decisionK141643 · Product code: **KPI** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k141643/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Non-implantable, For Incontinence (KPI)
Date received	Jun 19, 2014
Decision date	Apr 24, 2015
Days to decision	309 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>International Trade Group, Inc.</b>
Location	Oxford, OH, US
Contact	BRENT C REIDER
510(k) history	4 submissions · 4 cleared · 2010-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k141643/> 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 1, 2026