

**K895346 DR. DRY**Oct 24, 1989  
53 days to decisionK895346 · Product code: **KPN** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k895346/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alarm, Conditioned Response Enuresis (KPN)
Date received	Sep 1, 1989
Decision date	Oct 24, 1989
Days to decision	53 days
Third-party review	No

**APPLICANT**

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Company	<b>Intellitech</b>
Location	La Jolla, CA, US
Contact	MARK WIESNER
510(k) history	1 submissions · 1 cleared · 1989-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895346/>; 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 July 2, 2026