

**K782093 DETECTOR PAD**Feb 21, 1979  
69 days to decisionK782093 · Product code: **KPN** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k782093/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alarm, Conditioned Response Enuresis (KPN)
Date received	Dec 14, 1978
Decision date	Feb 21, 1979
Days to decision	69 days
Third-party review	No

**APPLICANT**

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Company	<b>Halperin Institute, Ltd. The</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1979-1979

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k782093/> 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 4, 2026