

**K901907 PROFESSIONAL MONITOR**Sep 12, 1990  
138 days to decisionK901907 · Product code: **KPN** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k901907/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alarm, Conditioned Response Enuresis (KPN)
Date received	Apr 27, 1990
Decision date	Sep 12, 1990
Days to decision	138 days
Third-party review	No

**APPLICANT**

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Company	<b>Travis Industries, Inc.</b>
Location	Coos Bay, OR, US
Contact	KEITH A BROWN
510(k) history	3 submissions · 3 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901907/>; 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