

K901906 WET-NO-MORE MONITORJul 11, 1990
75 days to decisionK901906 · Product code: **KPN** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k901906/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alarm, Conditioned Response Enuresis (KPN)
Date received	Apr 27, 1990
Decision date	Jul 11, 1990
Days to decision	75 days
Third-party review	No

APPLICANT

Company	Travis Industries, Inc.
Location	Coos Bay, OR, US
510(k) history	3 submissions · 3 cleared · 1990-1990

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Device record: <https://www.510kdatabase.net/k901906/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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