FDA 510(k) Premarket Notification

K901907 PROFESSIONAL MONITOR

CLEARED

Sep 12, 1990
138 days to decision

K901907 · Product code: **KPN** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k901907/

SUBMISSION DETAILS

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

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