FDA 510(k) Premarket Notification

K970077 LIBERTY PLUS SYSTEM(PFS-300)/VAGINAL PROBE, LIBERTY PLUS(PFS-044)

CLEARED

May 23, 1997
134 days to decision

K970077 · Product code: **KPI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k970077/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Non-implantable, For Incontinence (KPI)
Date received	Jan 9, 1997
Decision date	May 23, 1997
Days to decision	134 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Utah Medical Products, Inc.**
Location	Mchenry, IL, US
Contact	NADEEM AHMED
510(k) history	38 submissions · 38 cleared · 1979-2015