FDA 510(k) Premarket Notification

K810944 PENILE ERECTION MONITOR

CLEARED

Apr 23, 1981
16 days to decision

K810944 · Product code: **LIL** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k810944/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Penile Tumescence (LIL)
Date received	Apr 7, 1981
Decision date	Apr 23, 1981
Days to decision	16 days
Third-party review	No

APPLICANT

Company	**Medical Monitoring Systems, Inc.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1981-1984