

K810944 PENILE ERECTION MONITORApr 23, 1981
16 days to decisionK810944 · 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

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Device record: <https://www.510kdatabase.net/k810944/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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