

K831252 TUMISTORE RECORDER 1311May 18, 1983
30 days to decisionK831252 · Product code: **LIL** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k831252/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Penile Tumescence (LIL)
Date received	Apr 18, 1983
Decision date	May 18, 1983
Days to decision	30 days
Third-party review	No

APPLICANT

Company	Life-Tech Intl., Inc.
Location	Walker, MI, US
510(k) history	68 submissions · 66 cleared · 1982-2000

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Device record: <https://www.510kdatabase.net/k831252/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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