

K925931 RT2 RIGIDITY AND TUMESCENCE TESTDec 21, 1992
28 days to decisionK925931 · Product code: **LIL** · Gastroenterology & Urology
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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Penile Tumescence (LIL)
Date received	Nov 23, 1992
Decision date	Dec 21, 1992
Days to decision	28 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Compass Medical Technologies, Inc.
Location	Minneapolis, MN, US
Contact	PHILIP L BROOKS
510(k) history	1 submissions · 1 cleared · 1992-1992

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