FDA 510(k) Premarket Notification

K020154 VT-1

CLEARED

Apr 4, 2002
77 days to decision

K020154 · Product code: **LKY** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k020154/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, External Penile Rigidity (LKY)
Date received	Jan 17, 2002
Decision date	Apr 4, 2002
Days to decision	77 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Pos-T-Vac, Inc.**
Location	Dodge City, KS, US
Contact	DENNIS L BELL
510(k) history	12 submissions · 12 cleared · 1994-2002

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k020154/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026