

K020969 VACURECT VACUUM CONSTRICTION DEVICEJun 5, 2002
71 days to decisionK020969 · Product code: **LKY** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k020969/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, External Penile Rigidity (LKY)
Date received	Mar 26, 2002
Decision date	Jun 5, 2002
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Vacurect Manufacturing (Pty) , Ltd.
Location	North Attleboro, MA, US
Contact	MARY MCNAMARA-CULLINANE
510(k) history	2 submissions · 2 cleared · 2002-2003

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