

**K030913 NANMA VACUUM PUMP**Jun 3, 2003  
71 days to decisionK030913 · Product code: **LKY** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k030913/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, External Penile Rigidity (LKY)
Date received	Mar 24, 2003
Decision date	Jun 3, 2003
Days to decision	71 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Nanma Mfg Co., Ltd.</b>
Location	Crofton, MD, US
Contact	YOLANDA SMITH
510(k) history	9 submissions · 9 cleared · 2000-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030913/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 30, 2026