FDA 510(k) Premarket Notification

K030913 NANMA VACUUM PUMP

CLEARED

Jun 3, 2003
71 days to decision

K030913 · Product code: **LKY** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k030913/

SUBMISSION DETAILS

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

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