FDA 510(k) Premarket Notification

K860698 THE BOOSTER

CLEARED

Feb 3, 1989
1074 days to decision

K860698 · Product code: **LKY** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k860698/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, External Penile Rigidity (LKY)
Date received	Feb 25, 1986
Decision date	Feb 3, 1989
Days to decision	1074 days
Third-party review	No

APPLICANT

Company	**Knapps Corp.**
Location	Newark, CA, US
Contact	OLIVER L KNAPPS
510(k) history	1 submissions · 1 cleared · 1989-1989