

K860698 THE BOOSTER

Feb 3, 1989

1074 days to decision

K860698 · Product code: **LKY** · Gastroenterology & UrologySource: <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

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