

K900559 RESPONSE EXTERNAL PENILE RIGIDITY DEVICEMay 2, 1990
84 days to decisionK900559 · Product code: **LKY** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k900559/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, External Penile Rigidity (LKY)
Date received	Feb 7, 1990
Decision date	May 2, 1990
Days to decision	84 days
Third-party review	No

APPLICANT

Company	Smith-Collins Pharmaceutical, Inc.
Location	West Chester, PA, US
Contact	WILLIAM KNEPSHIELD
510(k) history	1 submissions · 1 cleared · 1990-1990

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