FDA 510(k) Premarket Notification

K782093 DETECTOR PAD

CLEARED

Feb 21, 1979
69 days to decision

K782093 · Product code: **KPN** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k782093/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alarm, Conditioned Response Enuresis (KPN)
Date received	Dec 14, 1978
Decision date	Feb 21, 1979
Days to decision	69 days
Third-party review	No

APPLICANT

Company	**Halperin Institute, Ltd. The**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1979-1979