FDA 510(k) Premarket Notification

# K802173 URIHESIVE STRIPS

**CLEARED**

Sep 26, 1980
17 days to decision

K802173 · Product code: **KNX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k802173/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Collector, Urine, (and Accessories) For Indwelling Catheter (KNX) |
| Date received | Sep 9, 1980 |
| Decision date | Sep 26, 1980 |
| Days to decision | 17 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **E. R. Squibb & Sons, Inc.** |
| Location | New York, NY, US |
| 510(k) history | 32 submissions · 32 cleared · 1977-1982 |