FDA 510(k) Premarket Notification

# K810690 CURITY 100% SILICONE SAFETY INDICATOR

**CLEARED**

Apr 23, 1981
41 days to decision

K810690 · Product code: **KOD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k810690/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Urological (KOD) |
| Date received | Mar 13, 1981 |
| Decision date | Apr 23, 1981 |
| Days to decision | 41 days |
| Third-party review | No |

## APPLICANT

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|---|---|
| Company | **The Kendal Co.** |
| Location | Mchenry, IL, US |
| 510(k) history | 63 submissions · 60 cleared · 1980-2000 |