FDA 510(k) Premarket Notification

# K060268 PEDIATRIC URINARY CATHETER

**CLEARED**

Feb 24, 2006
23 days to decision

K060268 · Product code: **GBM** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k060268/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Urethral (GBM) |
| Date received | Feb 1, 2006 |
| Decision date | Feb 24, 2006 |
| Days to decision | 23 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Clay Kennard** |
| Location | Oklahoma City, OK, US |
| Contact | CLAY KENNARD |
| 510(k) history | 1 submissions · 1 cleared · 2006-2006 |