FDA 510(k) Premarket Notification

# K880099 VAN-TEC BLADDER EVACUATOR

**CLEARED**

Mar 2, 1988
50 days to decision

K880099 · Product code: **KQT** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k880099/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Evacuator, Gastro-urology (KQT) |
| Date received | Jan 12, 1988 |
| Decision date | Mar 2, 1988 |
| Days to decision | 50 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Van-Tec, Inc.** |
| Location | Walker, MI, US |
| Contact | VANCE, JR. |
| 510(k) history | 52 submissions · 52 cleared · 1982-1988 |

**510k Database** - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k880099/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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