

K810368 VANCE PERCUTANEOUS MALECOT NEPHROSTOMYMar 20, 1981
36 days to decisionK810368 · Product code: **KOB** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k810368/>**SUBMISSION DETAILS**

| | |
|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Suprapubic (and Accessories) (KOB) |
| Date received | Feb 12, 1981 |
| Decision date | Mar 20, 1981 |
| Days to decision | 36 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Vance Products, Inc. |
| Location | Mchenry, IL, US |
| 510(k) history | 17 submissions · 17 cleared · 1978-1982 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k810368/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 3, 2026