

K070052 MET-HEAD MODEL# MHDD-XX:CC DIAMETER, CC CANNULATION

Mar 28, 2007
83 days to decision

K070052 · Product code: **KWD** · Orthopedic
Source: <https://www.510kdatabase.net/k070052/>

SUBMISSION DETAILS

| | |
|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Prosthesis, Toe, Hemi-, Phalangeal (KWD) |
| Date received | Jan 4, 2007 |
| Decision date | Mar 28, 2007 |
| Days to decision | 83 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Vilex, Inc. |
| Location | Pleasant Hills, PA, US |
| Contact | ABRAHAM LAVI |
| 510(k) history | 13 submissions · 12 cleared · 1998-2014 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k070052/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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