

**K801284 LOW PROFILE LEAD CONNECTOR MODIFICATION
(4002)**Dec 1, 1980
186 days to decisionK801284 · Product code: **DTB** · Cardiovascular
Source: <https://www.510kdatabase.net/k801284/>**SUBMISSION DETAILS**

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|-----------------------|-------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Permanent Pacemaker Electrode (DTB) |
| Date received | May 29, 1980 |
| Decision date | Dec 1, 1980 |
| Days to decision | 186 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Medtronic Vascular |
| Location | Walker, MI, US |
| Contact | KEITH J PROCTOR |
| 510(k) history | 475 submissions · 453 cleared · 1977-2023 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k801284/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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