

K020003 ELECTROCARDIOGRAPH (ECG) ELECTRODE, NON-STERILENov 7, 2002
309 days to decisionK020003 · Product code: **DRX** · Cardiovascular
Source: <https://www.510kdatabase.net/k020003/>**SUBMISSION DETAILS**

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|-----------------------|-------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Electrode, Electrocardiograph (DRX) |
| Date received | Jan 2, 2002 |
| Decision date | Nov 7, 2002 |
| Days to decision | 309 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Bio Protech, Inc. |
| Location | Sparks, NV, US |
| Contact | JANNA P TUCKER |
| 510(k) history | 17 submissions · 17 cleared · 2002-2019 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k020003/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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