

K800201 ESOPHAGEAL PILL-ELECTRODEFeb 26, 1980
27 days to decisionK800201 · Product code: **DRX** · Cardiovascular
Source: <https://www.510kdatabase.net/k800201/>**SUBMISSION DETAILS**

| | |
|-----------------------|-------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Electrode, Electrocardiograph (DRX) |
| Date received | Jan 30, 1980 |
| Decision date | Feb 26, 1980 |
| Days to decision | 27 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Arzbaecher & Co. |
| Location | Walker, MI, US |
| 510(k) history | 2 submissions · 2 cleared · 1979-1980 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k800201/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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