FDA 510(k) Premarket Notification

# K780117 CARDIOMETER

**CLEARED**

Feb 3, 1978
11 days to decision

K780117 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k780117/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT) |
| Date received | Jan 23, 1978 |
| Decision date | Feb 3, 1978 |
| Days to decision | 11 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Med General** |
| Location | Mchenry, IL, US |
| 510(k) history | 12 submissions · 12 cleared · 1977-1979 |