FDA 510(k) Premarket Notification

# K810331 DUO-TEL

**CLEARED**

Mar 20, 1981
39 days to decision

K810331 · Product code: **DRG** · Cardiovascular
Source: https://www.510kdatabase.net/k810331/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG) |
| Date received | Feb 9, 1981 |
| Decision date | Mar 20, 1981 |
| Days to decision | 39 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **General Electric Co.** |
| Location | Mchenry, IL, US |
| 510(k) history | 254 submissions · 254 cleared · 1976-2011 |