

K810174 RESPIRATION FUNCTION BLOCKSMar 17, 1981
54 days to decisionK810174 · Product code: **BZQ** · Anesthesiology
Source: <https://www.510kdatabase.net/k810174/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Monitor, Breathing Frequency (BZQ) |
| Date received | Jan 22, 1981 |
| Decision date | Mar 17, 1981 |
| Days to decision | 54 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Honeywell, Inc. |
| Location | Mchenry, IL, US |
| 510(k) history | 69 submissions · 69 cleared · 1976-1990 |

Honeywell, Inc. is an American multinational conglomerate headquartered in Charlotte, North Carolina. The company operates across aerospace, building automation, industrial automation, and energy solutions. Honeywell's medical device regulatory history spans from 1976 to 1990. The company received FDA 510(k) clearances from total submissions. Cardiovascular devices represented the dominant focus, accounting for approximately 75% of submissions. This historical record reflects the company's past involvement in patient monitoring systems, defibrillators, and related cardiov...

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