

K812667 MEDDARS/ULTRA IMAGER RESPIRATION MODULEOct 23, 1981
32 days to decisionK812667 · Product code: **BZQ** · AnesthesiologySource: <https://www.510kdatabase.net/k812667/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Breathing Frequency (BZQ)
Date received	Sep 21, 1981
Decision date	Oct 23, 1981
Days to decision	32 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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Device record: <https://www.510kdatabase.net/k812667/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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