

K830914 EXTERNAL VALVE FUNCTION BLOCKAug 12, 1983
141 days to decisionK830914 · Product code: **DRQ** · Cardiovascular
Source: <https://www.510kdatabase.net/k830914/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Amplifier And Signal Conditioner, Transducer Signal (DRQ)
Date received	Mar 24, 1983
Decision date	Aug 12, 1983
Days to decision	141 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/k830914/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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