

**K822590 PRESSURE ISOLATION AMPLIFIER #9813 517**Oct 4, 1982  
27 days to decisionK822590 · Product code: **DRJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k822590/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Signal Isolation (DRJ)
Date received	Sep 7, 1982
Decision date	Oct 4, 1982
Days to decision	27 days
Third-party review	No

**APPLICANT**

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Company	<b>Honeywell, Inc.</b>
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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