

**K853501 HONEYWELL ELECTRONIC VENTILATOR EV-A**Apr 4, 1986  
227 days to decisionK853501 · Product code: **CBK** · Anesthesiology  
Source: <https://www.510kdatabase.net/k853501/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Continuous, Facility Use (CBK)
Date received	Aug 20, 1985
Decision date	Apr 4, 1986
Days to decision	227 days
Third-party review	No

**APPLICANT**

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Company	<b>Honeywell, Inc.</b>
Location	Mchenry, IL, US
Contact	MARTIN KUTIK
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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