

K822926 MODEL LS-85 & MEDEL 1828 LINESCAN VISCONov 24, 1982
54 days to decision

K822926 · Cardiovascular

Source: <https://www.510kdatabase.net/k822926/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Oct 1, 1982
Decision date	Nov 24, 1982
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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Device record: <https://www.510kdatabase.net/k822926/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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