

K842210 IVENT 302VC DUAL CONTROLLERJul 25, 1984
51 days to decisionK842210 · Product code: **LDR** · General Hospital
Source: <https://www.510kdatabase.net/k842210/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Jun 4, 1984
Decision date	Jul 25, 1984
Days to decision	51 days
Third-party review	No

APPLICANT

Company	Ivent Corp.
Location	Walker, MI, US
510(k) history	3 submissions · 3 cleared · 1983-1985

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Device record: <https://www.510kdatabase.net/k842210/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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