FDA 510(k) Premarket Notification

K842210 IVENT 302VC DUAL CONTROLLER

CLEARED

Jul 25, 1984
51 days to decision

K842210 · 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