

K051253 MODIFICATION TO: MULTIRATE INFUSOR SV)WITH AND WITHOUT PCM), MULTIRATE INFUSOR LV (WITH AND WITHOUT PCM)

Jul 15, 2005
60 days to decision

K051253 · Product code: **MEA** · General Hospital
Source: <https://www.510kdatabase.net/k051253/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion, Pca (MEA)
Date received	May 16, 2005
Decision date	Jul 15, 2005
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Baxter Healthcare Corp
Location	Mchenry, IL, US
Contact	NANETTE HEDDEN
510(k) history	505 submissions · 496 cleared · 1977-2019

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k051253/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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