

**K002739 INFUSOR PATIENT CONTROL MODULE, 2ML, MODEL  
2C1067K**Sep 11, 2000  
10 days to decisionK002739 · Product code: **MEA** · General Hospital  
Source: <https://www.510kdatabase.net/k002739/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Pump, Infusion, Pca (MEA)
Date received	Sep 1, 2000
Decision date	Sep 11, 2000
Days to decision	10 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Baxter Healthcare Corp</b>
Location	Mchenry, IL, US
Contact	VICKI DREWS
510(k) history	505 submissions · 496 cleared · 1977-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k002739/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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