

**K030661 STRYKER PAINPUMP2**May 30, 2003  
88 days to decisionK030661 · Product code: **MEA** · General Hospital  
Source: <https://www.510kdatabase.net/k030661/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion, Pca (MEA)
Date received	Mar 3, 2003
Decision date	May 30, 2003
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Instruments</b>
Location	Kalamazoo, MI, US
Contact	NICOLE PETTY
510(k) history	73 submissions · 73 cleared · 1994-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030661/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 28, 2026