

**K944479 PM3300 INTERMITTENT VACUUM**Dec 22, 1994  
100 days to decisionK944479 · Product code: **KDP** · General Hospital  
Source: <https://www.510kdatabase.net/k944479/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Regulator, Vacuum (KDP)
Date received	Sep 13, 1994
Decision date	Dec 22, 1994
Days to decision	100 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Precision Medical, Inc.</b>
Location	Mchenry, IL, US
Contact	MICHAEL A KRUPA
510(k) history	30 submissions · 30 cleared · 1984-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k944479/> 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 May 24, 2026