

K922817 PM3000 & PM3100 VACUUM REGULATORMar 4, 1993
266 days to decisionK922817 · Product code: **KDP** · General Hospital
Source: <https://www.510kdatabase.net/k922817/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Regulator, Vacuum (KDP)
Date received	Jun 11, 1992
Decision date	Mar 4, 1993
Days to decision	266 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

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