FDA 510(k) Premarket Notification

K944479 PM3300 INTERMITTENT VACUUM

CLEARED

Dec 22, 1994
100 days to decision

K944479 · Product code: **KDP** · General Hospital
Source: https://www.510kdatabase.net/k944479/

SUBMISSION DETAILS

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

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