

K060095 MODIFICATION TO: VACLOK SYRINGEFeb 10, 2006
28 days to decisionK060095 · Product code: **FMF** · General HospitalSource: <https://www.510kdatabase.net/k060095/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Syringe, Piston (FMF)
Date received	Jan 13, 2006
Decision date	Feb 10, 2006
Days to decision	28 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Wrightmedicaltechnologyinc
Location	Arlington, TN, US
Contact	WESLEY L REED
510(k) history	302 submissions · 291 cleared · 1993-2023

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