

K960882 CONTROL SYRINGEJul 8, 1996
139 days to decisionK960882 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k960882/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Feb 20, 1996
Decision date	Jul 8, 1996
Days to decision	139 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Louisville Laboratories, Inc.
Location	Weston, MA, US
Contact	DAVID PHELPS
510(k) history	12 submissions · 12 cleared · 1989-1998

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