

K052445 DUOPROSS NEEDLEDec 5, 2005
89 days to decisionK052445 · Product code: **FMI** · General Hospital
Source: <https://www.510kdatabase.net/k052445/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Sep 7, 2005
Decision date	Dec 5, 2005
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Duopross Meditech Corporation
Location	Albuquerque, NM, US
Contact	FRANK FERGUSON
510(k) history	2 submissions · 2 cleared · 2005-2005

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