

K010958 FISTULOK FISTULA NEEDLE PROTECTIONMay 18, 2001
49 days to decisionK010958 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k010958/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Mar 30, 2001
Decision date	May 18, 2001
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Diasol, Inc.
Location	North Hollywood, CA, US
Contact	MONICA ABELES
510(k) history	10 submissions · 10 cleared · 1999-2014

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