

**K011004 NEEDLELOK HYPODERMIC NEEDLE PROTECTION**May 24, 2001  
50 days to decisionK011004 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k011004/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Apr 4, 2001
Decision date	May 24, 2001
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k011004/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 7, 2026