

**K981619 VACUETTE MULTIPLE SAMPLE LUER ADAPTER**Jul 15, 1998  
70 days to decisionK981619 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k981619/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	May 6, 1998
Decision date	Jul 15, 1998
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Greiner Meditech, Inc.</b>
Location	Bel Air, MD, US
Contact	DOUGLAS L HARRIS
510(k) history	10 submissions · 10 cleared · 1997-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k981619/> 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 29, 2026