

**K943137 B-D/BFI MAIL DISPOSAL SERVICE**Nov 7, 1994  
131 days to decisionK943137 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k943137/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jun 29, 1994
Decision date	Nov 7, 1994
Days to decision	131 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Med-Safe Systems, Inc.</b>
Location	Oceanside, CA, US
Contact	VIRGINIA SHILLINGTON
510(k) history	8 submissions · 8 cleared · 1994-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k943137/> 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 July 6, 2026