

**K955748 DOYLE EXTRACTOR**Mar 14, 1996  
90 days to decisionK955748 · Product code: **FMI** · General Hospital  
Source: <https://www.510kdatabase.net/k955748/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Dec 15, 1995
Decision date	Mar 14, 1996
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Safetech Intl., Inc.</b>
Location	North Attleboro, MA, US
Contact	DIANE MINEAR
510(k) history	1 submissions · 1 cleared · 1996-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k955748/> 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