

**K982018 LIGHT SABER ASPIRATION NEEDLE**Aug 7, 1998  
60 days to decisionK982018 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k982018/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jun 8, 1998
Decision date	Aug 7, 1998
Days to decision	60 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Minrad, Inc.</b>
Location	Washington, DC, US
Contact	JOHN MCNEIRNEY
510(k) history	14 submissions · 14 cleared · 1997-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k982018/> 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 May 19, 2026