

**K981140 MINRAD DRTS LIGHT SABER SYRINGE**Jun 25, 1998  
87 days to decisionK981140 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k981140/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Mar 30, 1998
Decision date	Jun 25, 1998
Days to decision	87 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k981140/> 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