

**K951808 MED-TEC REDI-FOAM**Jul 6, 1995  
78 days to decisionK951808 · Product code: **IYE** · Radiology  
Source: <https://www.510kdatabase.net/k951808/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Apr 19, 1995
Decision date	Jul 6, 1995
Days to decision	78 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Medtec, Inc.</b>
Location	Orange City, IA, US
Contact	DONALD RIIBE
510(k) history	41 submissions · 41 cleared · 1993-2014

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k951808/> 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 17, 2026