

**K062956 EZ-MIO, EZ-IO DISTAL TIBIA**Dec 5, 2006  
67 days to decisionK062956 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k062956/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Sep 29, 2006
Decision date	Dec 5, 2006
Days to decision	67 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Vidacare Corporation</b>
Location	Irvine, CA, US
Contact	GRACE HOLLAND
510(k) history	19 submissions · 19 cleared · 2004-2014

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k062956/> 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 June 7, 2026