

**K091140 EZ-MIO DISTAL TIBIA, EZ-IO DISTAL TIBIA, VIDAPORT
INTRAOSSEOUS INFUSION SYSTEM, EZ-IO**Oct 14, 2009
177 days to decisionK091140 · Product code: FMI · General Hospital
Source: <https://www.510kdatabase.net/k091140/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Apr 20, 2009
Decision date	Oct 14, 2009
Days to decision	177 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k091140/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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