FDA 510(k) Premarket Notification

K062956 EZ-MIO, EZ-IO DISTAL TIBIA

CLEARED

Dec 5, 2006
67 days to decision

K062956 · Product code: **FMI** · General Hospital
Source: 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	**Vidacare Corporation**
Location	Irvine, CA, US
Contact	GRACE HOLLAND
510(k) history	19 submissions · 19 cleared · 2004-2014