FDA 510(k) Premarket Notification

K062833 POWERED EZ-IO BONE MARROW ASPIRATION SYSTEM

CLEARED

Nov 30, 2006
70 days to decision

K062833 · Product code: **FCF** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k062833/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy, Mechanical, Gastrointestinal (FCF)
Date received	Sep 21, 2006
Decision date	Nov 30, 2006
Days to decision	70 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