FDA 510(k) Premarket Notification

CLEARED

K081713 VERTEBRAL ACCESS SYSTEM BY VIDACARE

Nov 21, 2008
157 days to decision

K081713 · Product code: **MOQ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k081713/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Battery, Replacement, Rechargeable (MOQ)
Date received	Jun 17, 2008
Decision date	Nov 21, 2008
Days to decision	157 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