FDA 510(k) Premarket Notification

CLEARED

K051848 SALUTE II DISPOSABLE FIXATION DEVICE, MODELS 0113070, 0113072, 0113073, 0113077, 0113079

Aug 11, 2005
35 days to decision

K051848 · Product code: **GDW** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k051848/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Implantable (GDW)
Date received	Jul 7, 2005
Decision date	Aug 11, 2005
Days to decision	35 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Davol Inc., Sub. C. R. Bard, Inc.**
Location	Canston, RI, US
Contact	LUCINDA L FOX
510(k) history	24 submissions · 24 cleared · 1994-2010