FDA 510(k) Premarket Notification

K083467 REBOUND HRDV

CLEARED

Apr 23, 2009
150 days to decision

K083467 · Product code: **FTL** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k083467/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Nov 24, 2008
Decision date	Apr 23, 2009
Days to decision	150 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Minnesota Medical Development, Inc.**
Location	Stillwater, MN, US
Contact	STEVE NUSS
510(k) history	3 submissions · 3 cleared · 2007-2009