FDA 510(k) Premarket Notification

K080393 MODIFICATION TO REBOUND HRD

CLEARED

Mar 13, 2008
29 days to decision

K080393 · Product code: **FTL** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k080393/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Feb 13, 2008
Decision date	Mar 13, 2008
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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