FDA 510(k) Premarket Notification

K083752 GLUSEAL 90, MODEL GLU9010

CLEARED

Feb 23, 2009
68 days to decision

K083752 · Product code: **KMF** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k083752/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bandage, Liquid (KMF)
Date received	Dec 17, 2008
Decision date	Feb 23, 2009
Days to decision	68 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Glustitch, Inc.**
Location	Delta, CA
Contact	DON BLACKLOCK
510(k) history	5 submissions · 5 cleared · 2001-2016