

K083752 GLUSEAL 90, MODEL GLU9010Feb 23, 2009
68 days to decisionK083752 · Product code: **KMF** · General & Plastic SurgerySource: <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 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k083752/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 3, 2026