

K100630 G-FORCE PEEK SUTURE ANCHOR SYSTEMJun 11, 2010
98 days to decisionK100630 · Product code: **MBI** · Orthopedic
Source: <https://www.510kdatabase.net/k100630/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fastener, Fixation, Nondegradable, Soft Tissue (MBI)
Date received	Mar 5, 2010
Decision date	Jun 11, 2010
Days to decision	98 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Wrightmedicaltechnologyinc
Location	Arlington, TN, US
Contact	KELLEN HILLS
510(k) history	302 submissions · 291 cleared · 1993-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k100630/> 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 May 18, 2026