

K101175 KFX APPIANFX PEEK FEMORAL IMPLANT WITH INSERTERMay 20, 2010
23 days to decisionK101175 · Product code: **MBI** · Orthopedic
Source: <https://www.510kdatabase.net/k101175/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Fastener, Fixation, Nondegradable, Soft Tissue (MBI)
Date received	Apr 27, 2010
Decision date	May 20, 2010
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Kfx Medical
Location	Carlsbad, CA, US
Contact	GAYLE HIROTA
510(k) history	7 submissions · 7 cleared · 2006-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k101175/> 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 14, 2026