FDA 510(k) Premarket Notification

K023088 BONE GRAFT SYRINGE, MODEL 8600-00X0

CLEARED

Oct 2, 2002
15 days to decision

K023088 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k023088/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Sep 17, 2002
Decision date	Oct 2, 2002
Days to decision	15 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Wrightmedicaltechnologyinc**
Location	Arlington, TN, US
Contact	ROGER D BROWN
510(k) history	302 submissions · 291 cleared · 1993-2023

REGULATORY CONSULTANT

Consulting firm	**Tuv America, Inc.**
Contact	MARK JOB

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov