FDA 510(k) Premarket Notification

K080088 BIOMET SPORTS MEDICINE ANCHOR DEVICES AND ZIPLOOP CONSTRUCTS

CLEARED

Jun 11, 2008
149 days to decision

K080088 · Product code: **HWC** · Orthopedic
Source: https://www.510kdatabase.net/k080088/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	Jan 14, 2008
Decision date	Jun 11, 2008
Days to decision	149 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Biomet Manufacturing, Inc.**
Location	Warsaw, IN, US
Contact	ROBERT R FRIDDLE
510(k) history	32 submissions · 32 cleared · 1999-2013

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k080088/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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