FDA 510(k) Premarket Notification

K082293 SURGVIEW INTEGRATED VISUALIZATION SYSTEM

CLEARED

Sep 9, 2008
28 days to decision

K082293 · Product code: **HRX** · Orthopedic
Source: https://www.510kdatabase.net/k082293/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Aug 12, 2008
Decision date	Sep 9, 2008
Days to decision	28 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Biovision Technologies, LLC**
Location	Golden, CO, US
510(k) history	3 submissions · 3 cleared · 2008-2015

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k082293/ 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 17, 2026