

**K081051 SPINEVU ENDOSCOPIC SPINE SYSTEM (SESS),
SPINEVU MINISCOPE**Aug 7, 2008
115 days to decisionK081051 · Product code: **HRX** · Orthopedic
Source: <https://www.510kdatabase.net/k081051/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Apr 14, 2008
Decision date	Aug 7, 2008
Days to decision	115 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Spine View, Inc.
Location	Fremont, CA, US
Contact	ROY CHIN
510(k) history	9 submissions · 9 cleared · 2008-2016

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k081051/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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