

K081165 ELLIPSE LUMBAR POSTERIOR OSTEOSYNTHESIS SYSTEMDec 17, 2008
237 days to decisionK081165 · Product code: **MNH** · Orthopedic
Source: <https://www.510kdatabase.net/k081165/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spondylolisthesis Spinal Fixation (MNH)
Date received	Apr 24, 2008
Decision date	Dec 17, 2008
Days to decision	237 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Spineart
Location	Geneva, CH
Contact	FRANCK PENNESI
510(k) history	44 submissions · 44 cleared · 2008-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k081165/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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