

**K141835 ROMEO POSTERIOR OSTEOSYNTHESIS SYSTEM  
PREBENT CROSS-CONNECTORS**Sep 10, 2014  
65 days to decisionK141835 · Product code: **MNI** · Orthopedic  
Source: <https://www.510kdatabase.net/k141835/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Orthosis, Spinal Pedicle Fixation (MNI)
Date received	Jul 7, 2014
Decision date	Sep 10, 2014
Days to decision	65 days
Third-party review	No
Summary / Statement	Summary
Other names	PREBENT RODS

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k141835/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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