

**K193580 VERASENSE for Zimmer Biomet Persona CR C-D/3-9  
Left, VERASENSE for Zimmer Biomet Persona CR C-D/3-9  
Right, VERASENSE for Zimmer Biomet Persona CR E-F/3-11  
Left, VERASENSE for Zimmer Biomet Persona CR E-F/3-11  
Right, VERASENSE for Zimmer Biomet Persona CR G-H/7-12  
Left**

Apr 1, 2020  
100 days to decision

K193580 · Product code: **ONN** · Orthopedic  
Source: <https://www.510kdatabase.net/k193580/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intraoperative Orthopedic Joint Assessment Aid (ONN)
Date received	Dec 23, 2019
Decision date	Apr 1, 2020
Days to decision	100 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Orthosensor, Inc.</b>
Location	Appollo Beach, FL, US
Contact	Deborah Johnson
510(k) history	8 submissions · 8 cleared · 2009-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k193580/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).

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