

**K253992 Veritas Reverse Total Shoulder System**May 7, 2026  
146 days to decisionK253992 · Product code: **PHX** · Orthopedic  
Source: <https://www.510kdatabase.net/k253992/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shoulder Prosthesis, Reverse Configuration (PHX)
Date received	Dec 12, 2025
Decision date	May 7, 2026
Days to decision	146 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Restor3d</b>
Location	Durham, NC, US
Contact	Brianna Prindle
510(k) history	12 submissions · 12 cleared · 2020-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k253992/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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