FDA 510(k) Premarket Notification

K253992 Veritas Reverse Total Shoulder System

CLEARED

May 7, 2026
146 days to decision

K253992 · Product code: **PHX** · Orthopedic
Source: https://www.510kdatabase.net/k253992/

SUBMISSION DETAILS

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

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