

K211359 Tornier Perform™ Patient-Matched Primary Reversed Glenoid and BLUEPRINT™ Patient Specific InstrumentationNov 12, 2021
193 days to decisionK211359 · Product code: **PHX** · Orthopedic
Source: <https://www.510kdatabase.net/k211359/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shoulder Prosthesis, Reverse Configuration (PHX)
Date received	May 3, 2021
Decision date	Nov 12, 2021
Days to decision	193 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Tornier, Inc.
Location	Beverly, MA, US
Contact	Lisa Stahl
510(k) history	51 submissions · 51 cleared · 2000-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k211359/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 13, 2026