

K103251 DUOCENTRIC REVERSEDOct 28, 2011
359 days to decisionK103251 · Product code: **PHX** · Orthopedic
Source: <https://www.510kdatabase.net/k103251/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shoulder Prosthesis, Reverse Configuration (PHX)
Date received	Nov 3, 2010
Decision date	Oct 28, 2011
Days to decision	359 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Aston Medical
Location	Santa Barbara, CA, US
Contact	CATHERINE GLOSTER
510(k) history	3 submissions · 3 cleared · 2011-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k103251/> 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 17, 2026