

K101704 ZIMMER DTO PIN PRESS INSTRUMENTAug 18, 2010
62 days to decisionK101704 · Product code: **NQP** · Orthopedic
Source: <https://www.510kdatabase.net/k101704/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Posterior Metal/polymer Spinal System, Fusion (NQP)
Date received	Jun 17, 2010
Decision date	Aug 18, 2010
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Zimmer Spine, Inc.
Location	Minneapolis, MN, US
Contact	TIM CRABTREE
510(k) history	38 submissions · 35 cleared · 2004-2016

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k101704/> 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