FDA 510(k) Premarket Notification

K101704 ZIMMER DTO PIN PRESS INSTRUMENT

CLEARED

Aug 18, 2010
62 days to decision

K101704 · 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