

**K111983 VISTA-S DEVICE MODEL 08/06-401-XXXXX,  
08/06-402-XXXXXX**

Nov 18, 2011  
129 days to decision

K111983 · Product code: **ODP** · Orthopedic  
Source: <https://www.510kdatabase.net/k111983/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Jul 12, 2011
Decision date	Nov 18, 2011
Days to decision	129 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Zimmer Trabecular Metal Technology</b>
Location	Parsippany, NJ, US
Contact	KATHLEEN RUTHERFORD
510(k) history	11 submissions · 11 cleared · 2007-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k111983/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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