

**K070218 VERTIFLEX OCTANE PEEK VERTEBRAL BODY REPLACEMENT (PEEK VBR)**Apr 6, 2007  
73 days to decisionK070218 · Product code: **MQP** · Orthopedic  
Source: <https://www.510kdatabase.net/k070218/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Jan 23, 2007
Decision date	Apr 6, 2007
Days to decision	73 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vertiflex (Tm), Incorporated</b>
Location	Carlsbad, CA, US
Contact	STEVE REITZLER
510(k) history	2 submissions · 2 cleared · 2007-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k070218/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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