

**K102401 HEMI IMPLANT**Jul 20, 2011  
330 days to decisionK102401 · Product code: **KWD** · Orthopedic  
Source: <https://www.510kdatabase.net/k102401/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Toe, Hemi-, Phalangeal (KWD)
Date received	Aug 24, 2010
Decision date	Jul 20, 2011
Days to decision	330 days
Third-party review	No
Summary / Statement	Summary
Other names	MINI HEMI IMPLANT

**APPLICANT**

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Company	<b>Vilex, Inc.</b>
Location	Pleasant Hills, PA, US
Contact	SYLVIA SOUTHARD
510(k) history	13 submissions · 12 cleared · 1998-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k102401/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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