

**K021025 MODIFICATION TO HEDROCEL VERTEBRAL BODY REPLACEMENT, MODEL XX-YYY-ZZZZ**

May 20, 2002  
52 days to decision

K021025 · Product code: **MQP** · Orthopedic  
Source: <https://www.510kdatabase.net/k021025/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Mar 29, 2002
Decision date	May 20, 2002
Days to decision	52 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Implex Corp.</b>
Location	Allendale, NJ, US
Contact	ROBERT POGGIE
510(k) history	65 submissions · 61 cleared · 1993-2005

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

Device record: <https://www.510kdatabase.net/k021025/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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