

**K020221 IQL BASIC FRAGMENT PLATE SET, IQL BASIC FRAGMENT SCREW SET, IQL SMALL FRAGMENT SET, IQL MINI FRAGMENT SET**

Apr 2, 2002  
70 days to decision

K020221 · Product code: **HRS** · Orthopedic  
Source: <https://www.510kdatabase.net/k020221/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Fixation, Bone (HRS)
Date received	Jan 22, 2002
Decision date	Apr 2, 2002
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Biomet Orthopedics, Inc.</b>
Location	Warsaw, IN, US
Contact	LOMNIE WITHAM
510(k) history	34 submissions · 34 cleared · 2000-2008

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

Device record: <https://www.510kdatabase.net/k020221/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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