

**K221920 ARIX Small Fragment System**Aug 24, 2022  
54 days to decisionK221920 · Product code: **HRS** · Orthopedic  
Source: <https://www.510kdatabase.net/k221920/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Fixation, Bone (HRS)
Date received	Jul 1, 2022
Decision date	Aug 24, 2022
Days to decision	54 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Jeil Medical Corporation</b>
Location	Deer Field, IL, US
Contact	Bora Kim
510(k) history	53 submissions · 53 cleared · 2002-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k221920/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 13, 2026