

**K894016 MAGNETIC RESONANCE DIAGNOSTIC DEVICE  
MRP-20**Nov 22, 1989  
169 days to decisionK894016 · Product code: **LNH** · Radiology  
Source: <https://www.510kdatabase.net/k894016/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jun 6, 1989
Decision date	Nov 22, 1989
Days to decision	169 days
Third-party review	No

**APPLICANT**

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Company	<b>Summit World Trade Corp.</b>
Location	Hudson, OH, US
Contact	WILLIAM J BISHOP
510(k) history	10 submissions · 10 cleared · 1989-1991

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k894016/>; 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 21, 2026