

**K983110 FLEXART,FLEXART/HYPER, VISART, VISART/HYPER**Feb 25, 1999  
174 days to decisionK983110 · Product code: **LNH** · Radiology  
Source: <https://www.510kdatabase.net/k983110/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Sep 4, 1998
Decision date	Feb 25, 1999
Days to decision	174 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Toshiba America Mri, Inc.</b>
Location	South San Francisco, CA, US
Contact	KEN NEHMER
510(k) history	68 submissions · 68 cleared · 1990-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k983110/> 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 19, 2026