

**K872468 JOVI MYELOGRAPHY HARNESS**Jul 1, 1987  
8 days to decisionK872468 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k872468/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Jun 23, 1987
Decision date	Jul 1, 1987
Days to decision	8 days
Third-party review	No

**APPLICANT**

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Company	<b>Jovi, Inc.</b>
Location	Dallas, TX, US
Contact	DENNIS T GRIGGS
510(k) history	1 submissions · 1 cleared · 1987-1987

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k872468/>; 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