

**K974112 VACUDAP, MODEL 2000, 2001**Jan 29, 1998  
90 days to decisionK974112 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k974112/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Oct 31, 1997
Decision date	Jan 29, 1998
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Victoreen, Inc.</b>
Location	Mchenry, IL, US
Contact	LINDA S NASH
510(k) history	40 submissions · 40 cleared · 1979-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k974112/> 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