

**K030870 WBR, MODEL HR**Apr 9, 2003  
21 days to decisionK030870 · Product code: **KPS** · Radiology  
Source: <https://www.510kdatabase.net/k030870/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Tomography, Computed, Emission (KPS)
Date received	Mar 19, 2003
Decision date	Apr 9, 2003
Days to decision	21 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>U.C.G. Technologies , Ltd.</b>
Location	Haifa, IL
Contact	DAN LAOR
510(k) history	2 submissions · 2 cleared · 2003-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030870/> 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