

**K031248 CONSULTIVA REPORT STATION, MODEL RS-1**Jun 20, 2003  
63 days to decisionK031248 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k031248/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Apr 18, 2003
Decision date	Jun 20, 2003
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Miramedica, Inc.</b>
Location	Los Gatos, CA, US
Contact	WIDO MENHARDT
510(k) history	1 submissions · 1 cleared · 2003-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k031248/> 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