

**K920099 GYMMAN**Mar 9, 1992  
60 days to decisionK920099 · Product code: **DRT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k920099/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jan 9, 1992
Decision date	Mar 9, 1992
Days to decision	60 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Lumiscop Co., Inc.</b>
Location	Edison, NJ, US
Contact	STEPHEN L WHITTENBU
510(k) history	13 submissions · 13 cleared · 1988-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920099/> 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 26, 2026