

**K912444 MICROBACT 12A SYSTEM, 12B SYSTEM, AND 24E SYSTEM**Feb 18, 1992  
260 days to decisionK912444 · Product code: **LGM** · General Hospital  
Source: <https://www.510kdatabase.net/k912444/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chamber, Patient Isolation (LGM)
Date received	Jun 3, 1991
Decision date	Feb 18, 1992
Days to decision	260 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Radiometer America, Inc.</b>
Location	Mchenry, IL, US
Contact	KATHLEEN M ONTKO
510(k) history	50 submissions · 50 cleared · 1983-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k912444/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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