

## Response Biomedical Corporation Announces Second Quarter Results

### FOR IMMEDIATE RELEASE

**Vancouver, British Columbia, August 10, 2011** – Response Biomedical Corporation (TSX: RBM, OTCBB: RPIBF) reported financial results for the second quarter ended June 30, 2011. For the three month period ended June 30, 2011, total revenues increased 23% to \$2,771,104 compared to \$2,255,513, gross margin percentage from product sales increased to 35% from 14%, operating expenses decreased 30% to \$1,490,930 from \$2,123,585. The increase in gross margin and decrease in operating expenses resulted in a 61% decrease in the loss to \$719,685 or \$0.02 per share compared to \$1,868,104 or \$0.07 per share in the same period last year.

For the six month period ended June 30, 2011, total revenues increased 38% to \$5,241,663 compared to \$3,804,992, gross margin on product sales increased to 24% from 13%, operating expenses decreased 34% to \$3,224,388 from \$4,871,200. Net loss decreased by 54% to \$2,101,891 or \$0.05 per share from \$4,528,152 or \$0.18 per share in the same period in the prior year.

As at June 30, 2011, the Company had a working capital balance of \$5,135,309 compared to \$6,611,283 as at December 31, 2010.

“We are pleased with the overall performance of the business in the second quarter,” said S. Wayne Kay, Chief Executive Officer. “Our continued focus on profitability and on managing working capital and overall costs has resulted in significantly improved results on a year-over-year and consecutive quarter basis. We are continuing to see the benefits of a reduced workforce as part of our company reorganization undertaken in 2010. In addition, our refocus on our cardiac business is reflected in increased sales volumes and dollars in China, Europe, and the Middle East.”

“Significant quarter to quarter fluctuation in revenue has been our history and we expect this to continue through the remaining quarters in 2011. The largest factor driving this fluctuation is the timing of the distribution of products to our major China market.”

For a further discussion of the Company’s financial results for the year ended December 31, 2010, please refer to the Company’s audited consolidated financial statements and related Management Discussion and Analysis, which can be found at [www.responsebio.com](http://www.responsebio.com), SEDAR (Canada) [www.sedar.com](http://www.sedar.com) or EDGAR (U.S.) [www.sec.gov/edgar/searchedgar/webusers.htm](http://www.sec.gov/edgar/searchedgar/webusers.htm). Information at these sites is typically available within 24 hours of the distribution of the news release.

## About Response Biomedical

Response Biomedical develops, manufactures and markets rapid on-site diagnostic tests for use with its RAMP<sup>®</sup> platform for clinical and environmental applications. RAMP<sup>®</sup> represents a new paradigm in diagnostics that provides high sensitivity and reliable information in minutes. It is ideally suited to both point of care testing and laboratory use.

*The RAMP<sup>®</sup> system consists of a Reader and single-use disposable test cartridges, and has the potential to be adapted to more than 250 medical and non-medical tests currently performed in laboratories. RAMP<sup>®</sup> clinical tests are commercially available for the early detection of heart attack and congestive heart failure through our commercial partners, Roche and Shionogi and through select international distributors.*

In the non-clinical market, RAMP<sup>®</sup> Tests are currently provided for the environmental detection of West Nile Virus, and Biodefense applications including the rapid on-site detection of anthrax, smallpox, ricin and botulinum toxin. Several other product applications are under development.

Response has achieved CE Marking for its Readers and clinical tests and its Quality Management System is registered to ISO 13485: 2003 and ISO 9001: ~~2000~~2008.

Response Biomedical is a publicly traded company listed on the TSX under the trading symbol "RBM" and quoted on the OTC Bulletin Board under the symbol "RPBFD". For further information, please visit the Company's website at [www.responsebio.com](http://www.responsebio.com).

*Statements contained in this press release relating to future results, events or developments, for example, statements regarding the private placement, future revenues, our expanding global network, growth possibilities and statements containing the words "believes," "may," "could", "plans," "will," "estimate," "continue," "anticipates," "intends," "expects", "goal" and similar expressions, are "forward-looking statements" or "forward-looking information" under applicable U.S. and Canadian securities laws. Forward-looking statements or information may involve, but are not limited to, comments with respect to our planned activities, business plan and strategies and their future implementation, and our expectations for our financial condition and the results of, or outlook for, our business operations generally. Forward-looking statements or information are subject to the related assumptions made by us and involve known and unknown risks, uncertainties and other factors that may cause actual results, events or developments to be materially different from those expressed or implied by such statements or information.*

*Many of such risks, uncertainties and other factors form part of our underlying assumptions, and include, among other things, financial risks that would affect our operations such as our limited available working capital and cash flows and whether and for how long available funds will be sufficient to fund our operations and our ability to raise additional capital as and when needed; our need for substantial additional funding to conduct research and development and commercialization activities; current financial market conditions which may negatively affect our ability to obtain financing; the ability to obtain regulatory approval and shareholder acceptance of planned financings, changing facility costs and other risks relating to our facilities expansion plans; our ability to establish, and our dependence upon, relationships with strategic alliance partners to develop and commercialize products; technological changes that impact our existing products or our ability to develop and commercialize our products; our ability to obtain and enforce timely patent and other intellectual property protection for our technology and products; our ability to obtain and maintain rights to technology from licensors; liability for patent, product liability and other claims asserted against us; commercialization limitations imposed by patents owned or controlled by third parties; technical risk in research and development; adverse results or unexpected delays in product development and clinical trials; our ability to retain, and our reliance upon, third party suppliers, manufacturers, distributors and alliance partners; our ability to attract and retain qualified personnel; our ability to effectively and efficiently manage the planned growth of our operations in Asia and in other international markets; our ability to obtain, and the timing of, necessary regulatory approvals; our ability to profitably sell our products at prices that would be acceptable to third-party reimbursement programs; competition including competition from others with significantly more resources; market acceptance of our products and the size of our markets; changes in business strategy or development plans; changes in, or the failure to comply with, governmental regulations; fluctuations in interest rates and foreign exchange rates; seasonality including government budget cycles; general economic and business conditions where we operate; and other factors referenced in our annual*

report on Form 20-F, our Annual Information Form (AIF) and other filings with Canadian and United States securities regulatory authorities.

Given these uncertainties, assumptions and risks, readers are cautioned not to place undue reliance on such forward-looking statements or information. We disclaim any obligation to update, or to publicly announce any revisions to, any such statements or information to reflect future results, events or developments, except as required by law.

SELECTED FINANCIAL DATA	Three Months Ended June 30		Six Months Ended June 30	
	2011	2010	2011	2010
	\$	\$	\$	\$
Product Sales	2,763,979	2,112,424	4,786,435	3,514,958
Cost of Sales	1,805,124	1,824,341	3,629,676	3,043,600
Gross Profit on Product Sales	958,855	288,083	1,156,759	471,358
Contract Service Revenue	7,125	143,089	455,228	290,034
Total Expenses	1,490,930	2,123,585	3,224,388	4,871,200
Loss for the Period	(719,685)	(1,868,104)	(2,101,891)	(4,528,152)
Loss per share	(0.02)	(0.07)	(0.05)	(0.18)

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