



**FOR IMMEDIATE RELEASE NASDAQ: CRME TSX: COM**

## **CARDIOME ENTERS COMMERCIALIZATION AGREEMENT WITH EUROLAB ESPECIALIDADES MEDICINALES FOR BRINAVESS™ IN ARGENTINA**

**Vancouver, Canada, Aug 25, 2014** -- Cardiome Pharma Corp. (NASDAQ: CRME / TSX: COM) today announced that its subsidiary, Correvio International Sàrl, has entered into an agreement with Eurolab Especialidades Medicinales de Eurofar S.R.L. to sell and distribute BRINAVESS™ (vernakalant intravenous) exclusively in Argentina. Under the terms of the agreement, Eurolab has agreed to specific annual commercial goals for BRINAVESS™. Financial details of the agreement have not been disclosed.

“This agreement with Eurolab allows for the continued availability of BRINAVESS to our customers in Argentina,” said Karim Lalji, Cardiome’s Chief Commercial Officer. “As the third largest country in South America in terms of population, Argentina remains a key market for BRINAVESS as we continue to build the brand in this region.”

“We are excited to expand our hospital product offerings to our customers with the addition of BRINAVESS to our portfolio,” said Gabriel Menendez, General Manager at Eurolab. “We are committed to making this important drug for the conversion of atrial fibrillation available to all of our customers throughout the country.”

In 2010, the Argentine pharmaceutical market was valued at \$5.1 billion with growth of 29.1% on a year-on-year basis.<sup>1</sup>

### **References:**

1. Scrip Insights. The pharmaceutical market in Argentina. Dec 2011.

### **About Cardiome Pharma Corp.**

Cardiome Pharma Corp. is a specialty pharmaceutical company dedicated to the development and commercialization of cardiovascular therapies that will improve the quality of life and health of patients suffering from heart disease. Cardiome has two marketed, in-hospital, cardiology products, BRINAVESS™ (vernakalant IV), approved in Europe and other territories for the rapid conversion of recent onset atrial fibrillation to sinus rhythm in adults, and AGGRASTAT® (tirofiban HCl) a reversible GP IIb/IIIa inhibitor indicated for use in patients with acute coronary syndrome.

Cardiome is traded on the NASDAQ Capital Market (CRME) and the Toronto Stock Exchange (COM). For more information, please visit our web site at [www.cardiome.com](http://www.cardiome.com).

### **About Eurolab Especialidades Medicinales**

Eurolab, headquartered in Buenos Aires, Argentina, supports the Multinational Pharmaceutical Industry in the areas of manufacturing, promotion and distribution of medicines in Argentina and Latin America. Eurolab also produces its own prescription and OTC products. For more information, please visit their web site at [www.laboratorioeurolab.com](http://www.laboratorioeurolab.com).

## **Forward-Looking Statement Disclaimer**

Certain statements in this news release contain forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995 or forward-looking information under applicable Canadian securities legislation that may not be based on historical fact, including without limitation statements containing the words “believe”, “may”, “plan”, “will”, “estimate”, “continue”, “anticipate”, “intend”, “expect” and similar expressions. Forward-looking statements may involve, but are not limited to, comments with respect to our objectives and priorities for the remainder of 2014 and beyond, our strategies or future actions, our targets, expectations for our financial condition and the results of, or outlook for, our operations, research and development and product and drug development. Such forward-looking statements involve known and unknown risks, uncertainties and other factors that may cause the actual results, events or developments to be materially different from any future results, events or developments expressed or implied by such forward-looking statements. Many such known risks, uncertainties and other factors are taken into account as part of our assumptions underlying these forward-looking statements and include, among others, the following: general economic and business conditions in the United States, Canada, Europe, and the other regions in which we operate; market demand; technological changes that could impact our existing products or our ability to develop and commercialize future products; competition; existing governmental legislation and regulations and changes in, or the failure to comply with, governmental legislation and regulations; availability of financial reimbursement coverage from governmental and third-party payers for products and related treatments; adverse results or unexpected delays in pre-clinical and clinical product development processes; adverse findings related to the safety and/or efficacy of our products or products; decisions, and the timing of decisions, made by health regulatory agencies regarding approval of our technology and products; the requirement for substantial funding to expand commercialization activities; and any other factors that may affect our performance. In addition, our business is subject to certain operating risks that may cause any results expressed or implied by the forward-looking statements in this presentation to differ materially from our actual results. These operating risks include: our ability to attract and retain qualified personnel; our ability to successfully complete pre-clinical and clinical development of our products; changes in our business strategy or development plans; intellectual property matters, including the unenforceability or loss of patent protection resulting from third-party challenges to our patents; market acceptance of our technology and products; our ability to successfully manufacture, market and sell our products; the availability of capital to finance our activities; and any other factors described in detail in our filings with the Securities and Exchange Commission available at [www.sec.gov](http://www.sec.gov) and the Canadian securities regulatory authorities at [www.sedar.com](http://www.sedar.com). Given these risks, uncertainties and factors, you are cautioned not to place undue reliance on such forward-looking statements and information, which are qualified in their entirety by this cautionary statement. All forward-looking statements and information made herein are based on our current expectations and we undertake no obligation to revise or update such forward-looking statements and information to reflect subsequent events or circumstances, except as required by law.

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