



## Insiders Purchasing Remaining Shares of Liquidating Fund and Clinical Update

**Sydney, Australia and San Diego, California (Friday, 8 July 2016 AEST)** – REVA Medical, Inc. (ASX: RVA) (“REVA” or the “Company”) announces that, following approval from the Company’s chairman, several members of its board of directors, along with institutional investors, are purchasing the remaining securities of a current REVA shareholder in a private transaction. The securities, traded in the form of CHESSE Depository Interests (“CDIs”) and representing an approximate 1.4% ownership position, are being acquired at A\$0.98 per CDI from the Visium Balanced Master Fund, an investment fund that is currently being liquidated. Following the private transaction, Visium has no remaining holdings in REVA.

REVA previously released positive data from its CE Mark clinical trial, FANTOM II Cohort A (see press release dated 18 May 2016). The Company reaffirms that it remains on track for final CE Mark submission during the third quarter of this year, with anticipated regulatory approval in the fourth quarter.

### About REVA

REVA is a clinical stage medical device company located in San Diego, California, USA, that is working to commercialize its proprietary bioresorbable scaffolds, as an alternative to metal stents, to treat coronary artery disease. Scaffolds provide restoration of blood flow, support the artery through the healing process, then disappear (or “resorb”) from the body over a period of time. This resorption allows the return of natural movement and function of the artery, a result not attainable with permanent metal stents. The Company’s *Fantom*<sup>®</sup> scaffold has been designed to offer an ideal balance of thinness and strength, with distinct ease-of-use features including complete scaffold visibility under x-ray, expansion with one continuous inflation, and no procedural time limitations. REVA will require regulatory approval before it can commercialize *Fantom* or any other product.

### Forward-Looking Statements

*This announcement contains or may contain forward-looking statements that are based on management's beliefs, assumptions and expectations and on information currently available to management. All statements that are not statements of historical fact, including those statements that address future operating performance and events or developments that we expect or anticipate will occur in the future, are forward-looking statements, such as those statements regarding our ability to obtain regulatory approvals, timely and successfully complete our clinical trials, protect our intellectual property position, commercialize our products if and when approved, develop and commercialize new products, recruit and retain our key personnel, and estimates regarding our capital requirements and financial performance. You should not place undue reliance on forward-looking statements. Although management believes forward-looking statements are reasonable as and when made, forward-looking statements are subject to a number of risks and uncertainties that may cause our actual results to vary materially from those expressed in forward-looking statements, including the risks and uncertainties that are described in the "Risk Factors" section of our Annual Report on Form 10-K filed with the US Securities and Exchange Commission (the "SEC") on March 10, 2016, and as may be updated in our periodic reports thereafter. Any forward-looking statements in this announcement speak only as of the date when made. REVA does not assume any obligation to publicly update or revise any forward-looking statements, whether as a result of new information, future events, or otherwise.*

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