CorMedix Successfully Completes and Passes ISO 13485 Certification Audit

The Associated Press

BRIDGEWATER, N.J.--(BUSINESS WIRE)--Oct 11, 2012--CorMedix Inc. (“CorMedix”) (NYSE Amex: CRMD), a pharmaceutical company focused on developing and commercializing therapeutic products for the prevention and treatment of cardiorenal disease, announced today receipt of ISO 13485:2003 certification. ISO 13485 is awarded by TÜV SÜD and is the globally recognized standard that outlines consistent international processes for the design and manufacturing of medical devices, including many supply chain functions such as assembly, packaging, warehousing and distribution. Compliance with ISO 13485 is often seen as a step towards achieving compliance with European regulatory requirements. CorMedix anticipates a final review and determination regarding its CE Mark application by the end of 4Q 2012.

“This certification represents a major achievement and milestone for CorMedix. The successful completion assists us in moving forward with regulatory registrations,” said Randy Milby, the Company’s Chief Operating Officer.

The ISO 13485:2003 certification is a stand-alone standard developed by the International Organization for Standardization that provides harmonized quality management systems requirements for manufacturers of medical devices. The conformity of Medical Devices and In-vitro Diagnostic Medical Device according to EEC-decrees 93/42/EEC, 90/385/EEC and 98/79/EEC must be assessed before sale is permitted. The preferred method to prove conformity is the certification of the Quality Management System according ISO 9001 and/or ISO 13485 and ISO 14971 by a Notified Body. The result of a positive assessment is the certificate of conformity allowing the CE Mark and the permission to sell the medical device in the European Union.

About CorMedix CorMedix Inc. is a pharmaceutical company that seeks to in-license, develop and commercialize therapeutic products for the prevention and treatment of cardiac and renal dysfunction, also known as cardiorenal disease. CorMedix most advanced product candidate: CRMD003 (Neutrolin ®) for the prevention of catheter related bloodstream infections and maintenance of catheter patency in tunneled, cuffed, central venous catheters used for vascular access in hemodialysis patients. Please see www.cormedix.com [1] for additional information.

Forward-Looking Statements This press release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995 that are subject to risks and uncertainties. All statements, other than statements of historical facts, regarding management's expectations, beliefs, goals, plans or CorMedix's prospects, future financial position, future revenues and projected costs should be considered forward-looking. Readers are cautioned that actual results

may differ materially from projections or estimates due to a variety of important factors, including the outcome of clinical trials of CorMedix's product candidates and whether they demonstrate these candidates' safety and effectiveness; the risks and uncertainties associated with: CorMedix’s ability to manage its limited cash resources, obtaining additional financing to support CorMedix's research and development and clinical activities and operations, including any subsequent closings of CorMedix’s private placement; obtaining regulatory approvals to conduct clinical trials and to commercialize CorMedix's product candidates, including CE Marking for Neutrolin ® in Europe; CorMedix's ability to enter into and maintain collaborations with third parties for its development programs; CorMedix's dependence on its collaborations and its license relationships; achieving milestones under CorMedix's collaborations; CorMedix's' dependence on preclinical and clinical investigators, preclinical and clinical research organizations, manufacturers and consultants; protecting the intellectual property developed by or licensed to CorMedix; and CorMedix's ability to maintain listing on NYSE-Amex. These and other risks are described in greater detail in CorMedix's filings with the Securities and Exchange Commission. CorMedix may not actually achieve the goals or plans described in its forward-looking statements, and investors should not place undue reliance on these statements. CorMedix disclaims any intent or obligation to update any forward-looking statements as a result of developments occurring after the date of this press release.

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Randy Milby, 908-517-9489 Chief Operating Officer KEYWORD: UNITED STATES NORTH AMERICA NEW JERSEY INDUSTRY KEYWORD: HEALTH BIOTECHNOLOGY CARDIOLOGY CLINICAL TRIALS HOSPITALS MEDICAL DEVICES PHARMACEUTICAL RESEARCH FDA MEDICAL SUPPLIES SCIENCE SOURCE: CorMedix Inc. Copyright Business Wire 2012 PUB: 10/11/2012 09:00 AM/DISC: 10/11/2012 09:00 AM http://www.businesswire.com/news/home/20121011005432/ [2]

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