



FOR: CARDIOCOMM SOLUTIONS, INC.
TSX VENTURE SYMBOL: EKG

January 29, 2019

CARDIOCOMM SOLUTIONS' HEARTCHECK™ DEVICE ENTERS FINAL FDA REVIEW PHASE

HeartCheck™ CardiBeat and GEMS™ Mobile review results expected in late February

TORONTO, ONTARIO - CardioComm Solutions, Inc. (TSX VENTURE:EKG) ("**CardioComm**" or the "**Company**"), a leading global provider of consumer heart monitoring and electrocardiogram ("**ECG**") acquisition and management software solutions, confirms it has completed a request for additional information from the US Food and Drug Administration ("**FDA**") for the Company's premarket notification 510(k), Class II medical device clearance application for the HeartCheck™ CardiBeat and GEMS™ Mobile Application.

The Company had submitted a letter of revocation of their supplementary information submission on December 26, 2018 in compliance with the FDA's directive. The Company has now provided the FDA a restatement of their response for additional information as of January 23, 2019, which the FDA has confirmed received. The FDA will now have 31 days to complete the 510(k) review of CardioComm's restated submission.

To learn more about CardioComm's products and for further updates regarding HeartCheck™ ECG device integrations please visit the Company's websites at www.cardiocommsolutions.com and www.theheartcheck.com.

About CardioComm Solutions

CardioComm Solutions' patented and proprietary technology is used in products for recording, viewing, analyzing and storing electrocardiograms for diagnosis and management of cardiac patients. Products are sold worldwide through a combination of an external distribution network and a North American-based sales team. CardioComm Solutions has earned the ISO 13485:2016 certification, is HIPAA compliant and holds clearances from the European Union (CE Mark), the USA (FDA) and Canada (Health Canada).

FOR FURTHER INFORMATION PLEASE CONTACT:

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Forward-looking statements

This release may contain certain forward-looking statements and forward-looking information with respect to the financial condition, results of operations and business of CardioComm Solutions and certain of the plans and objectives of CardioComm Solutions with respect to these items. Such statements and information reflect management's current beliefs and are based on information currently available to management. By their nature, forward-looking statements and forward-looking information involve risk and uncertainty because they relate to events and depend on circumstances that will occur in the future and there are many factors that could cause actual results and developments to differ materially from those expressed or implied by these forward-looking statements and forward-looking information.

In evaluating these statements, readers should not place undue reliance on forward-looking statements and forward-looking information. The Company does not assume any obligation to update the forward-looking statements and forward-looking information contained in this release other than as required by applicable laws, including without limitation, Section 5.8(2) of National Instrument 51-102 (*Continuous Disclosure Obligations*).

Neither TSX Venture Exchange nor its Regulation Services Provider (as that term is defined in policies of the TSX Venture Exchange) accepts responsibility for the adequacy or accuracy of this release.