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Clear Data. Trusted Results.

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PRESS RELEASE

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Duke Clinical Research Institute's Trial of Recom's Heart Monitor is Underway

LOS ANGELES--(BUSINESS WIRE)--Jan. 6, 2005--Recom Managed Systems, Inc. (OTCBB: RECM - News) in conjunction with the Duke Clinical Research Institute, of Durham, NC, announced today that its Sponsored Research trial is underway. The purpose of the trial is to conduct a series of independent market preference tests that will establish the Model 100's performance against currently marketed ECG monitoring devices.

The first patient recording in this evaluation was completed on 12/15/2004 by the eECG Core Laboratory at the Duke Clinical Research Institute. Commenting on the Study, Marvin H. Fink, Recom's CEO stated, "While it is just the beginning of this study, I am confident that our Model 100's performance will meet or exceed that of other ambulatory ECG monitoring devices currently available in the marketplace."

The Recom ECG monitor system utilizes the company's patented signal technology that produces a high fidelity ECG signal in the presence of ambient noise. This research is intended to demonstrate that a Recom 12-lead ambulatory recording can produce ECG data on a par with other 12-lead modalities such as resting ECG and exercise stress testing.

About Recom Managed Systems, Inc.

Recom Managed Systems, Inc. is an emerging life sciences company focused on the monitoring and detection of disease through continuous biomedical signal monitoring. Recom Managed Systems, Inc. uses its patented signal technology to design and develop medical devices that simplify and reduce the costs of diagnostic testing and patient monitoring in an ambulatory setting. With our patented signal technology platform, Recom brings clinical quality physiological signal monitoring to the ambulatory setting.

Caution Regarding Forward-Looking Statements

Statements in this release that are not strictly historical are "forward-looking" statements. Forward-looking statements involve known and unknown risks, which may cause Recom's actual results in the future to differ materially from expected results. Factors which could cause or contribute to such differences include, but are not limited to failure to complete the development and introduction of new products or services, failure to obtain federal or state regulatory approvals governing medical devices, monitoring and other related services or products, inability to obtain physician, patient or insurance acceptance of Recom's products or services, adverse equity market conditions and declines in the value of Recom's common stock, and the unavailability of financing to complete management's plans and objectives. These risks are qualified in their entirety by cautionary language and risk factors set forth and to be further described in Recom's filings with the Securities and Exchange Commission.

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