Latest Test Results Bolster RenovaCare Strategy to Spray Stem Cells on Wounds and Burns for Rapid Healing

New York & Pittsburgh – May 31, 2016 - RenovaCare, Inc., (OTCQB: RCAR) today announced positive spray test results demonstrating that its novel SkinGun™ technology achieves exceptionally-uniform distribution of fluids with 200 times greater coverage than conventional methods. These outcomes are especially promising to scientists developing patented RenovaCare CellMist™ and SkinGun™ technologies* for spraying fluids containing a patient’s own stem cells onto burns and wounds for rapid scar-free healing.

“These results mark important technical milestones in our pathway to spraying patient wounds with their own stem cells to promote fast and natural skin repair,” said Thomas Bold, President and CEO of RenovaCare, Inc. “Data from ongoing preclinical work supports our long-held conviction that our SkinGun™ technology is superior to standard methods for delivering fluids and stem cells to target sites, achieving excellent coverage while being extremely gentle to the cells.”

Click here to see video of ultra-gentle RenovaCare SkinGun™ sprayer.

In addition to these tests showing high distribution and concentration patterns, the SkinGun™ recently demonstrated the ability to spray powerful yet delicate skin stem cells ultra-gently. Scientists reported impressive 97.3% cell viability after SkinGun™ spraying. Cell viability is essential to regenerating skin for burns, wounds, and cosmetic applications.

Specifically, scientists at one of the world’s largest university hospitals, Berlin-Brandenburg Center for Regenerative Therapies (BCRT), a translational research center at Charité – Universitätsmedizin Berlin, sprayed human skin stem cells using the RenovaCare™ SkinGun™. Among technical aims of the study, was evaluation of several factors important to the regeneration of human skin, including cell yield, viability, metabolic activity, and cell growth.

Positive results were reported from experiments related to each of these investigations, and cell growth was comparable to pipetting, the industry’s widely accepted ‘gold-standard’ for the deposition of cells.
In these latest tests, scientists repeatedly sprayed fluids at various airflow rates using the RenovaCare SkinGun™, resulting in 200-times more droplets than conventional ‘syringe deposition’. For example, in an 8cm diameter surface area the SkinGun™ delivered more than 20,000 evenly distributed droplets versus only 91 droplets by conventional needle and syringe methods.

Scientists evaluated droplet size, distribution, and density alongside spray velocity and fluid viscosity in experiments conducted at Stem cell Systems GmbH (Berlin, Germany). Additional testing of the SkinGun™ remains ongoing.

The RenovaCare medical-grade spray device gently sprays a well-dispersed mist of wound care and/or irrigation fluids using a patented liquid-into-air stream system. The device targets the estimated $45 billion plus wound care market in the U.S.

*RenovaCare products are currently in development. They are not available for sale in the United States. There is no assurance that the company’s planned or filed submissions to the U.S. Food and Drug Administration, if any, will be accepted or cleared by the FDA.

About RenovaCare
RenovaCare, Inc. is developing first-of-their-kind autologous (self-donated) stem cell therapies for the regeneration of human organs, and novel medical grade liquid sprayer devices.

The company’s pipeline products under development target the body’s largest organ, the skin. The RenovaCare CellMist™ System will use the patented SkinGun™ to spray a liquid suspension of a patient’s stem cells – the CellMist™ Solution – onto wounds. RenovaCare is developing its CellMist™ System as a promising new alternative for patients suffering from burns, chronic and acute wounds, and scars. In the U.S. alone, this $45 billion market is greater than the spending on high-blood pressure management, cholesterol treatments, and back pain therapeutics.

In addition to the SkinGun™, the company is developing a medical-grade sprayer that uses a liquid-into-air stream system to disperse a gentle spray of wound care and/or irrigation fluids over a wound area.

For additional information, please call Drew Danielson at: 888-398-0202 or visit: http://renovacareinc.com

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increases for supplies and components, litigation and administrative proceedings involving the 
Company, the possible acquisition of new businesses or technologies that result in operating losses or 
that do not perform as anticipated, unanticipated losses, the possible fluctuation and volatility of the 
Company's operating results, financial condition and stock price, losses incurred in litigating and settling 
cases, dilution in the Company's ownership of its business, adverse publicity and news coverage, 
inability to carry out research, development and commercialization plans, loss or retirement of key 
executives and research scientists, and other risks. There can be no assurance that further research and 
development will validate and support the results of our preliminary research and studies. Further, 
there can be no assurance that the necessary regulatory approvals will be obtained or that the Company 
will be able to develop commercially viable products on the basis of its technologies. In addition, other 
factors that could cause actual results to differ materially are discussed in the Company's most recent 
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Exchange Commission at 100 F Street, N.E., Washington, D.C. 20549. You can obtain information about 
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The outcomes described herein are from preclinical studies and single case observations from work 
conducted under Innovative Practice Approach (IPA) guidelines with Institutional Review Board 
approval. The IPA work was undertaken by parties other than RenovaCare, Inc. or its affiliates at that 
time. This treatment was provided with a prototype version of the device, subsequently acquired by 
RenovaCare, Inc. The company has announced plans to conduct FDA-regulated clinical studies in the 
future. There is no assurance that such studies will be successfully undertaken, and even if or when 
initiated, that such studies will be completed or yield clinical outcomes.

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