Revive Therapeutics and South Carolina Research Foundation Enter into Exclusive License Agreement for Cannabinoids-Based Therapeutics in the Treatment of Liver Diseases

TORONTO, ONTARIO--(Marketwired – September 19, 2017) – Revive Therapeutics Ltd. ("Revive" or the "Company") (TSX VENTURE: RVV) (OTCQB: RVVTF), a company focused on the research, development and commercialization of novel treatments for serious and unmet medical needs, today announced that it has entered into a license agreement with South Carolina Research Foundation ("SCRF"), under which Revive will acquire an exclusive license from SCRF to develop and commercialize a portfolio of patents based on cannabinoid-based therapeutics, such as Cannabidiol, in the treatment of liver diseases.

"The licensing of the patent portfolio of cannabinoids in the treatment of liver diseases is an important milestone for Revive, as we continue to realize our objective of commercializing treatments to address unmet medical needs in liver diseases," said Craig Leon, Chief Executive Officer of Revive. "We will continue focusing on commercializing our candidates of cannabinoid-based therapies and on expanding our product pipeline in the field of liver disease such as autoimmune hepatitis, liver cirrhosis, non-alcoholic steatohepatitis (NASH), and non-alcoholic fatty liver disease (NAFLD)."

Revive will pay SCRF a nominal one time fee for entering into the license. All other future payments are consistent with industry standard and will be based on clinical trial and revenue milestones reached by Revive in the future.

About Liver Disease

Liver disease is described by irregular functioning of liver, causing disorders like hepatitis, fatty liver (NASH), and cirrhosis. There are over 100 described diseases of the liver affecting at least 30 million people alone in the U.S. A number of factors are driving the liver disease treatment market, which includes rapidly changing lifestyle patterns such as increasing alcohol consumption, unhealthy diets, and increasing prevalence of liver diseases. Liver diseases can result from injury to the liver caused by hepatitis C virus (HCV), hepatitis B virus (HBV), obesity, chronic excessive alcohol use or autoimmune diseases. Major drug categories used in the treatment of liver diseases includes anti-rejection drugs, vaccines, immunosuppressant, chemotherapy drugs and antiviral drugs. According to Allied Market Research, titled, "World Liver Disease Treatment Market – Opportunities and Forecast, 2014 - 2022", the global market for liver disease treatment is projected to reach \$19,536 million by 2022.

About South Carolina Research Foundation

Formed in 1997, the South Carolina Research Foundation (SCRF) is a major initiative to increase the support for faculty doing research at USC. The SCRF provides greater flexibility and competitiveness in implementing research grants and contracts, building infrastructure, and developing intellectual property for the faculty. Management

responsibilities of the SCRF include USC's intellectual technology licensing program and the USC Columbia Technology Incubator.

About Revive Therapeutics Ltd.

Revive Therapeutics Ltd. (TSX VENTURE:RVV) (OTCQB:RVVTF) is focused on the research, development and commercialization of novel treatments for serious and unmet medical needs by identifying and investigating potential therapies targeting the endocannabinoid system, such as cannabinoids, that may be repurposed for new indications, be delivered in a different way, combined with existing drugs, or be developed as new chemical entities. Additional information on Revive is available at www.ReviveThera.com.

For more information please contact:

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CAUTIONARY NOTE REGARDING FORWARD-LOOKING STATEMENTS

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

This news release includes certain information and statements about management's view of future events, expectations, plans and prospects that constitute "forward-looking information" that involves known and unknown risks and uncertainties, which are not comprised of historical facts, and most of which are beyond the control of Revive. Forward-looking statements include estimates and statements that describe Revive's future plans, objectives or goals, including words to the effect that Revive or its management expects a stated condition or result to occur. Forward-looking statements may be identified by such terms as "believes", "anticipates", "intends", "expects", "estimates", "may", "could", "would", "will", or "plan", and similar expressions. Specifically, forward-looking statements in this news release include, without limitation, statements regarding: the granting of a patent for Bucillamine for the treatment of gout; the potential efficacy and commercial viability of Bucillamine for treatment of gout and Bucillamine for the treatment of Cystinuria; expansion of the Bucillamine clinical testing program; the Company's drug research and development, and commercialization plans; the Company's research, development and commercialization plans for plant-based therapies, including cannabinoids; the timing of operations; and estimates of market conditions. These statements involve known and unknown risks, uncertainties, and other factors that may cause actual results or events, performance, or achievements of Revive to differ materially from those anticipated or implied in such forward-looking statements. Since forward-looking statements are based on assumptions and address future events and conditions, by their very nature they involve inherent risks and uncertainties. The Company believes that the expectations reflected in these forward-looking statements are reasonable, but there can be no assurance that actual results will meet management's expectations. In formulating the forward-looking statements contained herein, management has assumed: that business and economic conditions affecting Revive will continue substantially in the ordinary course and will be favourable to Revive; that clinical testing results will justify commercialization of the Company's drug candidates; that Revive will be able to obtain all requisite regulatory approvals to commercialize its drug candidates; that such approvals will be received on a timely basis; and, that Revive will be able to find suitable partners for development and commercialization of its drug repurposing candidates on favourable terms. Although these assumptions were considered reasonable by management at the time of preparation, they may prove to be incorrect and no assurance can be given that such events will occur in the disclosed time frames or at all.

Factors that may cause actual results to differ materially from those anticipated by these forward-looking statements include: uncertainties associated with obtaining regulatory approval to perform clinical trials and market products; the need to establish additional corporate collaborations, distribution or licensing arrangements; the Company's ability to raise additional capital if and when necessary; intellectual property disputes; increased competition from pharmaceutical and biotechnology companies; changes in equity markets, inflation, and changes in exchange rates; and other factors as described in detail in Revive's Management's Discussion & Analysis for the period ended June 30, 2016 and Revive's other public filings, all of which may be viewed on SEDAR (www.sedar.com). Given these risks and uncertainties, readers are cautioned not to place undue reliance on such forward-looking statements and information, which are qualified in their entirety by this cautionary statement. Except as required by law, Revive disclaims any intention and assumes no obligation to update or revise any forward-looking statements to reflect actual results, whether as a result of new information, future events, changes in assumptions, changes in factors affecting such forward-looking statements or otherwise.