NERVGEN PHARMA PRESENTING AT THE 9TH ANNUAL LD MICRO INVITATIONAL

Vancouver, Canada. June 4, 2019 – NervGen Pharma Corp. (TSX-V: NGEN) (OTCQB: NGENF) (“NervGen” or the “Company”), a regenerative medicine company dedicated to creating innovative solutions for the treatment of nerve damage, including spinal cord and peripheral nerve injury, today announced that Ernest Wong, NervGen Pharma’s President & CEO, will be presenting at the 9th Annual LD Micro Invitational on Wednesday, June 5 at 8:40AM PDT / 11:40AM EDT. In addition, NervGen representatives will be available for investor meetings during the event.

The 9th Annual LD Micro Invitational, which will feature over 200 companies and attended by over 1,000 individuals, is taking place on June 4th and 5th at the Luxe Sunset Bel Air Hotel in Los Angeles.

About NervGen
NervGen is restoring life’s potential by creating innovative solutions for the treatment of nerve damage, including spinal cord injuries and peripheral nerve injuries. The Company also continues to research secondary applications such as multiple sclerosis, stroke, acute myocardial infarction induced arrhythmia (“AMI”, commonly known as a heart attack) and other neurodegenerative diseases.

NervGen plans to initiate a Phase 1 human clinical trial for its lead compound, NVG-291, in early 2020 under an Investigational New Drug application with the US Food and Drug Administration. NervGen is advancing NVG-291 for the treatment of spinal cord injury as the Company believes this indication is a significant opportunity due to the current lack of non-surgical solutions in the market, the dramatic impact on quality of life and the high cost burden to the healthcare system. The Company believes NVG-291 as a therapy could alleviate or improve upon the symptoms and conditions associated with spinal cord injury and empower these patients to live more active and productive lives.

For further information, please contact: Bill Radvak, Executive Chairman bradvak@nervgen.com

Follow NervGen on Twitter (@NervgenC) and LinkedIn (NervGen Pharma Inc.) for the latest news on the Company.

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

Cautionary Note Regarding Forward-Looking Statements
This news release may contain “forward-looking information” and “forward-looking statements” within the meaning of applicable Canadian and United States securities legislation. Such forward-looking statements and information herein include, but are not limited to, the Company’s current and future plans, expectations and intentions, results, levels of activity, performance, goals or achievements, or any other future events or developments constitute forward-looking statements, including, without limitation, statements regarding
advancement of NVG-291 toward clinical development and commercialization, the timing of human trials and regulatory approval, the potential efficacy of the Company’s products and technology, and the potential to identify, evaluate and develop other drug candidates. The words “may”, “will”, “would”, “should”, “could”, “expect”, “plan”, “intend”, “trend”, “indication”, “anticipate”, “believe”, “estimate”, “predict”, “likely” or “potential”, or the negative or other variations of these words or other comparable words or phrases, are intended to identify forward-looking statements.

Forward-looking statements are based on estimates and assumptions made by the Company in light of management’s experience and perception of historical trends, current conditions and expected future developments, as well as other factors that the Company believes are appropriate and reasonable in the circumstances. Many factors could cause the Company’s actual results, level of activity, performance or achievements or future events or developments to differ materially from those expressed or implied by the forward-looking statements, including without limitation, a lack of revenue, insufficient funding, reliance upon key personnel, the uncertainty of the clinical development process, competition, and other factors set forth in the “Risk Factors” section of the Company’s Prospectus, financial statements and Management Discussion and Analysis which can be found on SEDAR.com.

Readers should not place undue reliance on forward-looking statements made in this document. Furthermore, unless otherwise stated, the forward-looking statements contained in this document are made as of the date of this document, and the Company has no intention and undertakes no obligation to update or revise any forward-looking statements, whether as a result of new information, future events or otherwise, except as required by applicable law. The forward-looking statements contained in this document are expressly qualified by this cautionary statement.