FOR IMMEDIATE RELEASE

SUSTAINED NANO SYSTEMS TO PRESENT AT THE AMERICAN ACADEMY OF OPHTHALMOLOGY ANNUAL CONVENTION

New York, NY – July 17, 2018 (NEWSFRONT MEDIA). Sustained Nano Systems, LLC (“SNS”), a biopharmaceutical company utilizing its platform technology of bio-resorbable nano and microparticles for long term drug delivery, is pleased to announce that its abstract, "Long-Term Sustained Delivery Of Bevacizumab Using A Novel Delivery Platform", has been selected for presentation at the American Academy of Ophthalmology convention to be held Oct 27 – 30, 2018, in Chicago. The American Academy of Ophthalmology is the leading professional medical association of ophthalmologists, with a membership of 32,000 physicians including more than 90 percent of practicing ophthalmologists in the United States as well as over 7,000 members abroad.
"This selection indicates the significant interest in SNS research and its ability to create long term delivery of large proteins like monoclonal antibodies," said Dr. Barry Libin, CEO of SNS. Monoclonal antibodies are used in retinal disease (macular degeneration) as well as in the new immunotherapies in cancer research and other disease states.

ABOUT SUSTAINED NANO SYSTEMS (SNS)

SNS is a privately held biopharmaceutical company focused on the development and commercialization of innovative therapies for diseases and conditions in ophthalmology, oncology and other areas. The Company’s research facility is located on the campus of Stony Brook University as part of the Long Island High Technology Incubator and Center for Biotechnology. The SNS platform is the result of a comprehensive research effort that allows long term micro-dosing for continuous release of drugs to targeted tissues. The basis of the SNS pipeline is its patented technology that incorporates bio-
resorbable nano and micro carriers to deliver sustained delivery of important therapeutics.

SNS is preparing for clinical trials for its proprietary drug: DEX-SA, a single, peri-ocular administration of dexamethasone to reduce inflammation of the eye following cataract surgery. This drug is to replace the need for daily and multiple eye drops to prevent ocular inflammation.

SNS has also reported positive results for the in-vitro presence of Latanaprost for the treatment of glaucoma over a duration of 159 days. Latanaprost, a prostaglandin analogue, is the #1 glaucoma drug in the world, requiring dosing of up to four drops every day for the remainder of a patient’s life. At the present time, there is no long-acting formulation for glaucoma. SNS LAT-LA (Latanaprost Long Acting) is designed to be administered by a single periocular injection with a duration of six months. “The potential for SNS LAT-LA to sustain delivery for six months, would fulfill an important unmet need in replacing the more burdensome and potentially less compliant regimen of daily eye drops,” said Dr. Libin. LAT-LA will now undergo further studies to indicate its efficacy in the long term reduction of intra-ocular pressure. Glaucoma is the second leading cause of blindness in the world, in which abnormally high intraocular pressure may cause damage to the optic nerve, resulting in irreversible vision loss. In the U.S. alone, 2.7 million people suffer from glaucoma. According to IMS Health data, there were 34 million prescriptions and sales of over $2.7 billion of drugs administered by eye drops for the treatment of glaucoma in the U.S. in 2017, and global spending was over $6 billion. Compliance is the biggest problem with existing therapies for glaucoma, with more than 50% of patients on topical prostaglandin analogs not compliant within the first six months of treatment

For further information, visit: www.sustainednanosystems.net

Forward-Looking Statements
SNS Disclosure Notice: This press release contains “forward-looking statements” as that term is defined in the Private Securities Litigation Reform Act of 1995. For this purpose, any statements contained herein or which are otherwise made by or on behalf of the Company that are not statements of historical facts may be deemed forward looking statements. Without limiting the generality of the foregoing, words such as “may,” “will,” “to,” “plan,” “expect,” “believe,” “anticipate,” “intend,” “could,” “should,” “would,” “estimate,” or “continue,” or the negative or other variations thereof or comparable terminology are intended to identify forward looking statements. Investors are cautioned that all forward looking statements involve risk and uncertainties which may cause results to differ materially from those set forth in the statements. Such risks and uncertainties include, but are not limited to the following: the success of research and development activities and the speed with which regulatory authorizations and product launches may be achieved; government regulation generally; competitive developments; the ability to successfully market products domestically and internationally; difficulties or delays in manufacturing or issues relating to manufacturing capacity; commercial obstacles to the successful introduction of brand products generally; legal defense costs, insurance expenses, settlement costs, and the risk of an adverse decision or settlement relating to product liability, patent protection,
governmental investigations, and other legal proceedings; the Company’s ability to acquire and protect patents and other intellectual property both domestically and internationally; the absence of certainty regarding the receipt of required regulatory approval or the timing or terms of such approvals; any changes in business, political and economic conditions; business interruption due to hurricanes or other events outside of the Company’s control.

Investors are cautioned not to place reliance on these forward-looking statements, which are valid only as of the date they were made. The Company undertakes no obligation to update or revise any forward-looking statements to reflect new information or the occurrence of unanticipated events or otherwise, except as expressly required by law.

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