FOR IMMEDIATE RELEASE

Company Contacts:
Nuvectra Corporation
Walter Berger, Chief Financial Officer
(972) 668-4106
wberger@nuvectramed.com

Jennifer Armstrong, Media Relations
(214) 618-4823
jarmstrong@nuvectramed.com

Investor Contacts:
The Ruth Group
Brian Johnston
(646) 536-7028
bjohnston@theruthgroup.com

Zack Zubow
(646) 536-7020
zkubow@theruthgroup.com

Nuvectra Files Regulatory Submission with FDA for Algovita® MRI-Conditional Approval

Plano, Texas, June 21, 2017 – Nuvectra Corporation (NASDAQ:NVTR), a neurostimulation medical device company, announced today that it has filed its regulatory submission with the U.S. Food and Drug Administration (FDA) for full-body MRI-conditional approval for the Company's Algovita SCS system.

Scott Drees, Chief Executive Officer, commented, "This submission to the FDA represents our continued commitment to the advancement of the Algovita system. The addition of a full body MRI-compatibility label will further enhance the product features making the Algovita system more attractive to potential customers and improving our competitive positioning. We look forward to working collaboratively with FDA through the review period, and upon approval, will be excited to offer new MRI capabilities for patients that choose the Algovita SCS system."

This submission, pending regulatory approval from the FDA, would position the Company to achieve MRI-conditional approval at or around year end 2017 following a 180 day review process.

About Nuvectra Corporation

Nuvectra™ is a neurostimulation company committed to helping physicians improve the lives of people with chronic neurological conditions. The Algovita® Spinal Cord Stimulation (SCS) System is our first commercial offering and is CE marked and FDA approved for the treatment of chronic pain of the trunk and/or limbs. Our innovative technology platform also has capabilities under development to support other neurological indications such as sacral nerve stimulation (SNS), and deep brain stimulation (DBS). In addition, our NeuroNexus subsidiary designs, manufactures and markets leading-edge neural-interface technologies for the neuroscience clinical research market. Visit the Nuvectra website at www.nuvectramed.com.

Cautionary Note Regarding Forward-Looking Statements

This press release contains "forward-looking statements," including statements we make regarding the outlook for Nuvectra as an independent publicly-traded company. Forward-looking statements are based only on our current beliefs, expectations and assumptions regarding the future of our business, future plans and strategies, projections, anticipated events and trends, the economy and other future conditions, and therefore they are subject to inherent

uncertainties, risks and changes in circumstances that are difficult to predict and may be outside of our control. Our actual performance may differ materially from those indicated in the forward-looking statements. Therefore, you should not rely on any of these forward-looking statements. Any forward-looking statement made by us is based only on information currently available to us and speaks only as of the date on which it is made. Important factors that could cause our actual results to differ materially from those indicated in the forward-looking statements include: (i) the timing of the commercial launch of Algovita in the United States; (ii) our ability to successfully commercialize Algovita and develop and commercialize enhancements to Algovita; (iii) the uncertainty of obtaining regulatory approvals in the United States and Europe for our Virtis SNS system, (iv) our ability to successfully launch and commercialize the Virtis SNS system if it receives regulatory approval, (v) the outcome of our development plans for our neurostimulation technology platform, including our ability to identify additional indications or conditions for which we may develop neurostimulation medical devices or therapies and seek regulatory approval thereof; (vi) our ability to identify business development and growth opportunities and to successfully execute on our strategy, including our ability to seek and develop strategic partnerships with third parties to, among other things, fund clinical and development costs for new product offerings; (vii) the performance by our development partners, including Aleva Neurotherapeutics, S.A., of their obligations under their agreements with us; (viii) the scope of protection for our intellectual property rights covering Algovita and other products using our neurostimulation technology platform, along with any product enhancements; (ix) our ability to successfully build an effective commercial infrastructure and sales force in the United States; (x) our compliance with all regulatory and legal requirements regarding implantable medical devices and interactions with healthcare professionals; and (xi) any product recalls or the receipt of any warning letters from any governmental or regulatory agency. Please see the sections entitled "Cautionary Statement Concerning Forward-Looking Statements" and "Risk Factors" in Nuvectra's Registration Statement on Form 10 for a description of these and other risks and uncertainties. We undertake no obligation to publicly update any forward-looking statement, whether written or oral, that may be made from time to time, whether as a result of new information, future developments or otherwise.