argenx to Present Additional Data from Global Phase 3 ADAPT Trial of Efgartigimod for Myasthenia Gravis at Upcoming Virtual Medical Meetings

Breda, the Netherlands / Ghent, Belgium – argenx (Euronext & Nasdaq: ARGX), a global immunology company committed to improving the lives of people suffering from severe autoimmune diseases and cancer, today announced that it will present new data from the pivotal Phase 3 ADAPT trial at the Myasthenia Gravis Foundation of America (MGFA) 2020 Virtual Scientific Session. These data will be presented again at the American Association of Neuromuscular & Electrodiagnostic Medicine (AANEM) 2020 Virtual Annual Meeting.

The ADAPT trial evaluated efgartigimod, an FcRn antagonist, in patients with generalized myasthenia gravis (gMG). argenx previously announced positive topline efficacy and safety data from the ADAPT trial in May 2020, which will be submitted as part of a Biologics License Application (BLA) to the U.S. Food and Drug Administration (FDA) by the end of 2020.

Details for the presentations are as follows:

MGFA 2020 Virtual Scientific Session
Date and Time: Saturday, October 3, 2020 at 3:20pm Eastern Time
Session: MG Clinical Trials
Title: *Treatment of Patients with Myasthenia Gravis with Efgartigimod: Results of the Phase 3 ADAPT Study*
Presenter: James F. Howard Jr., M.D., Professor of Neurology (Neuromuscular Disease), Medicine and Allied Health, Department of Neurology, The University of North Carolina at Chapel Hill School of Medicine

AANEM 2020 Virtual Annual Meeting
Date and Time: Wednesday October 7, 2020 at 3:15pm Central Time
Session: Abstract Poster Session I, Abstract #142
Title: *ADAPT: A Phase 3 Study of FcRn Antagonist, Efgartigimod, in Myasthenia Gravis*
Presenter: James F. Howard Jr., M.D., Professor of Neurology (Neuromuscular Disease), Medicine and Allied Health, Department of Neurology, The University of North Carolina at Chapel Hill School of Medicine

Note that the date and time of these virtual presentations are subject to change by meeting organizers. For the most up-to-date information, please visit the Company's website at www.argenx.com/investors.

Phase 3 ADAPT Trial
The Phase 3 ADAPT trial was a randomized, double-blind, placebo-controlled, multi-center, global trial evaluating the safety and efficacy of efgartigimod in patients with gMG. A total of 167 adult patients with gMG in North America, Europe and Japan enrolled in the trial and were treated. Patients were eligible to enroll in ADAPT regardless of antibody status, including patients with AChR antibodies (AChR-Ab+) and patients where AChR antibodies were not detected. Patients were randomized in a 1:1 ratio to receive efgartigimod or placebo for a total of 26 weeks. ADAPT was designed to enable an individualized treatment approach with an initial treatment cycle followed by a variable number of subsequent treatment cycles. The primary endpoint was the number of AChR-Ab+ patients who achieved a response on the MG-ADL score defined by at least a two-point improvement for four or more consecutive weeks.

About argenx
argenx is a global immunology company committed to improving the lives of people suffering from severe autoimmune diseases and cancer. Partnering with leading academic researchers through its Immunology Innovation Program (IIP), argenx is translating immunology breakthroughs into a world-class portfolio of novel antibody-based medicines. argenx is evaluating efgartigimod in multiple serious autoimmune diseases, and cusatuzumab in hematological cancers in collaboration with Janssen. argenx is also advancing several earlier stage experimental medicines within its therapeutic franchises. argenx has offices in Belgium, the United States and Japan. For more information, visit www.argenx.com and follow us on LinkedIn at https://www.linkedin.com/company/argenx/.

Forward-looking Statements

The contents of this announcement include statements that are, or may be deemed to be, "forward-looking statements." These forward-looking statements can be identified by the use of forward-looking terminology, including the terms "believes," "estimates," "anticipates," "expects," "intends," "may," "will" or "should" and include statements argenx makes concerning the timing of its BLA submission to the FDA and the therapeutic potential of its product candidates. By their nature, forward-looking statements involve risks and uncertainties and readers are cautioned that any such forward-looking statements are not guarantees of future performance. argenx's actual results may differ materially from those predicted by the forward-looking statements as a result of various important factors, including argenx's expectations regarding its the inherent uncertainties associated with competitive developments, preclinical and clinical trial and product development activities and regulatory approval requirements; argenx's reliance on collaborations with third parties; estimating the commercial potential of argenx's product candidates; argenx's ability to obtain and maintain protection of intellectual property for its technologies and drugs; argenx's limited operating history; and argenx's ability to obtain additional funding for operations and to complete the development and commercialization of its product candidates. A further list and description of these risks, uncertainties and other risks can be found in argenx's U.S. Securities and Exchange Commission (SEC) filings and reports, including in argenx's most recent annual report on Form 20-F filed with the SEC as well as subsequent filings and reports filed by argenx with the SEC. Given these uncertainties, the reader is advised not to place any undue reliance on such forward-looking statements. These forward-looking statements speak only as of the date of publication of this document. argenx undertakes no obligation to publicly update or revise the information in this press release, including any forward-looking statements, except as may be required by law.

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