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ASIA EDITION

Argenx Breaks New Ground in Autoimmune Therapy with 2024 Milestones

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FDA approvals, innovative treatments, and strategic global expansion propel Argenx to the forefront of autoimmune disease management.

Argenx, a global leader in immunology, has made significant strides in 2024, bolstering its position as a pioneer in the treatment of severe autoimmune diseases. The company's recent achievements span critical clinical developments, regulatory approvals, and strategic financial performance, underscoring its dedication to delivering innovative therapies to patients worldwide.

One of the most notable milestones for Argenx this year was the FDA approval of VYVGART Hytrulo, a groundbreaking neonatal Fc receptor (FcRn) blocker for the treatment of chronic inflammatory demyelinating polyneuropathy (CIDP). This approval marks the first new therapeutic option for CIDP in over three decades, providing hope for patients who have long faced limited treatment options. The approval was supported by compelling data from the ADHERE study, which demonstrated that VYVGART Hytrulo significantly reduces the risk of relapse and enhances patient functionality.

Argenx further showcased its commitment to advancing autoimmune therapies with the presentation of promising data for empasiprubart at the 2024 Peripheral Nerve Society Annual Meeting. The treatment for multifocal motor neuropathy (MMN) demonstrated significant reductions in the need for IVIg retreatment and improvements in muscle strength, signaling a potential new standard of care for MMN patients.

Financially, Argenx has also reported robust results for the first quarter of 2024, with global net product sales reaching $398 million, primarily driven by the strong performance of VYVGART SC. The company remains in a powerful financial position with $3.1 billion in cash reserves, positioning it well to continue advancing its pipeline. Key pipeline developments include the progression of empasiprubart and efgartigimod for Sjogren’s disease to Phase 3 trials. However, in a strategic decision based on risk assessments, Argenx has chosen not to pursue efgartigimod for ANCA-associated vasculitis (AAV).

Argenx's global expansion efforts were also highlighted by the approval of VYVGART (efgartigimod alfa) in Japan for the treatment of primary immune thrombocytopenia (ITP), offering a new, much-needed treatment option for ITP patients in the region. This approval reflects Argenx's broader strategy to expand its therapeutic reach across different geographies.

Looking ahead, Argenx has laid out its long-term strategic vision, "Vision 2030," with the ambitious goal of extending its life-changing therapies to 50,000 patients by the end of the decade. This vision includes a focus on expanding its innovative pipeline and the development of new treatment candidates to address unmet needs in autoimmune diseases.

In summary, Argenx's recent achievements underscore its unwavering commitment to advancing the treatment landscape for autoimmune diseases. With a robust pipeline, solid financial foundation, and strategic vision for the future, Argenx is poised to continue its leadership in the field, bringing hope and new treatment options to patients around the world.