GENOMICS | GENETICS | R&D | DIAGNOSTIC TESTS

GENOMIC VISION SIGNS A DISTRIBUTION AGREEMENT WITH AMCARE GENOMICS LABORATORY IN CHINA FOR ITS FSHD DIAGNOSTIC ASSAY

The company accelerates its strategic development in China, the largest market in the world with 70.000 to 140.000 people suffering from Facioscapulohumeral muscular dystrophy

Bagneux (France), Guangzhou (China), October 19, 2017 – 7.30 am (CEST) - Genomic Vision (FR0011799907 – GV), a company specialized in the development of diagnostic tests for the early detection of cancers and hereditary diseases, and applications for life sciences research, today announced that it has signed an exclusive distribution agreement with AmCare Genomics Laboratory to market the FSHD diagnostic assay in China.

"We are thrilled to have acquired Genomic Vision's technology and signed an exclusive distribution agreement for the FSHD diagnostic assay in China. Thanks to our large network and connections within the neurology and FSHD communities in China, we are confident in raising significant awareness among physicians and patients on the most powerful diagnostic test for FSHD", **commented Dr. Victor Wei Zhang, CEO of AmCare Genomics Laboratory.**

"Following our distribution deal with APG Bio Ltd last June, this marketing agreement with AmCare represents a second major milestone in our global strategy to enter new markets in Asia. In China, it is estimated that 70,000 to 140,000 people are affected with FSHD. Moreover, the field of genomics and genetic testing is extremely developed, notably in large cities. We are convinced that our robust diagnostic technology will provide great benefits to families and patients. We look forward to deploying our FSHD solution in China and to strengthening our presence in the country through this valuable partnership", **added David Del Bourgo, Marketing & Sales Director of Genomic Vision.**

Facioscapulohumeral muscular dystrophy (FSHD) is the third most prevalent muscular hereditary myopathy worldwide. This genetic disease manifests as atrophy and weakness in the face, shoulders, and ambulatory muscles. There is great variability in clinical severity, from a severe infantile form to individuals who remain asymptomatic throughout their lives. This autosomal dominant disease is thought to affect 1 in 10,000 to 1 in 20,000 people. The actual number of individuals with FSHD worldwide is 870,000, which could be significantly higher due to undiagnosed cases. About 70% of FSHD patients inherit the disease from a parent, while 30% of the cases are sporadic and associated to de novo mutations.

ABOUT GENOMIC VISION

GENOMIC VISION is a company specialized in the development of diagnostic solutions for the early detection of cancers and serious genetic diseases and tools for life sciences research. Through the DNA Molecular Combing, a strong proprietary technology allowing to identify genetic abnormalities, GENOMIC VISION stimulates the R&D productivity of the pharmaceutical companies, the leaders of the diagnostic industry and the research labs. The Company develops a robust portfolio of diagnostic tests (breast, ovarian and colorectal cancers, myopathies) and analysis tools (DNA replication, biomarkers discovery, gene editing quality control). Based near Paris, in Bagneux, the Company has approximately 60 employees. GENOMIC VISION is a public listed company listed

in compartment C of Euronext's regulated market in Paris (Euronext: GV - ISIN: FR0011799907). For further information, please visit www.genomicvision.com

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FORWARD LOOKING STATEMENT

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Such forward-looking statements are based on assumptions that Genomic Vision considers to be reasonable. However, there can be no assurance that such forward-looking statements will be verified, which statements are subject to numerous risks, including the risks set forth in the "Risk Factors" section of the annual financial report dated April 29, 2016, available on the web site of Genomic Vision (www.genomicvision.com) and to the development of economic conditions, financial markets and the markets in which Genomic Vision operates. The forward-looking statements contained in this press release are also subject to risks not yet known to Genomic Vision or not currently considered material by Genomic Vision. The occurrence of all or part of such risks could cause actual results, financial conditions, performance or achievements of Genomic Vision to be materially different from such forward-looking statements.

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