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DIAGNOS Successfully Completes ISO 13485:2016 MDSAP Audit

Brossard, Quebec, Canada – April 23, 2019 - DIAGNOS Inc. ("DIAGNOS", "the Corporation" or "we") (TSX Venture: ADK) (OTCQB: DGNOF), a leader in early detection of critical health issues through the use of its *FL**AI**RE* platform based on Artificial Intelligence (**AI**), announces today that its quality management system continues to fully comply with the requirements of ISO 13485:2016 - Medical devices quality standard.

As part of the requirements for the commercialization of our flagship product CARA from Health Canada, the Food and Drug Agency in the US and CE in Europe, DIAGNOS must undergo thorough statutory annual quality compliance audits, such as the Medical Device Single Audit Program (MDSAP). The MDSAP is a comprehensive approach to quality system auditing and includes the development of an international coalition of countries devoted to pooling resources, technology, and services to enhance the safety and oversight of medical devices worldwide.

"I would like to take this opportunity to personally thank each of our employees for their hard work and commitment to the quality of our products. Our clients expect our healthcare solutions to perform well while being safe and DIAGNOS is able to meet their expectations", said Mr. André Larente, President of DIAGNOS.

About DIAGNOS

DIAGNOS is a publicly-traded Canadian corporation with a mission of early detection of critical health issues through the use of its Artificial Intelligence ("AI") tool CARA (Computer Assisted Retina Analysis). CARA is a tele-ophthalmology platform that integrates with existing equipment (hardware and software) and processes at the point of care. CARA's Artificial Intelligence image enhancement algorithms make standard retinal images sharper, clearer and easier to read. CARA is accessible securely over the internet, and is compatible with all recognized image formats and brands of fundus cameras, and is EMR compatible. CARA is a cost-effective tool for screening large numbers of patients in real-time and has been cleared for commercialization by several regulatory authorities such as Health Canada, the U.S. Food and Drug Administration and the European Union.

Additional information is available at www.diagnos.com and www.sedar.com

For further information about this press release, please contact:

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