

PRESS RELEASE 11/14/12

Ortho Kinematics Passes FDA Inspection With No Observations

AUSTIN, Texas – November 12, 2012 — Ortho Kinematics, a privately held healthcare diagnostics company focused on spine imaging informatics, announced today that it has been inspected by the U.S. Food and Drug Administration (FDA) without the issuance of a Form 483, which is used to report any non-compliance issues after a site inspection. The inspection took place from October 10-15, 2012.

Paul Gunnoe, CEO, says “this represents our company's first audit by the U.S. FDA and we are very pleased that we meet or exceed FDA requirements. We have invested heavily in preparation for this, and are very proud of the outcome.”

Adam Deitz, the company's Chief Technology Officer, continued “Quality is our watchword, and continuously improving the quality of our products and services is the foundation of our success.”

The Company's lead product, the VMA, is positioned to revolutionize the diagnostic side of the spine industry. The VMA technology provides clinicians with more accurate and reliable data, allowing them to make more informed diagnosis and treatment decisions.”

About Ortho Kinematics

Ortho Kinematics is a privately held diagnostic technology company, focused on spine imaging informatics and committed to the idea that spine motion matters. The company is working with a group of leading clinicians, researchers and developers who are passionate about leveraging spine biomechanical data to improve the diagnosis and treatment of back pain.

Ortho Kinematics is located in Austin, Texas and is on the web at www.orthokinematics.com. For more information, contact Ortho Kinematics Inc.

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