



Research Highlighting K2M's Lamellar 3D Titanium Technology™, BACS™ Surgical Planner, & RAVINE® Lateral Access System to Be Presented at EUROSPINE 2017

K2M to showcase differentiated product portfolio, 3D-printed offerings, & Balance ACS™ platform

LEESBURG, Va., Oct. 10, 2017 (GLOBE NEWSWIRE) -- [K2M Group Holdings, Inc.](#) (NASDAQ:KTWO) (the "Company" or "K2M"), a global leader of complex spine and minimally invasive solutions focused on achieving three-dimensional Total Body Balance™, today announced that clinical data related to K2M spinal solutions will be presented at [EUROSPINE 2017](#), occurring October 11—13, 2017, in Dublin. The Company will also showcase its comprehensive product portfolio, 3D-printed spinal solutions, and [Balance ACS™](#) (BACS™) platform at Booth F20.

"Taking a comprehensive approach to spine surgery allows me to address my patients' individual treatment needs more precisely," said Robert Lee, FRSC, TR & Orth, a spine surgeon at Royal National Orthopaedic Hospital NHS Trust in Stanmore, UK, and author of five studies on K2M products to be presented this week. "Our research supported favorable findings on surgical planning software, minimally invasive surgical techniques, and 3D porous CASCADIA™ interbody cages that utilize Lamellar 3D Titanium Technology™."

Data from five clinical studies on K2M's CASCADIA Interbody Systems featuring Lamellar 3D Titanium Technology, BACS Surgical Planner, and RAVINE® Lateral Access System will be presented:

- 1 3D Porous Lamellar Titanium Lateral Interbody Cages: Subsidence Rates Compared to PEEK and Early Clinical Outcomes, Lee R (Poster 26)
- 1 Clinical and Radiographic Results Following the Use of Lateral Interbody Cages to Restore Lumbar Lordosis in Revision Fusion Surgery, Lee R (Poster 27)
- 1 3D Porous Lamellar Titanium versus PEEK and Solid Titanium Interbody TLIF Cages: A Prospective Clinical and Radiographic Comparison, Lee R, Wilson L (Poster 28)
- 1 Minimally Invasive Anterior Column Reconstruction in Primary Adult Degenerative Scoliosis Correction Surgery, Lee R (Poster 84)
- 1 Reliability of Pre-Operative Surgical Planning Software in Predicting Postoperative Alignment in Patients Undergoing Minimally Invasive Multilevel Anterior Column Reconstruction for Positive Sagittal Balance Deformity, Lee R (Poster 85)

K2M Chairman, President, and CEO Eric Major said, "We are pleased by the amount of research on K2M products being presented at this year's EUROSPINE meeting. K2M is leading the way in 3D spinal innovation—as evidenced by our market-leading 3D-printed product portfolio and comprehensive Balance ACS platform—to help surgeons treat many different spinal pathologies both in European countries and across the globe."

At the meeting, K2M will showcase its comprehensive portfolio of complex spine, minimally invasive, and degenerative solutions, including the [CASCADIA Interbody Systems](#) featuring [Lamellar 3D Titanium Technology](#), the [EVEREST® Minimally Invasive XT Spinal System](#), the [EVEREST Deformity Spinal System](#), the [RAVINE Lateral Access System](#), the [MESA® Deformity Spinal System](#), the RHINE™ Cervical Disc System*, and more.

K2M will also feature its Balance ACS platform, which provides solutions focused on achieving balance of the spine by addressing each anatomical vertebral segment with a 360-degree approach to the axial, coronal, and sagittal planes, emphasizing Total Body Balance as an important component of surgical success. The Company will demo the [BACS Surgical Planner](#), a surgical image measuring technology that assists in planning and preoperative implant selection.

For more information on K2M's complete product portfolio, visit www.K2M.com. For more information on Balance ACS, visit www.BACS.com.

*This product is intended for export and is not sold or offered for sale in the United States.

About K2M

K2M Group Holdings, Inc. is a global leader of complex spine and minimally invasive solutions focused on achieving three-dimensional Total Body Balance. Since its inception, K2M has designed, developed, and commercialized innovative complex spine and minimally invasive spine technologies and techniques used by spine surgeons to treat some of the most complicated spinal pathologies. K2M has leveraged these core competencies into [Balance ACS](#), a platform of products, services, and research to help surgeons achieve three-dimensional spinal balance across the axial, coronal, and sagittal planes, with the goal of supporting the full continuum of care to facilitate quality patient outcomes. The Balance ACS platform, in combination with the Company's technologies, techniques and leadership in the 3D-printing of spinal devices, enable K2M to compete favorably in the global spinal surgery market. For more information, visit www.K2M.com and connect with us on [Facebook](#), [Twitter](#), [Instagram](#), [LinkedIn](#) and [YouTube](#).

Forward-Looking Statements

This press release contains forward-looking statements that reflect current views with respect to, among other things, operations and financial performance. Forward-looking statements include all statements that are not historical facts such as our statements about our expected financial results and guidance and our expectations for future business prospects. In some cases, you can identify these forward-looking statements by the use of words such as, "outlook," "guidance," "believes," "expects," "potential," "continues," "may," "will," "should," "could," "seeks," "predicts," "intends," "plans," "estimates," "anticipates" or the negative version of these words or other comparable words. Such forward-looking statements are subject to various risks and uncertainties including, among other things: our ability to achieve or sustain profitability in the future; our ability to demonstrate to spine surgeons the merits of our products; pricing pressures and our ability to compete effectively generally; collaboration and consolidation in hospital purchasing; inadequate coverage and reimbursement for our products from third-party payors; lack of long-term clinical data supporting the safety and efficacy of our products; dependence on a limited number of third-party suppliers; our ability to maintain and expand our network of direct sales employees, independent sales agencies and international distributors and their level of sales or distribution activity with respect to our products; proliferation of physician-owned distributorships in the industry; decline in the sale of certain key products; loss of key personnel; our ability to enhance our product offerings through research and development; our ability to manage expected growth; our ability to successfully acquire or invest in new or complementary businesses, products or technologies; our ability to educate surgeons on the safe and appropriate use of our products; costs associated with high levels of inventory; impairment of our goodwill and intangible assets; disruptions in our main facility or information technology systems; our ability to ship a sufficient number of our products to meet demand; our ability to strengthen our brand; fluctuations in insurance cost and availability; our ability to comply with extensive governmental regulation within the United States and foreign jurisdictions; our ability to maintain or obtain regulatory approvals and clearances within the United States and foreign jurisdictions; voluntary corrective actions by us or our distribution or other business partners or agency enforcement actions; recalls or serious safety issues with our products; enforcement actions by regulatory agencies for improper marketing or promotion; misuse or off-label use of our products; delays or failures in clinical trials and results of clinical trials; legal restrictions on our procurement, use, processing, manufacturing or distribution of allograft bone tissue; negative publicity concerning methods of tissue recovery and screening of donor tissue; costs and liabilities relating to environmental laws and regulations; our failure or the failure of our agents to comply with fraud and abuse laws; U.S. legislative or Food and Drug Administration regulatory reforms; adverse effects of medical device tax provisions; potential tax changes in jurisdictions in which we conduct business; our ability to generate significant sales; potential fluctuations in sales volumes and our results of operations over the course of the year; uncertainty in future capital needs and availability of capital to meet our needs; our level of indebtedness and the availability of borrowings under our credit facility; restrictive covenants and the impact of other provisions in the indenture governing our convertible senior notes and our credit facility; continuing worldwide economic instability; our ability to protect our intellectual property rights; patent litigation and product liability lawsuits; damages relating to trade secrets or non-competition or non-solicitation agreements; risks associated with operating internationally; fluctuations in foreign currency exchange rates; our ability to comply with the Foreign Corrupt Practices Act and similar laws; our ability to implement and maintain effective internal control over financial reporting; potential volatility in our stock due; our lack of current plans to pay cash dividends; our ability to take advantage of certain reduced disclosure requirements and exemptions as a result of being an emerging growth company; increased costs and additional regulations and requirements as a result of no longer qualifying as an emerging growth company as of December 31, 2017; potential dilution by the future issuances of additional common stock in connection with our incentive plans, acquisitions or otherwise; anti-takeover provisions in our organizational documents

and our ability to issue preferred stock without shareholder approval; potential limits on our ability to use our net operating loss carryforwards; and other risks and uncertainties, including those described under the section entitled "Risk Factors" in our most recent Annual Report on Form 10-K filed with the SEC and our Quarterly Report filed with the SEC on August 2, 2017, as such factors may be updated from time to time in our periodic filings with the SEC, which are accessible on the SEC's website at www.sec.gov. Accordingly, there are or will be important factors that could cause actual outcomes or results to differ materially from those indicated in these statements. These factors should not be construed as exhaustive and should be read in conjunction with the other cautionary statements that are included in this release and our filings with the SEC.

We operate in a very competitive and challenging environment. New risks and uncertainties emerge from time to time, and it is not possible for us to predict all risks and uncertainties that could have an impact on the forward-looking statements contained in this release. We cannot assure you that the results, events and circumstances reflected in the forward-looking statements will be achieved or occur, and actual results, events or circumstances could differ materially from those described in the forward-looking statements.

The forward-looking statements made in this press release relate only to events as of the date on which the statements are made. We undertake no obligation to publicly update or review any forward-looking statement, whether as a result of new information, future developments or otherwise, except as required by law. We may not actually achieve the plans, intentions or expectations disclosed in our forward-looking statements and you should not place undue reliance on our forward-looking statements. Unless specifically stated otherwise, our forward-looking statements do not reflect the potential impact of any future acquisitions, mergers, dispositions, joint ventures, investments or other strategic transactions we may make.

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