Zimmer Biomet Spine Announces Mobi-C® Cervical Disc Seven-Year Follow-Up Data Published in International Journal of Spine Surgery

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Results Demonstrate Clinical Superiority of Mobi-C Two-Level Cervical Total Disc Replacement Procedures Compared to Anterior Cervical Discectomy and Fusion

WARSAW, Ind., March 28, 2018 /PRNewswire/ -- Zimmer Biomet Holdings, Inc. (NYSE and SIX: ZBH), a global leader in musculoskeletal healthcare, today announced that the full seven-year follow-up data from a study evaluating the Company's Mobi-C® Cervical Disc for total disc replacement has been published in the peer-reviewed *International Journal of Spine Surgery*, the official scientific journal of the International Society for the Advancement of Spine Surgery (ISASS). The Mobi-C Cervical Disc is a leading option for cervical total disc replacement (cTDR) that is designed to maintain patient range of motion post-surgery. The paper can be accessed online at the following web address: http://ijssurgery.com/10.14444/4031. Complete disclosure of surgeon affiliations and potential conflicts of interest can be found in the article.

The article, titled "*Long-term Evaluation of Cervical Disc Arthroplasty with the Mobi-C® Cervical Disc: A Randomized, Prospective, Multicenter Clinical Trial with Seven-Year Follow-up,*" reports on the continuation of a prospective, randomized clinical trial of cTDR with Mobi-C compared to the previous standard of care, anterior cervical discectomy and fusion (ACDF), and follows the nearly 600 patients through seven years. Researchers found that cTDR with Mobi-C was associated with lower pain scores, maintained range of motion, less adjacent level degeneration and adjacent level subsequent surgery, as well as a lower rate of secondary surgery compared to ACDF.

"This study further validates the strengths of Mobi-C as a solution for cervical total disc replacement. The positive clinical outcomes remain consistent with previous findings, mainly that Mobi-C continues to deliver positive patient outcomes including efficacy, safety, patient satisfaction and fewer follow-up surgeries," said Joe Ross, General Manager of Zimmer Biomet's Spine division. "It is particularly important that the clinical effectiveness of cTDR versus ACDF becomes more apparent when treatment increases from one to two levels, indicating a significant benefit for Mobi-C in two-level procedures, which affects an estimated one in three cervical disc patients."

The published article discusses the Investigational Device Exemption/Post-Approval Study that evaluated 599 patients, including 164 treated with one-level cTDR, 225 treated with two-level cTDR, 81 treated with one-level ACDF and 105 treated with two-level ACDF. At seven years, follow-up rates ranged from 73.5 percent to 84.4 percent (overall 80.2 percent). Overall success rates for two-level patients were 60.8 percent for cTDR and 34.6 percent for ACDF. Overall success rates for one-level patients were 55.2 percent for cTDR and 50.0 percent for ACDF. Researchers concluded that the composite success analysis demonstrated clinical superiority of two-level cTDR over ACDF and non-inferiority of one-level cTDR versus ACDF. For two-level procedures, the Neck Disability Index (NDI) success rate was significantly greater in the cTDR group (79.0 percent) compared to ACDF (58.0 percent), marking significant patient-reported improvements with cTDR. In addition, overall, more than 86.0 percent of patients who received cTDR (two-level) and 73.9 percent of patients who underwent ACDF (two-level) reported they were "very satisfied" after seven years. Finally, the rate of subsequent surgery at the index level was significantly lower in the two-level cTDR group compared to the ACDF group (cTDR: 4.4 percent vs. ACDF:16.2 percent; p=0.001); and similarly, the rate of adjacent level secondary surgery was significantly lower in the two-level cTDR (4.4 percent) patients compared to the ACDF (11.4 percent; p=0.03) patients. The clinical data for the seven-year results and associated product labeling updates are currently under review with the U.S. Food and Drug Administration (FDA).

	Mobi-C	ACDF	
575 patients included in the 7-year analysis:			
Two-level	225	105	
One-level	164	81	
Overall Success Rate:			
Two-level	60.8%	34.6%	Clinical Superiority of Mobi-C versus ACDF (p<0.0001)
One-level	55.2%	50.0%	Non-inferiority of Mobi-C versus ACDF (p>0.05)
Two-level Neck Disability Index (NDI) success rate			
Two-level	79.0%	58.0%	p<0.05

One-level	76.5%	77.8%	p=NS
Patients (from both groups) reporting "very satisfied" with the procedure			
Two-level	86.0%	73.9%	p=0.039
One-level	90.9%	77.8%	p=0.028
Rate of secondary surgery at the index level			
Two-level	4.4%	16.2%	p=0.001
One-level	3.0%	12.3%	p=0.008
Rate of secondary surgery at an adjacent level			
Two-level	4.4%	11.4%	p=0.03
One-level	3.7%	13.6%	p=0.007

Mobi-C Cervical Disc

Mobi-C is the first cervical disc prosthesis approved by the FDA for reconstruction of a cervical disc at *both* one and two levels (C3-C7). Mobi-C is a cobalt chromium alloy and polyethylene mobile-bearing prosthesis that is inserted in a single step, without requiring bone chiseling to accommodate vertebral anchorage such as screws or keels. The Mobi-C Cervical Disc Prosthesis is indicated in skeletally mature patients for reconstruction of the disc from C3-C7 following discectomy at one level or two contiguous levels for intractable radiculopathy (arm pain and/or neurological deficit) with or without neck pain or myelopathy due to abnormality localized to the level of the disc space and at least one of the following conditions confirmed by radiographic imaging (CT, MRI or X-rays): herniated nucleus pulposus, spondylosis (defined by the presence of osteophytes) and/or visible loss of disc height compared to adjacent levels. The Mobi-C Cervical Disc Prosthesis is implanted using an anterior approach. Patients should have failed at least six weeks of conservative treatment or demonstrated progressive signs or symptoms despite nonoperative treatment prior to implantation of the Mobi-C Cervical Disc Prosthesis.

About Zimmer Biomet

Founded in 1927 and headquartered in Warsaw, Indiana, Zimmer Biomet is a global leader in musculoskeletal healthcare. We design, manufacture and market orthopaedic reconstructive products; sports medicine, biologics, extremities and trauma products; office based technologies; spine, craniomaxillofacial and thoracic products; dental implants; and related surgical products.

We collaborate with healthcare professionals around the globe to advance the pace of innovation. Our products and solutions help treat patients suffering from disorders of, or injuries to, bones, joints or supporting soft tissues. Together with healthcare professionals, we help millions of people live better lives.

We have operations in more than 25 countries around the world and sell products in more than 100 countries. For more information, visit www.zimmerbiomet.com or follow Zimmer Biomet on Twitter at www.twitter.com/zimmerbiomet.

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This release contains forward-looking statements within the meaning of the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. Forward-looking statements include, but are not limited to, statements concerning Zimmer Biomet's expectations, plans, prospects, and product and service offerings, including new product launches and potential clinical successes. Such statements are based upon the current beliefs and expectations of management and are subject to significant risks, uncertainties and changes in circumstances that could cause actual outcomes and results to differ materially. For a list and description of some of such risks and uncertainties, see Zimmer Biomet's periodic reports filed with the U.S. Securities and Exchange Commission (SEC). These factors should not be construed as exhaustive and should be read in conjunction with the other cautionary statements that are included in Zimmer Biomet's filings with the SEC. Forward-looking statements speak only as of the date they are made, and Zimmer Biomet disclaims any intention or obligation to update or revise any forward-looking statements, whether as a result of new information, future events or otherwise. Readers of this release are cautioned not to rely on these forward-looking statements, since there can be no assurance that these forward-looking statements will prove to be accurate. This cautionary statement is applicable to all forward-looking statements contained in this release.

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