

BIOMET INC

Form 424B3

August 29, 2013

Filed Pursuant to Rule 424(b)(3)

Registration No. 333-188262

PROSPECTUS SUPPLEMENT

(to prospectus dated June 21, 2013 and the prospectus supplements dated July 11, 2013, July 18, 2013 and August 29, 2013)

BIOMET, INC.

\$1,825,000,000 6.500% Senior Notes due 2020

\$800,000,000 6.500% Senior Subordinated Notes due 2020

This prospectus supplement updates and supplements the prospectus dated June 21, 2013 and the prospectus supplements dated July 11, 2013, July 18, 2013 and August 29, 2013.

See the "Risk Factors" section beginning on page 6 of the prospectus and the "Risk Factors" section in our Annual Report on Form 10-K filed with the Securities and Exchange Commission on August 29, 2013 for a discussion of certain risks that you should consider before investing in the notes.

Neither the Securities and Exchange Commission nor any state securities commission has approved or disapproved of these securities or passed upon the adequacy or accuracy of this prospectus. Any representation to the contrary is a criminal offense.

This prospectus supplement and the accompanying prospectus have been prepared for and may be used by Goldman, Sachs & Co. and any affiliates of Goldman, Sachs & Co. in connection with offers and sales of the notes related to market-making transactions in the notes affected from time to time. Goldman, Sachs & Co. or its affiliates may act as principal or agent in such transactions, including as agent for the counterparty when acting as principal or as agent for both counterparties, and may receive compensation in the form of discounts and commissions, including from both counterparties, when it acts as agents for both. Such sales will be made at prevailing market prices at the time of sale, at prices related thereto or at negotiated prices. We will not receive any proceeds from such sales.

You should rely only on the information contained or incorporated by reference in this prospectus supplement and the accompanying prospectus. We have not authorized any person to provide you with any information or represent anything about us or this offering that is not contained in this prospectus supplement and the accompanying prospectus. If given or made, any such other information or representation should not be relied upon as having been authorized by us. This prospectus supplement and the accompanying prospectus does not offer to sell nor ask for offers to buy any of the securities in any jurisdiction where it is unlawful, where the person making the offer is not qualified to do so, or to any person who cannot legally be offered the securities. You should not assume that the information contained or incorporated by reference in this prospectus supplement and the accompanying prospectus is accurate as of any date other than the date on the front cover of this prospectus supplement and the accompanying prospectus or the date of any document incorporated by reference herein.

The date of this prospectus supplement is August 29, 2013.

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UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 10-K

(Mark One)

ANNUAL REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934

For the fiscal year ended May 31, 2013.

OR

TRANSITION REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934

For the transition period from _____ to _____
Commission File Number 001-15601

LVB ACQUISITION, INC.

BIOMET, INC.

(Exact name of registrant as specified in its charter)

| | |
|--|--------------------------------------|
| Delaware | 26-0499682 |
| Indiana | 35-1418342 |
| (State or other jurisdiction of incorporation or organization) | (I.R.S. Employer Identification No.) |

| | |
|--|------------|
| 56 East Bell Drive, Warsaw, Indiana | 46582 |
| (Address of principal executive offices) | (Zip Code) |

(574) 267-6639

(Registrant's telephone number, including area code)

Securities registered pursuant to Section 12(b) of the Act: None

Securities registered pursuant to Section 12(g) of the Act: LVB Acquisition, Inc. common stock, par value \$0.01 per share

Indicate by check mark if the registrant is a well-known seasoned issuer, as defined in Rule 405 of the Securities Act

| | | | |
|-----------------------|-----|----|---|
| LVB ACQUISITION, INC. | Yes | No | x |
|-----------------------|-----|----|---|

| | | | |
|--------------|-----|----|---|
| BIOMET, INC. | Yes | No | x |
|--------------|-----|----|---|

Indicate by check mark if the registrant is not required to file reports pursuant to Section 13 or Section 15(d) of the Act.

| | | | |
|-----------------------|-----|----|---|
| LVB ACQUISITION, INC. | Yes | No | x |
|-----------------------|-----|----|---|

| | | | |
|--------------|-----|----|---|
| BIOMET, INC. | Yes | No | x |
|--------------|-----|----|---|

Indicate by check mark whether the registrant (1) has filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was required to file such reports), and (2) has been subject to such filing requirements for the past 90 days.

| | | | |
|-----------------------|-----|---|----|
| LVB ACQUISITION, INC. | Yes | x | No |
|-----------------------|-----|---|----|

| | | | |
|--------------|-----|---|----|
| BIOMET, INC. | Yes | x | No |
|--------------|-----|---|----|

Indicate by check mark whether the registrant has submitted electronically and posted on its corporate Website, if any, every Interactive Data File required to be submitted and posted pursuant to Rule 405 of Regulation S-T (§ 232.405 of

this chapter) during the preceding 12 months (or for such shorter period that the registrant was required to submit and post such files).

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| | | | |
|-----------------------|-----|---|----|
| LVB ACQUISITION, INC. | Yes | x | No |
| BIOMET, INC. | Yes | x | No |

Indicate by check mark if disclosure of delinquent filers pursuant to Item 405 of Regulation S-K (§ 229.405 of this chapter) is not contained herein, and will not be contained, to the best of registrant’s knowledge, in definitive proxy or information statements incorporated by reference in Part III of this Form 10-K or any amendment to this Form 10-K.

| | |
|-----------------------|---|
| LVB ACQUISITION, INC. | |
| BIOMET, INC. | ý |

Indicate by check mark whether the registrant is a large accelerated filer, an accelerated filer, a non-accelerated filer, or a smaller reporting company. See the definitions of “large accelerated filer,” “accelerated filer” and “smaller reporting company” in Rule 12b-2 of the Exchange Act. (Check one):

| | | |
|-------------------------|---|---------------------------|
| LVB ACQUISITION, INC. | | |
| Large accelerated filer | | Accelerated filer |
| Non-accelerated filer | x (Do not check if a smaller reporting company) | Smaller reporting company |
| BIOMET, INC. | | |
| Large accelerated filer | | Accelerated filer |
| Non-accelerated filer | x (Do not check if a smaller reporting company) | Smaller reporting company |

Indicate by check mark whether the registrant is a shell company (as defined in Rule 12b-2 of the Act).

| | | | |
|-----------------------|-----|----|---|
| LVB ACQUISITION, INC. | Yes | No | x |
| BIOMET, INC. | Yes | No | x |

As of May 31, 2013, there was no established public trading market for any of the common stock of the registrants. The number of shares of the registrants’ common stock outstanding as of July 31, 2013:

| | |
|-----------------------|------------------------------------|
| LVB ACQUISITION, INC. | 552,359,416 shares of common stock |
| BIOMET, INC. | 1,000 shares of common stock |

DOCUMENTS INCORPORATED BY REFERENCE
None.

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FORWARD-LOOKING STATEMENTS

This annual report contains forward-looking statements within the meaning of the U.S. federal securities laws. Statements that are not historical facts, including statements about our beliefs and expectations, are forward-looking statements. Forward-looking statements include statements generally preceded by, followed by, or that include the words “believe,” “could,” “expect,” “forecast,” “intend,” “may,” “anticipate,” “plan,” “predict,” “possibly,” “project,” “potential,” “should,” “will” or similar expressions. These statements include, but are not limited to, statements related to:

- the timing and number of planned new product introductions;
- the effect of anticipated changes in the size, health and activities of the population or on the demand for our products;
- assumptions and estimates regarding the size and growth of certain market categories;
- our ability and intent to expand in key international markets;
- the timing and anticipated outcome of clinical studies;
- assumptions concerning anticipated product developments and emerging technologies;
- the future availability of raw materials;
- the anticipated adequacy of our capital resources to meet the needs of our business;
- our continued investment in new products and technologies;
- the ultimate marketability of products currently being developed;
- our ability to successfully implement new technologies and transition certain manufacturing operations, including transitions to China;
- our ability to manage working capital and generate adequate cash flows to service outstanding debt;
- our ability to sustain sales and earnings growth;
- our success in achieving timely approval or clearance of our products with domestic and foreign regulatory entities;
- our success in implementing our operational improvement programs;
- the stability of certain foreign economic markets;
- the effect of foreign currency fluctuations on our results;
- the impact of anticipated changes in the musculoskeletal industry and our ability to react to and capitalize on those changes;
- our ability to successfully implement desired organizational changes;
- the impact of our managerial changes; and
- our ability to take advantage of technological advancements.

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Forward-looking statements reflect our current expectations and are not guarantees of performance. These statements are based on our management’s beliefs and assumptions, which in turn are based on currently available information. Important assumptions relating to these forward-looking statements include, among others, assumptions regarding demand for our products, expected pricing levels, raw material costs, the timing and cost of planned capital expenditures, future regulatory reforms affecting the healthcare industry, expected outcomes of pending litigation and regulatory matters, the solvency of our insurers and the ultimate resolution of allocation and coverage issues with those insurers, competitive conditions and general economic conditions. Readers of this annual report are cautioned that reliance on any forward-looking statement involves risks and uncertainties.

Although we believe that the assumptions on which the forward-looking statements contained herein are based are reasonable, any of those assumptions could prove to be inaccurate given the inherent uncertainties as to the occurrence or nonoccurrence of future events. There can be no assurance that the forward-looking statements contained in this annual report will prove to be accurate. The inclusion of a forward-looking statement in this annual report should not be regarded as a representation by us that our objectives will be achieved. Forward-looking statements also involve risks and uncertainties, which could cause actual results to differ materially from those projected by any forward-looking statement. Many of these factors are beyond our ability to control or predict and could, among other things, cause actual results to differ from those contained in forward-looking statements made or incorporated by reference in this annual report and presented elsewhere by management from time to time. Such factors, among others, may have a material adverse effect upon our business, financial condition, results of operations and cash flows and may include, but are not limited to, factors discussed under the heading “Risk Factors” and the following:

- changes in general economic conditions and interest rates;
- changes in the availability of capital and financing sources;
- changes in competitive conditions and prices in our markets;
- changes to the regulatory environment for our products, including national health care reform;
- the effects of incurring or having incurred a substantial amount of indebtedness under our 6.5% senior notes, 6.5% senior subordinated notes and senior secured credit facilities;
- the effects upon us of complying with the covenants contained in our senior secured credit facilities and the indentures governing our 6.5% senior notes and 6.5% senior subordinated notes;
- restrictions that the terms and conditions of indentures governing our 6.5% senior notes and 6.5% senior subordinated notes and our senior secured credit facilities may place on our ability to respond to changes in our business or take certain actions;
- changes in the relationship between supply of and demand for our products;
- fluctuations in costs of raw materials and labor;
- the effect of foreign currency fluctuations on our results;
- changes in other significant operating expenses;
- decreases in sales of our principal product lines;
- slowdowns or inefficiencies in our product research and development efforts;
- increases in expenditures related to increased government regulation of our business;
- developments adversely affecting our sales activities inside or outside the United States;
- decreases in reimbursement levels by our customers, including certain of our foreign government customers that are experiencing financial distress;

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- difficulties in transitioning certain manufacturing operations to China and other locations;
- challenges in effectively implementing restructuring and cost saving initiatives;
- increases in cost-containment efforts from managed care organizations and other third-party payors;
- loss of our key management and other personnel or inability to attract such management and other personnel;
- increases in costs of retaining existing independent sales agents of our products;
- potential future goodwill and/or intangible impairment charges;
- unanticipated expenditures related to litigation; and
- failure to comply with the terms of the Deferred Prosecution Agreement.

We caution you not to place undue reliance on these forward-looking statements, which speak only as of the date they were made. We do not undertake any obligation to publicly release any revisions to these forward-looking statements to reflect events or circumstances after the date of this annual report or to reflect the occurrence of unanticipated events. We intend to take advantage of the safe harbor provisions of the Private Securities Litigation Reform Act of 1995 regarding our forward-looking statements, and are including this sentence for the express purpose of enabling us to use the protections of the safe harbor with respect to all forward-looking statements.

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Part I.

Explanatory Note

This Form 10-K is a combined annual report being filed separately by two registrants: LVB Acquisition, Inc. (“LVB” and “Parent”) and its wholly owned subsidiary, Biomet, Inc. Each registrant hereto is filing on its own behalf all of the information contained in this annual report that relates to such registrant. Each registrant hereto is not filing any information that does not relate to such registrant, and therefore makes no representation as to any such information.

Item 1. Business.

General

Currently, the principal asset of LVB is the ownership of 100% of the common stock of Biomet, Inc., which is an operating company. Biomet, Inc., an Indiana corporation incorporated in 1977, is one of the largest orthopedic medical device companies in the United States and worldwide with operations in more than 50 locations throughout the world and distribution in approximately 90 countries. Biomet, Inc.’s principal operating subsidiaries include Biomet U.S. Reconstruction, LLC; Biomet Orthopedics, LLC; Biomet Manufacturing, LLC; Biomet Europe BV; EBI, LLC; Biomet 3i, LLC; Biomet International Ltd.; Biomet Microfixation, LLC; Biomet Sports Medicine, LLC; Biomet Trauma, LLC; and Biomet Biologics, LLC. Unless the context requires otherwise, the term “LVB,” “Biomet,” “Company,” “we,” “our”, or “us” refers to LVB Acquisition, Inc. and all of its subsidiaries. We design, manufacture and market a comprehensive range of both surgical and non-surgical products used primarily by orthopedic surgeons and other musculoskeletal medical specialists. For over 30 years, we have applied advanced engineering and manufacturing technology to the development of highly durable joint replacement systems.

Transactions with the Sponsor Group

On December 18, 2006, Biomet, Inc. entered into an Agreement and Plan of Merger with LVB Acquisition, LLC, a Delaware limited liability company, which was subsequently converted to a corporation, LVB Acquisition, Inc., and LVB Acquisition Merger Sub, Inc., an Indiana corporation and a wholly-owned subsidiary of LVB (“Purchaser”), which agreement was amended and restated as of June 7, 2007 and which we refer to as the “Merger Agreement.” Pursuant to the Merger Agreement, on June 13, 2007, Purchaser commenced a cash tender offer (the “Offer”) to purchase all of Biomet, Inc.’s outstanding common shares, without par value (the “Shares”) at a price of \$46.00 per Share (the “Offer Price”). Approximately 82% of the outstanding Shares were tendered to Purchaser in the Offer. At Biomet, Inc.’s special meeting of shareholders held on September 5, 2007, more than 91% of its shareholders voted to approve the proposed merger, and LVB acquired Biomet, Inc. on September 25, 2007 through a reverse subsidiary merger, with Biomet, Inc. being the surviving company (the “Merger”). Subsequent to the acquisition, Biomet, Inc. became a subsidiary of LVB, which is controlled by LVB Acquisition Holding, LLC, or “Holding,” an entity controlled by a consortium of private equity funds affiliated with the Sponsors (as defined below) and their co-investors.

Competitive Strengths

We believe we have a number of competitive strengths that will enable us to further enhance our position in the orthopedic medical device market.

Broad Market Leadership. We believe we are the fourth largest player in the U.S. orthopedic reconstructive market and have maintained this position for over a decade. In addition, we are a leading provider of dental reconstructive devices worldwide and maintain market leadership positions in the electrical stimulation and craniomaxillofacial fields.

Strong Relationships with Surgeon Customers. As a result of their satisfaction with our products, we enjoy long-standing relationships with our surgeon customers, many of which commence during the surgeons’ residency training programs. Our support of medical education programs provides important training opportunities for orthopedic surgeons early in their careers. Supporting “hands-on” training provides opportunities for residents, fellows and attending surgeons to experience the clinical benefits of our products. Surgeons have historically exhibited limited willingness to switch manufacturers, as successful patient outcomes are related to the practitioners’ familiarity with the procedural characteristics and instrumentation of certain implants.

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Consistently Strong Operating Cash Flow Generation. Our business is characterized by consistently strong operating cash flows due to our robust operating history and moderate capital intensity. We have continually increased revenues, with fiscal year 2013 representing our 35th consecutive year of year-over-year net sales growth. Over the last 20 years, from fiscal year 1993 through fiscal year 2013, we increased net sales at a compounded annual growth rate of approximately 12%. We have sustained growth through multiple macro-economic cycles, demonstrating a stable business profile. In addition, we have historically had modest capital expenditures and working capital requirements, providing for strong operating cash flow conversion.

Experienced and Dedicated Management Team. We have a highly experienced management team at both the corporate and operational level. Our team is led by Jeffrey R. Binder, a 22-year veteran of the orthopedic medical device industry, who was appointed President and Chief Executive Officer in February 2007. Daniel P. Florin was appointed Senior Vice President and Chief Financial Officer in June 2007 and brings 21 years of financial officer/controller experience in the medical device industry and five years of public accounting and auditing experience to us. In February 2008, Jon C. Serbousek was appointed President of Biomet Orthopedics and was recently appointed as President, Biomet Biologics, having spent 8 years with Medtronic and 13 years with DePuy, for a total of 26 years in the medical device industry. Adam Johnson was appointed Senior Vice President and President of EBI, LLC, d/b/a Biomet Spine & Bone Healing Technologies in June 2012, having previously served and continuing to serve as President of Biomet Microfixation and brings 13 years of experience in the medical device industry.

Premier Equity Sponsorship. The Blackstone Group, Goldman, Sachs & Co., Kohlberg Kravis Roberts & Co. and TPG (together the “Sponsors”) are among the most well-known and respected financial sponsors in the world. The Sponsors have collectively made investments in over 950 companies. The Sponsors have considerable experience in the healthcare sector with investments in companies such as HCA Holdings, Inc., IASIS Healthcare Corporation, Quintiles Transnational Corp., DJO Global and Vanguard Health Systems, Inc., among others.

Economic Uncertainties

Our results of operations could be substantially affected not only by global economic conditions, but also by local operating and economic conditions, which can vary substantially by market. Unfavorable conditions can depress sales in a given market and may result in actions that adversely affect our margins, constrain our operating flexibility or result in charges which are unusual or non-recurring.

Regulatory and Other Uncertainties

In the United States, healthcare providers that purchase our products (e.g., hospitals, physicians, dentists and other health care providers) generally rely on payments from third-party payors (principally federal Medicare, state Medicaid and private health insurance plans) to cover all or a portion of the cost of our musculoskeletal products. In March 2010, comprehensive health care reform legislation was enacted through the passage of the Patient Protection and Affordable Health Care Act (H.R. 3590) and the Health Care and Education Reconciliation Act (H.R. 4872). Among other initiatives, these laws impose a 2.3% excise tax on domestic sales of medical devices following December 31, 2012, which is estimated to contribute approximately \$20 billion to healthcare reform. Various healthcare reform proposals have also emerged at the state level. Outside of the excise tax, which has impacted our results of operations and cash flows following December 31, 2012, we cannot predict with certainty what healthcare initiatives, if any, will be implemented at the state level, or what the ultimate effect of federal health care reform or any future legislation or regulation will have on us. However, an expansion in government’s role in the U.S. healthcare industry may lower reimbursements for our products, reduce medical procedure volumes and adversely affect our business, results of operations and cash flows, possibly materially.

Outside the United States, reimbursement systems vary significantly from country to country. If adequate levels of reimbursement from third-party payors outside the United States are not obtained, international sales of our products may decline. Many foreign markets, including Canada and some European and Asian countries, have tightened reimbursement rates. Our ability to continue to sell certain products profitably in these markets may diminish if the government-managed healthcare systems continue to reduce reimbursement rates, which can decrease pricing and procedural volume.

Business Strategy

We intend to enhance our position as a leading orthopedic medical device company by pursuing the following strategic initiatives:

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Continue to Develop and Launch New Products and Technologies. The New Product Introduction (“NPI”) process is a global portfolio and project management approach that helps bring visibility and control to all commercial aspects of new product development projects. The process, which is managed by each of our global Product organizations, breaks each project down into six stages of work and further divides these stages by formal review gates. We have a single database of all of our development projects that is easily filtered and sorted to generate customized project roadmaps that serve as communication tools providing visibility to all functional teams. The database is designed to prioritize and focus the portfolio and also ensure that the workload is properly resourced and managed across the business. Projects are assessed against pre-determined gate criteria. The global Product organizations select and prioritize projects that can be adequately resourced and help deliver product category growth targets, satisfy specific hurdle rates and strategic drivers and provide a balanced product portfolio.

Enhance Surgeon Customer Relationships through Product Performance and Innovation. We intend to continue to meet the needs of our surgeon customers and hospital customers by providing clinically successful and innovative products that offer a cost-effective means of treating patients. Our success has been built on responsiveness to the needs