BIOMET INC
Form 424B3
July 18, 2013

Filed Pursuant to Rule 424(b)(3)
Registration No. 333-188262
PROSPECTUS SUPPLEMENT
(to prospectus dated June 21, 2013 and the prospectus supplement dated July 11, 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 supplement dated July 11, 2013.

See the "Risk Factors" section beginning on page 6 of the prospectus 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 July 18, 2013.

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

FORM 8-K

CURRENT REPORT
Pursuant to Section 13 or 15(d) of
the Securities Exchange Act of 1934
Date of Report (Date of earliest event reported): July 16, 2013

LVB ACQUISITION, INC.
BIOMET, INC.
(EXACT NAME OF REGISTRANT AS SPECIFIED IN ITS CHARTER)

Delaware	000-54505	26-0499682
Indiana	001-15601	35-1418342
(State or other jurisdiction of incorporation)	(Commission File Number)	(IRS Employer Identification No.)

56 East Bell Drive
Warsaw, Indiana 46582
(Address of Principal Executive Offices, Including Zip Code)
(574) 267-6639
(Registrant's Telephone Number, Including Area Code)
Not Applicable
(Former Name or Former Address, if Changed Since Last Report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

¨ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
¨ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
¨ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
¨ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Item 8.01. Other Events.

On July 16, 2013, Biomet, Inc. issued a press release announcing that the Company will close its manufacturing facility in Le Locle, Switzerland.

The press release announcing the facility closure is filed herewith as exhibit 99.1, and is incorporated by reference herein.

Item 9.01. Financial Statements and Exhibits.

Exhibit No.	Document
99.1	Press Release issued July 16, 2013 (translated into English from the French original).

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrants have duly caused this report to be signed on their behalf by the undersigned hereunto duly authorized.

Date: July 18, 2013

LVB ACQUISITION, INC.

/s/ Daniel P. Florin
By: Daniel P. Florin
Its: Senior Vice President and Chief Financial Officer

BIOMET, INC.

/s/ Daniel P. Florin
By: Daniel P. Florin
Its: Senior Vice President and Chief Financial Officer

Exhibit 99.1

Biomet announces conclusion of consultation process at Le Locle, Switzerland manufacturing facility

16 July, 2013--Biomet, Inc., a global leader in the manufacture of orthopaedic and biotechnology products, announced today that it will close its manufacturing facility in Le Locle, Switzerland, which employs approximately 230 workers. The facility will cease a majority of its production by June, 2014. Employees will be given appropriate notice, in compliance with applicable laws and no dismissals related to the plant closure will occur prior to June, 2014. Biomet intends to transfer the production to other facilities within its global manufacturing network and to other Biomet-certified sources.

The closure is in response to declining global prices for medical devices and a challenging economic environment, which necessitates that the Company continually and sustainably improves the cost-efficiency of its global manufacturing operations in the changing medical device market, and to ensure that the Company is able to cost-effectively provide the highest-quality products to healthcare professionals and their patients.

Where possible, the Company will explore other employment options for affected team members. In cases where other positions are not available, the Company intends to provide severance/social plan and outplacement assistance to support the team members in their transitions.

About Biomet

Biomet, Inc. and its subsidiaries design, manufacture and market products used primarily by musculoskeletal medical specialists in both surgical and non-surgical therapy. Biomet's product portfolio encompasses large joint reconstructive products, including orthopedic joint replacement devices, and bone cements and accessories; sports medicine, extremities and trauma products, including internal and external orthopedic fixation devices; spine and bone healing products, including spine hardware, spinal stimulation devices, and orthobiologics, as well as electrical bone growth stimulators; dental reconstructive products; and other products, including microfixation products and autologous therapies. Headquartered in Warsaw, Indiana, Biomet and its subsidiaries currently distribute products in approximately 90 countries.

Contacts

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Forward-Looking Statements

This press release contains “forward-looking statements” within the meaning of Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934, as amended. Those statements are often indicated by the use of words such as “will,” “intend,” “anticipate,” “estimate,” “expect,” “plan” and similar expressions. Forward-looking statements involve certain risks and uncertainties. Actual results may differ materially from those contemplated by the forward looking statements due to, among others, the following factors: the success of the Company’s principal product lines; the results of the ongoing investigation by the United States Department of Justice; the ability to successfully implement new technologies; the Company’s ability to sustain sales and earnings growth; the Company’s success in achieving timely approval or clearance of its products with domestic and foreign regulatory entities; the impact to the business as a result of compliance with federal, state and foreign governmental regulations and with the Deferred Prosecution Agreement; the impact to the business as a result of the economic downturn in both foreign and domestic markets; the impact of federal health care reform; the impact of anticipated

changes in the musculoskeletal industry and the ability of the Company to react to and capitalize on those changes; the ability of the Company to successfully implement its desired organizational changes and cost-saving initiatives; the ability of the Company to successfully integrate the Trauma Acquisition; the impact to the business as a result of the Company's significant international operations, including, among others, with respect to foreign currency fluctuations and the success of the Company's transition of certain manufacturing operations to China; the impact of the Company's managerial changes; the ability of the Company's customers to receive adequate levels of reimbursement from third-party payors; the Company's ability to maintain its existing intellectual property rights and obtain future intellectual property rights; the impact to the business as a result of cost containment efforts of group purchasing organizations; the Company's ability to retain existing independent sales agents for its products; the impact of product liability litigation losses; and other factors set forth in the Company's filings with the SEC, including the Company's most recent annual report on Form 10-K and quarterly reports on Form 10-Q. Although the Company believes 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 non-occurrence of future events. There can be no assurance as to the accuracy of forward-looking statements contained in this press release. The inclusion of a forward-looking statement herein should not be regarded as a representation by the Company that the Company's objectives will be achieved. The Company undertakes no obligation to update publicly or revise any forward-looking statements, whether as a result of new information, future events or otherwise. Accordingly, the reader is cautioned not to place undue reliance on forward-looking statements which speak only as of the date on which they were made.