



Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485: 2003

This is to certify that:

Biomet Biologics
Also trading as
Biomet Biologics, Inc.
56 East Bell Drive
P.O. Box 587
Warsaw
Indiana
46581
USA

Holds Certificate No: **FM 547157**

and operates a Quality Management System which complies with the requirements of ISO 13485: 2003 for the following scope:

The design, manufacture and distribution of sterile disposable medical equipment for collecting and processing blood and blood constituents for autologous use, calcium based bone void fillers, sterile operating room supplies, invasive surgical equipment, sterile surgical instruments, non-surgical instruments and reusable surgical instruments which can be connected to an active medical device.

Trading names which apply at this location include: Biomet Biologics; Biomet Biologics, Inc.; Cell Factor Technologies, Inc.

For and on behalf of BSI:

VP Regulatory Affairs, BSI Group America Inc.

Originally Registered: **02/18/2009**

Effective Date: **10/19/2011**

Expiry Date: **10/18/2014**



CMDCAS
Recognized
Registrar



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This certificate remains the property of BSI and shall be returned immediately upon request.
An electronic certificate can be authenticated [online](http://www.bsigroup.com/ClientDirectory). Printed copies can be validated at www.bsigroup.com/ClientDirectory
To be read in conjunction with the scope above or the attached appendix.

Americas Headquarters: 12110 Sunset Hills Road, Suite 200, Reston, VA 20190, USA.

