



Production Quality Assurance

No. CE 69332

Issued to:

Biomet Biologics
also trading as **Cell Factor Technologies, Inc.**
56 East Bell Drive
Warsaw
Indiana
46582
USA

In respect of:

The manufacture of sterile, disposable medical devices for collecting and processing blood and blood constituents for autologous use and bone graft mixing and delivery systems

on the basis of our examination under the requirements of Council Directive 93/42/EEC, Annex V, Section 3.2.

For and on behalf of the British Standards Institution, a Notified Body for the above Directive (Notified Body Number 0086):

David Ford, Managing Director, BSI Product Services

First Issued: 2 Oct 2002

Date: 1 Dec 2008

Expiration Date: 1 Oct 2012

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Conditions of Approval

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate unless specifically agreed with BSI.

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