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

01-50-1454
Revision A
Date: 2012-06

Blood Draw Kit

ATTENTION ADMINISTERING PERSONNEL

NOTE: FOR SINGLE USE ONLY. Discard the entire disposable kit after one use, using acceptable disposal method for products potentially contaminated with blood.

DESCRIPTION

Blood Draw Kit

The Blood Draw Kit contains syringes, needles and other accessories required for the collection of a patient's blood for processing in Biomet Biologics devices.

MATERIALS

Blood-draw components are packaged, labeled and sterilized as indicated by the manufacturer's labeling**.** All components in this kit are latex free.

Contents include:

60ml Luer-Lok syringe	Qty. 1
30ml Luer-Lok syringe	Qty. 2
20ml Luer-Lok syringe	Qty. 1
10ml Luer-Lok syringe	Qty. 2
2"x 2" Gauze	Qty. 2
Fistula Needle Set w/instructions	Qty. 1
18ga Needles	Qty. 5
Blank labels	Qty. 3

WARNINGS AND PRECAUTIONS

1. Users should exercise caution when handling surgical needles to avoid inadvertent needle sticks. Discard used needles in "sharps" containers.
2. Do not use sterile components in this kit if package is opened or damaged.
3. Single use device. Do not reuse.
4. The patient is to be made aware of general risks associated with treatment and the possible adverse effects.

POSSIBLE ADVERSE EFFECTS FROM DRAWING BLOOD

1. Damage to blood vessels, hematoma, delayed wound healing and/or infection.
2. Temporary or permanent nerve damage that may result in pain or numbness.
3. Infection associated with blood draw location.

STERILITY

All components supplied in this kit are sterilized by the respective suppliers using irradiation, ethylene oxide gas (ETO) or Heat. Do not re-sterilize. Do not use after expiration date. Do not use any component from an opened or damaged package. Single Use Only. Do not Reuse.

INSTRUCTIONS FOR USE

Refer to the Package Insert (IFU) of the Biomet Biologics kit being used for Instructions.

CAUTION: Federal Law (USA) restricts this device to sale by or on the order of a physician.

Comments regarding this kit can be directed to Attn: Regulatory Dept, Biomet, Inc., P.O. Box 587, Warsaw, IN 46581 USA. FAX: 574-372-3968.

CE Mark on the package insert (IFU) is not valid unless there is a CE Mark on the product (description) label.

Authorized Representative: Biomet U.K., Ltd.
Waterton Industrial Estate
Bridgend, South Wales
CF31 3XA UK

Symbol Legend

 Manufacturer

 Date of manufacture

 Do not reuse

 Caution, see instructions for use

 Sterilized using ethylene oxide

 Sterilized using irradiation

 Sterile

 Sterilized using aseptic processing techniques

 Sterilized using steam or dry heat

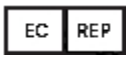
 Use by date

 WEEE device

 Catalogue number

 Batch code

 Flammable

 Authorized representative in the European Community