

# For an easy-to-use solution, choose the Codman<sup>®</sup> EDS 3<sup>™</sup> EXTERNAL **DRAINAGE SYSTEM**

- Entirely disposable, complete system
- Simple set up
- Laser accuracy
- Needle-free access
- Latex & PVC free



The Codman® EDS 3™ External

Drainage System was developed

with input from more than 200

nurses and physicians.

High-visibility pressure scales in mmHg (-3 to 20) and cmH $_{_2}O$  (-5 to 27)

Drip Chamber with hydrophobic & haemorepellent filter marked for o-100ml capacity, funnel bottom, and sampling site (outlet)

Improved, larger tubing connecting the drip chamber and the drainage bag

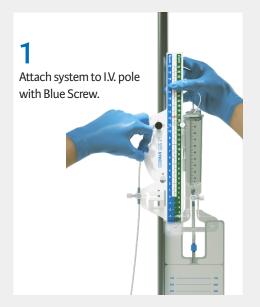
Reusable Leveling Device incorporates bubble level with laser lamp

Pressure-resistant striped CSF tubing line

Redesigned drainage bag with needle-free access port

# CSF EXTERNAL DRAINAGE SYSTEM DESIGNED WITH AND FOR CLINICIANS

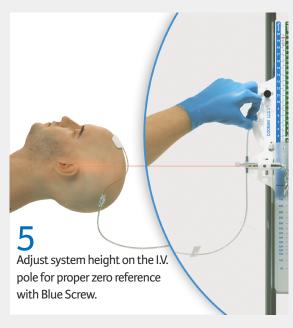
# **Easy Setup**













# Confidence in providing a TOTAL DRAINAGE SOLUTION

## **OUR PLEDGE TO YOU**

Purchase our BACTISEAL® EVD Catheter with the Codman® EDS 3™ System.

If any part of the system requires a revision due to infection or component failure,

Codman Specialty Surgical will replace those components free of charge.

## BACTISEAL EVD + EDS 3 = CONFIDENCE

| PART NO.           | DESCRIPTION  |
|--------------------|--|
|                    | 1.5mm Inner Lumen Diameter   |
| 821745             | - 0 0 0 0 4  |
|                    | 1.9mm Inner Lumen Diameter   |
| 821749             | - 0 0 0 0 4  |
|                    | 1.9mm Clear Inner Lumen Diameter                                     |
| 821750             |  |
| PART NO.           | DESCRIPTION  |
| 821730C            | Codman® External Drainage System EDS 3™<br>with ventricular catheter |
|                    | CODMANDE . LD . C . FDC TA   |
| 821731C            | CODMAN® External Drainage System EDS 3™<br>No Ventricular Catheter   |
| 821731C<br>821732C | <b>5</b> ,   |

#### Indications

The Codman® BACTISEAL® EVD Catheter and Codman® BACTISEAL® Clear EVD Catheter Sets (BACTISEAL® EVD Catheters), are indicated for gaining access to the ventricles of the brain and can be used with dimensionally compatible devices for draining cerebrospinal fluid (CSF) and other fluids of similar physical characteristics as a means of reducing intracranial pressure and CSF volume.

Use of the Codman® EDS 3™ CSF External Drainage System (EDS 3) is indicated for draining cerebrospinal fluid (CSF) and other fluids of similar physical characteristics as a means of reducing intracranial pressure and CSF volume. 1–4

#### Contraindications

Codman® EDS 3™: Use of this device is contraindicated if scalp infection is present. Use of this device is contraindicated for patients receiving anticoagulants or for patients who are known to have a bleeding diathesis. Use of this device is contraindicated where 24-hour supervision from trained personnel is not available.

Bactiseal®: Do not implant this device in patients with active infections, such as ventriculitis meningitis, or skin infections at or near the implantation site. Treat the infection before implanting this device. Do not implant this device in patients with known hypersensitivity to rifampin or clindamycin hydrochloride. Use of this device is contraindicated in patients receiving anticoagulants or known to have a bleeding diasthesis.

- 1. Ingraham FD and Campbell JB: An apparatus for closed drainage of the ventricular system. Ann Surg 114: 1096–1098, 1941
- 2. Bering EA: A simplified apparatus for constant ventricular drainage. J Neurosurg 8: 450–452, 1951.
- 2. Detroit Dev. A simplifined applications to constant in eventual in claimage. I recursing 0. 450 4,251.351.

  3. White RJ, Dakters J, Voung H, et al: continuous control of CSF volume and pressure with an externalized valve-drainage system. Trans Am Soc Artif Intern Organs 13: 332–333, 1967.
- 4. White RK, Dakters JG, et al: Temporary control of cerebrospinal fluid volume and pressure by means of an externalized valve-drainage system. J Neurosurg 30: 264–269, 1969.

Availability of these products might vary from a given country or region to another, as a result of specific local regulatory approval or clearance requirements for sale in such country or region.

- Non contractual document. The manufacturer reserves the right, without prior notice, to modify the products in order to improve their quality.
- Warning: Applicable laws restrict these products to sale by or on the order of a physician.
- Consult product labels and inserts for any indication, contraindications, hazards, warnings, precautions, and instructions for use.

Products mentioned in this document are CE risk class I and III devices. Please contact Integra customer service should you need any additional information on devices classification. All the medical devices mentioned on this document are CE marked in accordance with the applicable European laws, unless specifically identified as "NOT CE MARKED".

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