CoreTainer®

Stereotactic Core Biopsy Image and Transport System

With **A/B Divider** for Basket Collection

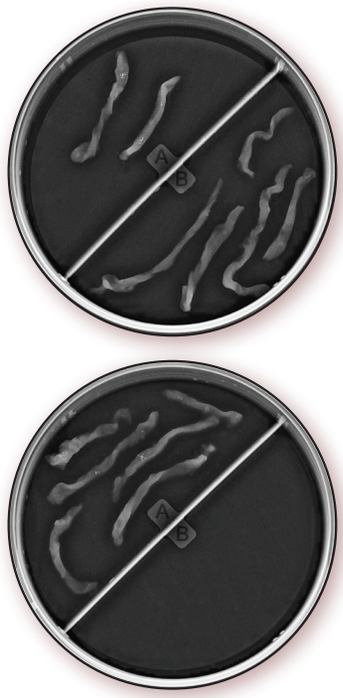
Protect Tissue Integrity: All-in-one system eliminates excessive handling of delicate core specimens by imaging and transporting in the same container.

Transport Safely: Accommodates formalin and is safe for transport to pathology with its watertight lid.

Image Clearly: Helps to eliminate potentially obscuring artifact caused by other makeshift methods. Micro-calcifications are clearly visible on the image.

Enhance Communication: Provides a section specifically for patient information, helping to reduce potential confusion by keeping cores and patient information together.

Save Time: Divided A/B compartments and radiolucent letters quickly allow correlation of core location to the image.



REF **303** QTY **12 / Box**

- All-in-one system for separating, imaging, and transporting stereotactic core needle biopsy specimens with minimal handling
- Two separate compartments labeled A and B
- 3.5″ diameter

Instructions for Use:

1. Use one CoreTainer® with A/B Divider for Basket Collection per biopsy site.
2. Complete patient information on CoreTainer label.
3. Place specimens into A and/or B compartments as desired.
4. Take image of specimens in CoreTainer*.
5. Indicate use of A and/or B compartments for pathology.
6. Lightly moisten specimens with saline or fix with formalin.
7. Seal lid on tightly before transporting CoreTainer to pathology.

*CoreTainer is designed for use with digital equipment. If foreign particles are introduced, they may appear during imaging.

Call 1.800.233.5539 • Fax 1.800.735.1234
Visit beekley.com • Email info@beekley.com

Customers outside the U.S. – Contact your local distributor for pricing and product availability.
To locate a distributor call +1.860.583.4700 or email international@beekley.com

Beekley Medical®
when your diagnosis must be right®

Manufactured by Beekley Corporation
One Prestige Lane, Bristol, CT 06010-7454 USA
Tel: 1.800.233.5539 or +1.860.583.4700
Fax: 1.800.735.1234 beekley.com

WARNINGS AND PRECAUTIONS:
- Single Patient Use Only; Do Not Reuse
- Non-Sterile

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If any serious incident occurs in relation to this device, report it to Beekley Medical and to the competent authority of the Member State in which the user and/or patient is established.

CE EC REP
EMERGO EUROPE
Westervoortsedijk 60
6827 AT Arnhem
The Netherlands

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