

**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.

™ 303 CoreTainer

## 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<sup>®</sup> 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®



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



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.

REV: 303-INF-IFU-WEB-0225