

CoreTainer[®]

Stereotactic Core Biopsy Image and Transport System

With **Clock-Face Divider** for Manual Core 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 clock-face compartments and radiolucent numbers quickly allow correlation of core location to the image.

REF 301 **QTY** 12 / Box

- All-in-one system for separating, imaging, and transporting stereotactic core needle biopsy specimens with minimal handling
- Five separate compartments; one center and four surrounding o'clock compartments
- 3.5" diameter



Instructions for Use:


1. Use one CoreTainer[®] with Clock-Face Divider for Manual Core Collection per biopsy site.
2. Complete patient information on CoreTainer label.
3. Place specimens into labeled compartments as desired.
4. Take image of specimens in CoreTainer*.
5. Lightly moisten specimens with saline or fix with formalin.
6. Seal lid on tightly before transporting CoreTainer to pathology.
7. Transport with communication for pathology on how compartments were utilized.

*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

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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.

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