BEEKLEY
CORPORATION

Declaration of Conformity

This European Declaration of Conformity is issued under the sole responsibility of the manufacturer.

MANUFACTURER		
Name of Company	**Address**	**SRN**
Beekley Corporation	One Prestige Lane Bristol, Connecticut 06010 USA	US-MF-000007055

AUTHORIZED REPRESENTATIVE			
Name of Company	**Address**	**SRN**	**Phone/email**
Emergo Europe	Prinsessegracht 20 2514 AP The Hague The Netherlands	NL-AR-000000116	+31.70.345.8570 EmergoEurope@ul.com

PRODUCT IDENTIFICATION		
Product Name	**Code / Catalog Number**	**Basic UDI-DI**
AccuGrid®	300	00815137021138BD
AccuGrid®	302	00815137020193BE
CoreTainer®	301	00815137021145BA
CoreTainer®	303	00815137022029BE
Intended Purpose	**Photo**	
AccuGrid is a sealed localizing grid system for imaging, transporting, and processing surgical breast specimens.		

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CoreTainer is a sealed, compartmentalized system for imaging and transporting core needle biopsy samples.	

RISK CLASS FOR MEDICAL DEVICES		
Device Classification		**Common Specifications / Standards**
Class:	A	Optional
Rule:	5	

Beekley Corporation declares that the above-mentioned products meet the provision of the following EU legislation:

- In Vitro Diagnostic Regulation (EU) 2017/746

COMPANY REPRESENTATIVE: Kate B. Chase

TITLE: Director of Quality & Regulatory **SIGNATURE:**

PLACE: Bristol, CT 06010 USA **DATE:** 26/01/2022

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