OvertureTi Knee Resurfacing System® Single Use Instruments INSTRUCTIONS FOR USE

DESCRIPTION

The **OvertureTi Knee Resurfacing System Single Use Instruments** are designed to be used in the sizing, spacing, and delivery of **OvertureTi Knee Resurfacing System Implants.** The **OvertureTi Knee Resurfacing System Single Use Instruments** are provided gamma sterilized and intended for single-use.

MATERIALS

Instrument Kit	Material
Femoral Oblong Instrument Kit	**Ultem HU2300, 17-4 PH Stainless Steel Per ASTM F899, 304 Stainless Steel, 420J2 Stainless Steel**
Femoral Round Instrument Kit	**Ultem HU2300, 17-4 PH Stainless Steel Per ASTM F899, 304 Stainless Steel, 420J2 Stainless Steel**
Tibial Instrument Kit	**Ultem HU2300, 304 Stainless Steel, 420J2 Stainless Steel**
Universal Sizing Kit	**Ultem HU2300**

DETAILED DIRECTIONS FOR USE

Step-by-step instructions on how to use the **OvertureTi Knee Resurfacing System Single Use Instruments** are provided in the Overture Orthopaedics' OvertureTi Knee Resurfacing System Single Use Instrument Surgical Technique Guide.

WARNINGS

- The **OvertureTi Knee Resurfacing System Single Use Instruments** should not be used in patients with severe osteoporosis.
- Remove instruments from packaging using sterile technique.

PRECAUTIONS

- Use of the **OvertureTi Knee Resurfacing System Single Use Instruments** should only be undertaken after the surgeon has become thoroughly knowledgeable about knee anatomy and biomechanics; has had experience with partial knee replacement; and has had sufficient training in the use of this device.
- Carefully inspect instruments for damage prior to use.
- If instruments are found to be damaged, contact the manufacturer.

STERILIZATION

All **OvertureTi Knee Resurfacing Single Use Instruments** are provided sterilized. Do not re-sterilize or re-process the instruments. Do not use the instrument kit if packaging appears damaged or open. Do not use the instrument kit if expired.

STORAGE

Sterilized Instrument kits should be stored at temperatures between -30°C-60°C and between 15% and 90% relative humidity.

FURTHER INFORMATION

- The surgical technique contains further information on the **OvertureTi Knee Resurfacing System Single Use Instruments** and may be obtained by contacting Overture Orthopaedics or visiting overtureortho.com
- **OvertureTi Knee Resurfacing System Single Use Instruments** should only be used with **OvertureTi Knee Resurfacing System Implants**.
- Refer to **OvertureTi Knee Resurfacing System Implants IFU** for **OvertureTi Knee Resurfacing System** contraindications, potential adverse events, and warnings.

Manufactured by *Overture Orthopaedics Inc.*
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CAUTION: USA law restricts this device to sale by or on the order of physician.

Symbols Glossary

Symbol No.	Symbol	Description
1		Indicates the manufacturer's catalogue number so that the medical device can be identified.
2		Indicates manufacturer's serial number so that a specific medical device can be identified.
3		Indicates date after which the medical device is not to be used.
4		Indicates medical device manufacturer.
5		Indicates quantity provided
6		Indicates used via prescription only
7		Indicates the medical device that has been sterilized using irradiation.
8		Indicates the medical device that is not to be resterilized.
9		Indicates the medical device is intended for one single use only.
10		Indicates user to consult the instructions for use and the instructions for use are available in an electronic format.
11		Indicates the medical device that should not be used if the package has been damaged or opened.