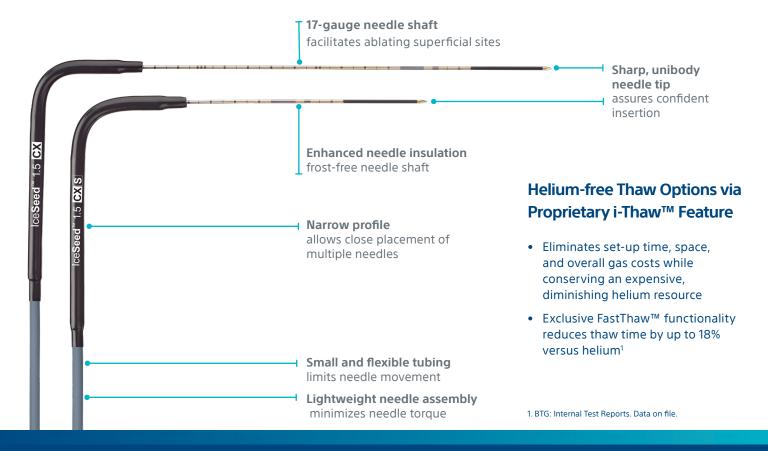




# $egin{aligned} \mathbf{ICESEED}^{^{\mathrm{TM}}} & 1.5 \text{ CX and } \mathbf{ICESEED}^{^{\mathrm{TM}}} & 1.5 \text{ CX Short} \\ & \text{Cryoablation Needles} \end{aligned}$

# WHEN PRECISION MATTERS



### **Precise Control...Refined Therapeutic Choice**



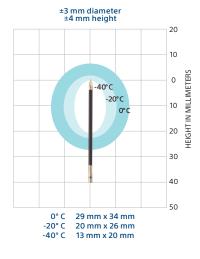
## **ICESEED**<sup>™</sup> 1.5 CX and **ICESEED**<sup>™</sup> 1.5 CX Short Cryoablation Needles

#### IceSeed™ 1.5 CX

IceSeed 1.5 CX Needle: H7493967433170

Shaft Length: 17.5 cm

Shaft Diameter: 1.5 mm / 17G

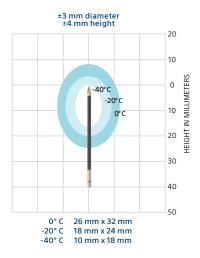


#### IceSeed™ 1.5 CX Short

IceSeed 1.5 CX Short Needle: H7493967233100

Shaft Length:

Shaft Diameter: 1.5 mm / 17G



#### Multi-Needle Isotherm Values (diameter x height in mm)

CRYOABLATION NEEDLES	SINGLE-NEEDLE ISOTHERM			TWO-NEEDLE ISOTHERM			THREE-NEEDLE ISOTHERM			FOUR-NEEDLE ISOTHERM		
	0° C	-20° C	-40° C	0° C	-20° C	-40° C	0° C	-20° C	-40° C	0° C	-20° C	-40° C
IceSeed™ 1.5 CX	29 x 34	20 x 26	13 x 20	41 x 39	31 x 29	23 x 21	48 x 49	38 x 39	29 x 31	52 x 52	42 x 42	33 x 34
IceSeed™ 1.5 CX S	26 x 32	18 x 24	10 x 18	39 x 36	29 x 26	21 x 17	43 x 45	34 x 36	25 x 27	49 x 51	40 x 41	31 x 33

**SKIN PROTECTION:** When performing cryoneurolysis on shallow nerves make sure to place a sterile glove filled with warm saline or warmed ultrasound gel at the surface of the skin. Also, avoid direct contact between skin and needle tubing with the use of towels or draping.

#### **Ordering Information** Cryoablation Portfolio of Needles

CRYOABLATION NEEDLES	PART NUMBER	CONFIGURATION	SHAFT DIAMETER (MM / GAUGE)	SHAFT LENGTH (CM)	HANDLE COLOR	TRACK ABLATION RADIAL WIDTH / LENGTH
IceSeed™ 1.5 CX	H7493967433170	Angled 90°	1.5 mm /17 G	17.5 cm	Black	1.6 / 14 mm
IceSeed™ 1.5 CX S	H7493967233100	Angled 90°	1.5 mm /17 G	10 cm	Black	1.6 / 14 mm

#### ICESEED™ CRYOABLATION NEEDLE

CAUTION: Federal law (USA) restricts this device to sale by or on the order of a physician. By only, Prior to use, please see the complete "Instructions for Use" for more information on Indications. Contraindications Warnings, Precautions, Adverse Events, and Operator's Instructions. INTENDED USE/INDICATIONS FOR USE: The Boston Scientific disposable (xelPod Cryoablation Needle, (xeSphere Cryoablation Needle, and (xeFore Cryoablation Needle, and (xeFore Cryoablation Needle), and (xeFore Cryoablation Needle), are meant to be connected to a Boston Scientific (Efx Cryoablation System or Visual-ICE Cryoablation System when performing cryoablative tissue destruction through application of extremely cold temperatures. The needles are intended to convert high-pressure gas to either a very cold Freezing application or to a warm Thawing application. The Boston Scientific (IcSposable LedeRod Cryoablation Needle, (eeSperce Cryoablation Needle, (eeSperce Cryoablation Needle, (eaSperce Cryoabl thorough understanding of the technical principles, clinical applications, and risk associated with cryoablation is necessary before using this product. Refer to the ICEfx Cryoablation System or Visual-ICE Cryoablation System User Manuals for optional education. \* Do not use this device for any purpose other than the stated intended use. \* Observe the expiration date of this product. Do not use past the listed expiration date. \* Select CX Needles appropriate for the application and tumor size. The iceball shape and size for CX Needles are described in the Device Description Section. \* Do not use CX Needles within a Magnetic Resonance Imaging (MRI) suite or environment. \* Use special care to ensure that a cryoablation needle does not come into contact with an implanted device. \* Cryoablation needles have sharp tips. Use care to ensure safe handling of a cryoablation needle to eliminate the risk of injury or possible exposure to blood-borne pathogens. \* The sterile field and sterility of the cryoablation needle should be maintained at all times. Do not contaminate the distal end of the sterile cryoablation needle. • CX Needles are designed and indicated for Freezing and Thawing applications. These needles are not designed, indicated or tested for thermal protection. • Inspect the packaging for damage. Do not use the needle if packaging appears opened or damaged; in the event of such occurrence, contact Boston Scientific Technical Assistance Center to arrange return of the complete package with the product. • Do not use the needle if it is bent or damaged while attempting to unpack or use it. Never use a defective needle for a recriminal Assistance center to arrange return of the compiete package with me product. • Up not use the needle if it is bent or alrange to while attempting to unpack or use it. never use a electrow needle for a croablation procedure. A defective cryabalition needle that has a gas leak can cause a gas embolism or pneumatosis in the patient. • Avoid bending the needles hard. Do not grass, paid damage to the needle shaft. • During use, avoid damage to the needle from other surgical instruments. • Do not kink, inch, cut or pull excessively on the needle tubing. Damage to needle handle or tubing may cause be needle to become unusable • Feature appropriate stability of thing to avoid inadvertent movement of tubing or needle shifting during a procedure. • Each needle imusable blocked into a needle channel before initiating a cryabalation procedure to avoid the risk of forceful ejection of the needles while under gas pressure. • Prior to starting a cryabalation procedure, set up the cryabalation system (reference the SVSTEM SETUP Section within the cryabalation of unique the receive phase while performing the Needle Integrity and Functionality Test. This isstemus the successfully completed in order to begin the procedure. • Do not use the needle if there is no reformation during the freezee phase while performing the Needle Integrity and Functionality Test. • Refer to cryabalation system user manual for troubleshooting, if the issue deep onter solve, obtain a new needle and repeat the testing procedure. • Do not use the needle if bubbles are seen escaping from the needle during Needle Integrity and Functionality Test. • Finsure adequate measures are taken to more activatives a distinct to the transferate times. • Commission undersome the order of the procedure of the protect organs and structures adjacent to the targeted tissue. • Continuously monitor needle insertion, needle insertion, needle positioning, it when the continuously monitor needle insertion, needle positioning, it when the continuously monitor needle insertion, needle positioning, it when the continuously monitor needle insertion, needle positioning, it when the continuously monitor needle insertion, needle positioning, it when the continuously monitor needle insertion, needle positioning, it when the continuously monitor needle insertion, needle positioning, it when the continuously monitor needle insertion, needle positioning, it when the continuously monitor needle insertion, needle positioning, it when the continuously monitor needle insertion, needle positioning, it when the continuously monitor needle insertion, needle positioning, it when the continuously monitor needle insertion, it was not a continuously monitor needle insertion, needle positioning, it when the continuously monitor needle insertion, needle positioning, it when the continuously monitor needle insertion, it was not a continuously monitor needle insertion, needle positioning, it when the continuously monitor needle insertion, it was not a continuously monitor needle insertion, it was not a continuously monitor needle positioning, it was not a conti from the patient's body and report such event to Boston Scientific. • If a needle unintentionally strikes bone, do not start or continue the Freezing process as needle integrity may be compromised. Replace the needle prior to continuing the procedure. • Needle handles and the needle shaft may frost during freezing. Avoid prolonged contact with frosted portions of the needle handle to avoid unintended thermal fissue damage to the patient or clinician. The patient's skin surface should be protected by warm saline irrigation or other means as determined by the physician. • Needle tubing may become extremely cold when conducting freeze cycles during a cryoablation procedure. It is important that a patient's skin is protected from direct contact with needle tubing to avoid the potential for thermal injury to the patient. Ensure an appropriate insulating barrier is placed as needed (such as towels) or other method is employed to prevent needle tubing from touching a patient's skin. • Ensure the Active Zone Indicator is not positioned outside the patient's skin when track ablation Cautery function) is activated. • The needle handle may become warm during active thawing. Prolonged contact with warm portions of the needle handle could cause unintended thermal tissue damage? burnt to the patient or clinician. • Active thawing produces heat along the distal needle shaft. Use care to avoid thermal injury/burn to nontargeted tissues. • Ensure adequate thawing or cooling before attempting to remove needles from the patient. • Discontinue all needle operation prior to needle removal to minimize risk of thermal injury and/or tissue injury. • When conducting FastThaw Function or track ablation (Cautery function), be alert for the Active Zone Indicator as when track ablation is activated and when the needle is withdrawn to prevent unintended tissue damage from the hot needle . • Nemove needles from the patient prior to disconnecting needles from the Boston Scientific cryoablation system. • Dispose of needle(s) in accordance with the Disposal Section. • No data regarding cryoablation in combination with other therapies is available from Boston Scientific. PRECAUTIONS: General • Boston Scientific CX Needles operate only with a Boston Scientific (Efx Cryoablation System or Visual-ICE Cryoablation System. • Boston Scientific recommends that only needles of the same type be placed together in a single channel. Using needles of differing types in a channel may affect the accuracy of the Gas Indicator. • Confirm availability of sufficient gas to conduct the planned procedure: the number of needles, needle operations activated, gas cylinder size, pressure and gas flow affect the required as a volume. Handling • Use of multiple needles is recommended to fully cover a target sire and provide a suitable margin. • Multiple needles placed in an adjacent configuration will typically create a large, coalesced iceball. Iceball formation must be monitored using image guidance to optimize a successful ablation procedure. • Availability of a back-up needle is recommended should a replacement or additional needle be required during a procedure. • If the Boston Scientific cryoablation system contains pressurized helium, track ablation (Cautery function), i-Thaw Function and FastThaw Function cannot be activated. Procedural • In a safe and controlled fashion, twist the protective sleeve in a rotational motion while simultaneously pulling it away from the device handle. Take extra care when removing the protective sleeve from the device to avoid contact with the distal end of the needle. ADVERSE EVENTS: The potential adverse events associated with the device and/or cryoablation procedure include, but are not limited to: • Allergic reaction (contrast, device, other) • Angina • Arrhythmia • Atelectasis • Bladder spasms • Bleeding/hemorrhage • Burn/frostbite • Cerebrovascular accident (CVA)/stroke • Cryoshock phenomenon (e.g. multi organ failure, severe coagulopathy, disseminated intravascular coagulation (DIC)) • Death • Distension • Edema/swelling • Ejaculatory dysfunction • Embolism (air, device, thrombus) • Feetcille dysfunction • Fewer • Fistula • Fracture • Gastrointestinal symptoms (e.g., nausea, vomitting, diarrhea, constipation) • Healing, impaired • Hematoma • Hematuria • Hemothorax • Hepatic dysfunction / failure • Hernia • Hypoternison • Hypoternison • Hypoternia • News • Precipital • News • News • Precipital • Parenchymal or capsule fracture • Respiratory distress/insufficiency/failure • Scrotaledema • Stenois/stricture • Subciss/stricture • Subciss/stri



#### Peripheral Interventions

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