

## ENGLISH

**BEFORE USING THE DEVICE, PLEASE READ THE FOLLOWING INSTRUCTIONS COMPLETELY AND CAREFULLY. CORRECT APPLICATION IS VITAL TO THE PROPER FUNCTIONING OF THE DEVICE.**

**IF YOU ARE USING THE CRYO/CUFF WITH THE AIRCAST CRYO/CUFF IC UNIT BE SURE YOU READ AND UNDERSTAND ALL INSTRUCTIONS LOCATED IN THE AIRCAST CRYO/CUFF IC INSTRUCTIONS FOR USE.**

**INTENDED USER PROFILE:** The intended user should be a licensed medical professional, the patient, the patient's caregiver or a family member providing assistance. Users should be able to:  
• Read, understand and be physically capable to perform all the directions, warnings and cautions provided in the information for use.

**OPERATING PRINCIPLE:** The AirCast Cryo/Cuff provides cold therapy and compression.

**INTENDED USE/INDICATIONS:** The Aircast Cryo/Cuff combines focal compression with cold to minimize swelling and pain.

**FREQUENCY OF USE:** The length of use and frequency of use of the Cryo/Cuff are determined by the health care professional depending on individual patient's needs. This device can be employed to cause serious injury be sure to check skin frequently.

**CONTRAINDICATIONS:** Cryotherapy should not be used on persons with Raynaud's or other vasospastic disease, cold hypersensitivity, decreased skin sensitivity, compromised local circulation, diabetes, stroke, cerebral aneurysm, arterial hypertension, peripheral vascular disease, aortic aneurysm, aortic dissection, peripheral vascular disease causing ischemia or poor local circulation, local tissue infection, cold allergy, cold hypersensitivity, previous cold injury and paronychia cold hemoglobinuria.

Do not use if you are allergic to any of the materials contained within this product.

### WARNINGS AND PRECAUTIONS:

This device can be cold enough to cause serious injury, including tissue damage. You must be able to check your skin condition under the device and if you cannot check your skin condition frequently. People are sensitive to cold in diverse ways and may react differently to cold treatment. Check for increased pain, burning, welts, numbness, tingling, increased redness, discoloration, itching, increased swelling, blisters, irritation or other changes in skin condition under the Cuff or around the treatment area. If you experience any of these conditions, immediately discontinue use of the device and contact your physician.

When using the device, you must occur every 1-2 hours on a routine basis. Caution must be taken by routinely monitoring the patient during use on children, elderly, incapacitated patients, severe cardiovascular diseases, those with decreased skin sensitivity or circulation due to poor nutrition, malnutrition, dehydration, alcohol or narcotic use, patients on medication for blood pressure, or to occur outside the presence of the licensed healthcare professional, her or she must ensure that the patient or patient caretaker understands how such monitoring is to be conducted.

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**NOT MADE WITH NATURAL RUBBER LATEX.**



**NOT FABRICADO CON LÁTEX DE CAUCHO NATURAL.**



**NON REALIZATO CON LATTICE DI GOMMA NATURALE.**



**BEVAT GEEN NATUURLIJK RUBBERLATEX.**



**NEI PRESENTI NEBOL POUŽITÝ PŘÍRODNÝ GUMOVÝ LATEX.**



**IBA PRE JEDNEHO PACIENTA.**



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