











NATRACURE™ ADVANCED COOLING GEL CAP 1 Piece 11.8" x 7.5" x 2.0"	REF 762-00NC1
NATRACURE™ AVANCERET KØLENDE GELHÆTTE 1 Stk. 11,8" x 7,5" x 2,0"	BONNET DE GEL DE REFROIDISSEMENT NATRACURE™ AVANCÉ 1 unité 11,8 po (30 cm) x 7,5 po (19 cm) x 2,0 po (5 cm)
NATRACURE™ AUSGEREIFTE KÜHLGELKAPPE 1 Stück 11,8" x 7,5" x 2,0"	CAPPELLO IN GEL A RAFFREDDAMENTO AVANZATO NATRACURE™ 1 unità 11,8" x 7,5" x 2,0"
NATRACURE™ AVANSERT KJØLEGEHETTE 1 del 30 x 19 x 5 cm	NATRACURE™ CAPUZ DE GEL DE REFRIGERAÇÃO AVANÇADA 1 Peça 11.8" x 7.5" x 2.0"
NATRACURE™ ADVANCED COOLING GEL CAP 1 Piece 11.8" x 7.5" x 2.0"	NATRACURE™ HUVUDBAND MED AVANCERAD KYLANDE GEL 1 st. 11,8" x 7,5" x 2,0"

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NATRACURE™ ADVANCED COOLING GEL CAP

REF 762-00NC1

1 Piece | 11.8" x 7.5" x 2.0"

Designed to help

- Reduce hair loss*
- Soothe throbbing headache and pain during and after chemotherapy

Key features:

- New GEL/ICETM formula for long cold therapy treatments
- Insulated outer fabric helps retain the cold
- Rounded top for a more targeted treatment
- Fits heads 22.5" to 25.5"
- Suitable for freezer or refrigerator

Intended Use(s): For Cold Only: Provide cold therapy to specific body parts to support healing post-injury, surgery, chronic pain, and to help ease the side effects of chemotherapy.

Intended User(s): For Cold Only: For use by individuals with chronic pain, arthritis, recent or acute injuries, post-operative pain or individuals undergoing chemotherapy treatment.

Directions: Place the cap in the refrigerator or freezer, either wrapped in a paper towel or placed inside the accompanying plastic cover. Chill for a minimum of 2 hours. Place cap on the head and use the top hook-and-loop to adjust the fit, and bottom hook-and-loop for ponytails to leave the cap. Keep on for 20 minutes at a time.

Care: Hand wash the outer and inner fabric with soap and water. Air or blot out until completely dry. Use soaps that have less scent and other possible additives to minimize any possible reactions. Dispose of in accordance with local regulations.

Contraindication(s): Individuals using pacemakers or insulin pumps should NEVER use these products. Do not use during pregnancy. Do not apply cold to area of lost sensation. Do not use where there is poor circulation.

Warning(s):

1. FOR COLD USE ONLY!
2. Do not place product on an open wound.
3. If product causes skin irritation, discomfort or poor circulation occurs, discontinue use and notify your physician.
4. If you have diabetes, neuropathy, or poor circulation, consult a medical professional before using.
5. DO NOT STEP ON PACKS! Excessive weight may cause gel pack(s) to leak or rupture.
6. USE OVER CLOTHING OR PLACE FABRIC INTERFACE BETWEEN THE COLD PACK AND THE SKIN, PRIOR TO USING COLD THERAPY. POSSIBLE INTERFACES INCLUDE: A light towel, cloth, sheet/pillowcase (cut to size), 6 layers of paper towel, or Stockinette. Either wrap the interface around the hot/cold pack prior to attaching to the support (if included) or hold the interface on the skin while placing the product on the body.

Caution(s):


1. Consult your physician before using this product.
2. Only use this product for 20 minutes at a time on the affected area.
3. If at any time soreness, rash, or an increase in pain occurs, discontinue use and consult your physician.
4. Only use this product for its intended purposes as described in the instructions for use.
5. Do not use this product while using transdermal drug delivery skin patches, skin creams, balms, liniments or lotions.
6. If removable pack has been torn or has been compromised (and is leaking) discard immediately.
7. Always use a fabric interface.
8. Keep out of reach of children and pets.

Materials: PE/Nylon Loop Velvet, PTFE-coated nylon, Proprietary Gel Formula

UMDNS: 25806

Made in China

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 PolyGel, LLC

1 Danforth Drive, Easton, PA 18045 USA

NATRACURE™ is a brand of PolyGel, LLC

NatraCure.com

* Studies that cite cold therapy for neuropathy pain and hair loss

<https://pubmed.ncbi.nlm.nih.gov/19171552/>

<https://pubmed.ncbi.nlm.nih.gov/32036471/>

<https://www.breastcancer.org/research-news/cold-therapy-helps-hair-skin-nail-effects>

<https://jamanetwork.com/journals/jamadermatology/article-abstract/2711245>

Not Made with Natural Rubber Latex



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ENGLISH



Symbol	Symbol Description (English)	Danish/dansk	French/Français	German/Deutsche	Italian/italiano	Norwegian	Portuguese/portugalski	Spanish/Español	Swedish/Sweco
	Indicates the medical device manufacturer, as defined in EU MDR 2017/745.	Angiver producenten af medicinsk udstyr, som defineret i EU MDR 2017/745.	Indique le fabricant du dispositif médical, tel que défini dans EU MDR 2017/745.	Zeigt den Hersteller von Medizinprodukten gemäß EU-MDR 2017/745 an.	Indica il produttore del dispositivo medico, come definito nell'MDR (dell'UE 2017/745).	Indikerer produsenten av medisinsk utstyr, som defineret i EU MDR 2017/745.	Indica o fabricante do dispositivo médico, conforme definido no MDR da UE 2017/745.	Indica el fabricante del dispositivo médico, tal como se define en EU MDR 2017/745.	Angiver tillverkaren av medicintekniska produkter, enligt definitionen i EU MDR 2017/745.
	Indicates the authorized representative in the European Community.	Angiver den autoriserede repræsentant i Det Europæiske Fællesskab.	Indique le représentant autorisé dans la Communauté européenne.	Gibt den Bevollmächtigten in der Europäischen Gemeinschaft an.	Indica il rappresentante autorizzato nella Comunità europea.	Angir autorisert representant i Det europeiske fellesskap.	Indica o representante autorizado na Comunidade Europeia.	Indica el representante autorizado en la Comunidad Europea.	Angiver den auktoriserade representanten i Europeiska gemenskapen.
	Indicates the date when the medical device was manufactured and the Country where the device was manufactured. The week-letter code is defined in ISO 3166-1 and can be viewed at https://www.iso.org/obp/ui/#search:Selection+Codes .	Angiver datoen, hvor det medicinske udstyr blev fremstillet, og det land, hvor enheden blev fremstillet. Udstyrsweekkoden er defineret i ISO 3166-1 og kan ses på https://www.iso.org/obp/ui/#search:Selection+Codes .	Indique la date de fabrication du dispositif médical et le pays dans lequel le dispositif a été fabriqué. Le code à deux lettres est défini dans la norme ISO 3166-1 et peut être consulté à l'adresse https://www.iso.org/obp/ui/#search:Selection+Codes de pays.	Gibt das Datum an, an dem das Medizinprodukt hergestellt wurde, und das Land, aus dem das Medizinprodukt hergestellt wurde. Der aus zwei Buchstaben bestehende Code ist in ISO 3166-1 definiert und kann unter https://www.iso.org/obp/ui/#search:Selection+Codes nachgesehen werden. Wählen Sie Ländercodes.	Indica la data in cui è stato prodotto il dispositivo medico e il paese in cui è stato prodotto il dispositivo. Il codice di due lettere è definito in ISO 3166-1 e può essere visualizzato su https://www.iso.org/obp/ui/#search:Selection+Codes paese.	Angir datoen da medisinsk utstyr ble produsert og landet der enheten ble produsert. To-bokstavskoden er defineret i ISO 3166-1 og kan vises på https://www.iso.org/obp/ui/#search:Selection+Codes .	Indica a data em que o dispositivo médico foi fabricado e o país onde o dispositivo foi fabricado. O código de duas letras é definido na ISO 3166-1 e pode ser visualizado em https://www.iso.org/obp/ui/#search:Selection+Codes de país.	Indica la fecha en la que se fabricó el dispositivo médico y el país donde se fabricó el dispositivo. El código de dos letras se define en ISO 3166-1 y puede ver en https://www.iso.org/obp/ui/#search:Selection+Codes de país.	Angiver datum da den medicinska utrustningen tillverkades och landet där enheten tillverkades. Koden med två bokstäver definieras i ISO 3166-1 och kan ses på https://www.iso.org/obp/ui/#search:Selection+Codes .
	Indicates the manufacturer's batch code so that the batch or lot can be identified.	Angiver producentens batchkode, så batch eller part kan identificeres.	Indique le code de lot du fabricant afin que le lot ou le lot puisse être identifié.	Gibt den Chargencode des Herstellers an, damit die Charge oder Charge identifiziert werden kann.	Indica il codice batch del produttore in modo che sia possibile identificare il batch o il lot.	Angir produsentens batchkode slik at batchen eller partiet kan identifiseres.	Indica o código do lote do fabricante para que o lote ou lote possa ser identificado.	Indica el código de lote del fabricante para que se pueda identificar el lote o lote.	Angiver tillverkarens batchkod så att partiet eller partiet kan identifieras.
	Indicates the manufacturer's catalog number so that the medical device can be identified.	Angiver producentens katalognummer, så det medicinske udstyr kan identificeres.	Indique le numéro de catalogue du fabricant afin que le dispositif médical puisse être identifié.	Gibt die Katalognummer des Herstellers an, damit das Medizinprodukt identifiziert werden kann.	Indica il numero di catalogo del produttore in modo che il dispositivo o medico possa essere identificato.	Angir produsentens katalognummer slik at medisinsk utstyr kan identifiseres.	Indica o número de catálogo do fabricante para que o dispositivo médico possa ser identificado.	Indica el número de catálogo del fabricante para que se pueda identificar el dispositivo o médico.	Angiver tillverkarens katalognummer så att den medicinska enheten kan identifieras.
	Indicates product's DEHP-Free and Phthalate-Free.	DEHP-fri og ftalati-udstyr.	Sans DEHP et sans phtalates médical.	DEHP-frei und Phthalat-frei.	Senza DEHP e senza ftalati.	Indikerer at produktet er DEHP-fritt og ftalati-fritt.	Indica que o item é um dispositivo médico.	Indica que el artículo es un dispositivo médico.	Anger att produkten är en medicinsk utrustning.
	Indicates the item is a medical device.	Angiver, at emnet er et medicinsk udstyr.	Indique que l'article est un dispositif médical.	Zeigt an, dass es sich bei dem Artikel um ein Medizinprodukt handelt.	Indica che l'articolo è un dispositivo o medico.	Angir at varen er et medisinsk utstyr.	Indica informações de contato do exportador.	Indica información de contacto del exportador.	Anger att produkten är en medicinsk utrustning.
	Indicates Exporter Contact Information.	Indikerer eksportørens kontaktoplysninger.	Indique les informations de contact de l'exporteur.	Zeigt die Kontaktinformationen des Exporteurs an.	Indica le informazioni di contatto dell'esportatore.	Indikerer eksportørens kontaktoplysning.	Indica informações de contato do exportador.	Indica información de contacto del exportador.	Anger att produkten är en medicinsk utrustning.
	Indicates Single user - multiple use.	Angiver Enkeltbruger - flere anvendelser.	Indique - Utilisateur unique - usage multiple.	Zeigt an Einzelbenutzer - Mehrfachnutzung.	Indica utente singolo - uso multiplo.	Indikerer Enkeltbruger - flertal brugere.	Indica Único usuario - uso multiplo.	Indica Usuario único - uso multiple.	Indikerer En användare - flera gånger.
	Hover your camera over the QR Code to view Instructions for Use in 25 languages.	Hold museen over QR-koden for at se brugsanvisningen på 25 sprog.	Passer votre caméra sur le code QR pour afficher les instructions d'utilisation en 25 langues.	Beugein Sie die Kamera über den QR-Code, um die Gebrauchsanweisung in 25 Sprachen anzuzeigen.	Posiziona la fotocamera sul codice QR per visualizzare le istruzioni per l'uso in 25 lingue.	Hold kameraet over QR-koden for å se bruksanvisningen på 25 språk.	Hold kameraet over QR-koden for å se bruksanvisningen på 25 språk.	Pase la cámara sobre el código QR para ver las instrucciones de uso en 25 idiomas.	Håll musekameran över QR-koden för att se bruksanvisningen på 25 språk.
	Indicates Single user - multiple use.	Angiver Enkeltbruger - flere anvendelser.	Indique - Utilisateur unique - usage multiple.	Zeigt an Einzelbenutzer - Mehrfachnutzung.	Indica utente singolo - uso multiplo.	Indikerer Enkeltbruger - flertal brugere.	Indica Único usuario - uso multiplo.	Indica Usuario único - uso multiple.	Indikerer En användare - flera gånger.