NEOSPLINT

DESCRIPTION
SplintArt are simple and practical splints created to immobilize limbs.
The set includes a complete range of splints for the treatment of sprained or fractured limbs, which can be shaped as needed and once applied according to the needs of the case, offer extraordinary rigidity.
Due to their lightness and adaptability, the complete set or the single pieces can be taken everywhere as they take up very little space.
Moreover, being made of neoprene, they maintain body temperature.

COMPOSITION
SplintArt structure in Royal Blue imitation neoprene laminated with MTP mm.5 D90 and black cotton jersey
Cold mouldable rigid aluminium core
Blue and orange Velcro closure and adjustment

USE DESTINATION
Sprained limbs
Fractured limbs

CODE	MEASURE	Code R.D.M.	Code C.N.D.
STE125	Leg	74683/R	M030599
STE123	Arm	74727/R	M030599
STE124	Elbow - Ankle	74731/R	M030599
STE122	Forearm	74732/R	M030599
STE126	Wrist	74737/R	M030599
STE127	Complete Set (bag + 1 piece to measure)	74761/R	M030599

METHOD OF APPLICATION
1.Place the injured limb on the SplintArt
2.Shape the metal core as desired
3.Adjust and close through Velcro (orange Velcro indicates top)
(To return the SplintArt to a straight position, straighten the metal core with your hands)

INDICATIONS
Post-traumatic or post-operative situations of the limbs where there is a need for effective support and/or protection of the skeletal system and muscle bundles.
When using the devices in the presence of skin lesions, protect the lesion with appropriate bandaging; in such situations, do not use the device in direct contact with the lesion.
Sanitary product to be applied under medical supervision.
Radiolucent – Latex Free – Non-Sterile – Single patient.

CONTRAINDICATIONS / WARNINGS
Disorders of the venous, arterial or lymphatic system. If a medical condition is suspected, consult your physician before use. In any case, use for prolonged periods of time only under medical supervision. It is advisable not to apply the product directly in contact with the skin.

PRESERVATION
Store in a dry place, away from sunlight, heat sources and dust.
The product has no expiration date, but the expected life span is 10 years for the materials and components of the device.

CLEANING
-Hand wash in water at 30° C with neutral soap or detergent
-Rinse well
-Do not wring -Let dry horizontally -Do not iron

DISPOSAL
Not dangerous material, assimilable to urban waste

PACKAGING
Primary: Pieces 1 in PVC bag
Secondary: box pieces 5
Disposable packaging and packing.

PRODUCT CONFORMITY/REFERENCE STANDARDS
REGULATION (EU) 2017/745
Class I Medical Device
ISO 9001:2015

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CE

NEOSPLINT

MANUFACTURER

Artsanity S.r.l. – Via della Borsa, 18 – 31033 Castelfranco Veneto (TV) – ITALY

PACKAGE LEAFLET ACCOMPANYING THE PRODUCT

SPLINTART

COMPOSIZIONE
Struttura SplintArt in simil-neoprene Blu Royal accoppiato con MTP mm.5 D90 e jersey di cotone nero
Anima rigida di alluminio modellabile a freddo - Chiusura e regolazione in velcro blu e arancio

COMPOSITION
SplintArt structure in Blue Royal neoprene imitation coupled with MTP 5mm D90 and black cotton jersey. Rigid cold moldable aluminum core - Blue and orange velcro closure and adjustment

DESTINAZIONE D'USO
Arti slogati – arti fratturati

INTENDED USE
Sprained limbs - fractured limbs

ISTRUZIONI PER L'USO
Appoggiare l'arto lesionato sullo SplintArt, Modellare l'anima in metallo come si desidera, Regolare e chiudere attraverso i velcri (il velcro di colore arancio indica la parte superiore) (Per riportare la SplintArt in posizione retta raddrizzare l'anima in metallo con le mani)

INSTRUCTION FOR USE
Rest the injured limb on the SplintArt, Model the metal core as desired, Adjust and close through the velcro straps (the orange velcro indicates the upper part) (To return the SplintArt to a straight position, straighten the metal core with the hands)

AVVERTENZE
Disturbi al sistema venoso, arterioso o linfatico. In caso di sospetta patologia in atto, consultare il proprio medico prima dell'uso. In ogni caso utilizzare per periodi prolungati nel tempo solo sotto controllo medico. Si consiglia di applicare il prodotto non direttamente a contatto con la cute Radiotrasparente - non sterile – monopaziente – Latex Free
"Si raccomanda di segnalare ad ArtsanitySrl e all'autorità Nazionale competente qualsiasi incidente grave verificatosi in relazione al dispositivo "
Prodotto fabbricato e distribuito conformemente al Regolamento (UE) 2017 / 745
Dispositivo Medico di Classe I

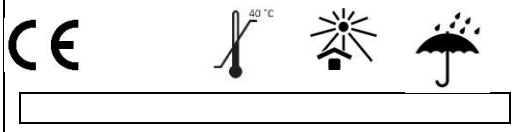
WARNING
Disorders of the venous, arterial or lymphatic system. In case of suspected ongoing disease, consult your doctor before use. In any case, use for prolonged periods of time only under medical supervision. It is recommended to apply the product not directly in contact with the skin. X-ray transparent - non sterile - single patient - Latex Free
"It is advisable to report to Artsanity Srl and the competent national authority any serious accident occurring in relation to the device"
Product manufactured and distributed in accordance with Regulatory (UE) 2017 / 745
Class I Medical Device

CONSERVAZIONE E LAVAGGIO
Conservare in luogo fresco, asciutto e pulito, lontano da luce solare diretta e fonti di calore.
Lavare a mano in acqua a 30°C con sapone o detersivo neutro non sciacquare bene, non strizzare.

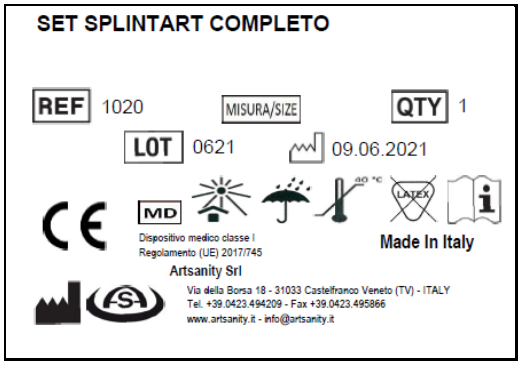
STORAGE AND CLEANING
Store in a cool, dry and clean place away from direct sunlight and heat sources. Hand wash in water at 30 ° C with soap or neutral detergent. Do not rinse well, do not wring

CODICI / *CODE*
REF 1001 Gamba – C.N.D.: M030599 – R.D.M.: 74683/R
REF 1002 Braccio – C.N.D.: M030599 – R.D.M.: 74727/R
REF 1003 Gomito-caviglia – C.N.D.: M030599 – R.D.M.: 74731/R
REF 1004 Avambraccio – C.N.D.: M030599 – R.D.M.: 74732/R
REF 1005 Polso – C.N.D.: M030599 – R.D.M.: 74737/R
REF 1020 Set completo – C.N.D.: M030599 – R.D.M.: 74761/R

Fabbricato e distribuito / *Manufactured and distributed*:



PRIMARY PACKAGING LABEL



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