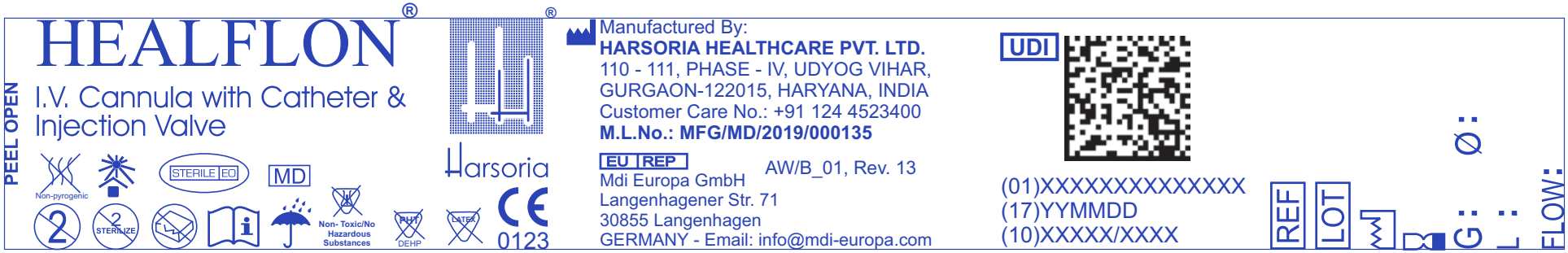
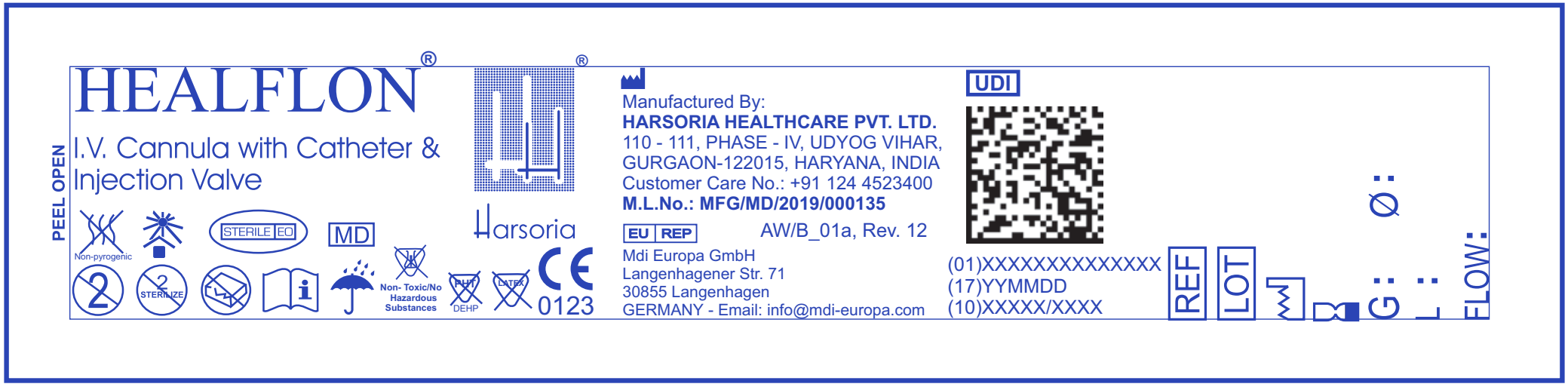
Size= 134X27.5 mm
Printing area=121.6X18.5 mm
Colour of print = Blue (Pantone 2728cvc)
Pitch=29mm, Cavity=6
Pitch=32mm, Cavity=5
Pitch=28.6mm, Cavity=14



FILE NAME/TYPE	HEALFLON (TF_01) / FLON TYPE		**HARSORIA HEALTHCARE PVT. LTD.** ○ CMYK
DESIGNER	BALBEER SINGH		**PANTONE COLORS USED** BLUE 2728
MINIMUM FONT TYPE AND SIZE	Current Size 3.5	Standard Size 4.5	
	Arial	Time New Roman/Arial	
DATE STARTED	12/09/2025		
DATE COMPLETE	01/10/2025		
FILE LOCATION	Z:\CANNULA\FLON TYPE\HEALFLON (TF-01)\B_01, Rev. 13.cdr		**SIZE (LXH) -134X27.5mm**

Size= 114x27.5 mm
Printing area= 104x18 mm
Color of print= Blue (Pantone2728 cvc)
Pitch= 29mm, Cavity=6
Pitch= 32mm, Cavity=5
Pitch= 28.6mm, Cavity=14



FILE NAME/TYPE	HEALFLON (TF_01) / FLON TYPE		HARSORIA HEALTHCARE PVT. LTD. ○ CMYK
DESIGNER	BALBEER SINGH		**PANTONE COLORS USED** BLUE 2728
MINIMUM FONT TYPE AND SIZE	**Current Size** 3	**Standard Size** 4.5	
	Arial	Time New Roman/Arial	
DATE STARTED	12/09/2025		
DATE COMPLETE	01/10/2025		
FILE LOCATION	D:\Design-Artwork\CANNULA\FLON TYPE\HEALFLON (TF-01)\OK\HEALFLON SMOLL (TF_01a)\HEALFLON\B_01a, Rev. 12.cdr		**SIZE (LXH) -114X27.5mm**

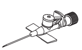
FILE NAME/TYPE	HEALFLON (TF_01) / FLON TYPE		HARSORIA HEALTHCARE PVT. LTD. ○ CMYK
DESIGNER	BALBEER SINGH		PANTONE COLORS USED: BLACK, 2728 CVC
MINIMUM FONT TYPE AND SIZE	Current Size 4.5 Arial	Standard Size 4 Time New Roman/Arial	
DATE STARTED	12/09/2025		
DATE COMPLETE	01/10/2025		
FILE LOCATION	Z:\CANNULA\FLON TYPE\HEALFLON (TF-01)OK\IC_01, Rev.19		SIZE (HxL) -168X136X150 mm

CUTT

Colour of print = Blue (Pantone 2728CVC) & Black
Colour Scheme = As per design
Size = 168X136X150 mm
Print Instruction sheet inside top flap

Without Varnish This area

HEALFLON®
I.V. CANNULA WITH CATHETER & INJECTION VALVE



USE IS RESTRICTED TO A QUALIFIED DOCTOR OR A TRAINED PARAMEDIC

STERILIZE Non-pyrogenic DEHP Non- Toxic/No Hazardous Substances MD STERILE EO

STRIP STICKER

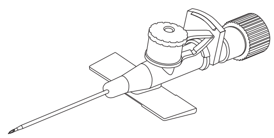
Manufactured By:
HARSORIA HEALTHCARE PVT. LTD.
110 -111, Phase-IV, Udyog Vihar,
Gurgaon - 122 015, Haryana, INDIA.
Email: info@harsoria.com
Website: www.harsoria.com
Customer Care No.: +91 124 4523400
M.L.No.: MFG/MD/2019/000135

EU REP
Mdi Europa GmbH
Langenhagener Str. 71
30855 Langenhagen - GERMANY
Email: info@mdi-europa.com
AW/IC_01,Rev.19

CE 0123

REF
LOT
Manufacturing Date
Expiry Date
UNITS 100
UDI (01)XXXXXXXXXXXXXX (17)YYMMDD (10)XXXXX/XXXX

HEALFLON®



Harsoria

- I.V. Canula
- I.V. Canule
- IV-kanyyli
- I.V. Kanül
- I.V. Kanyle
- Canula EV
- Infusionskanyl
- Kaniula dozylna
- Cathéter court IV
- Venenverweilkanüle
- Cannula Intravenoso
- Cathéter Intraveineux

100 UNITS

Non-pyrogenic DEHP Non- Toxic/No Hazardous Substances MD STERILE EO CE 0123

®
Harsoria

Harsoria

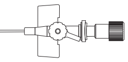
- Sterile unless packaging is damaged. **Caution:** Do not attempt to reinsert a partially or completely withdrawn needle. Take extreme care not to cut catheter. Do not use scissors. non-pyrogenic.
- Steril om förpackningen är oskadad. **Varning:** Återinför aldrig en helt eller delvis utdragen stålkanyl. Var uppmärksam så att plastkatetern ej skärs eller klipps av. Använd ej sax. Pyrogenfri.

CE 0123

REF	LOT	Use before	Do not reuse
Article number.	Lot number:	Use before:	Do not reuse.
Artikelnummer.	Lot nummer.	Används före.	För engångsbruk.
Número de artículo.	Número de Lote.	Utilizar antes.	No reutilizar.
Artikkelnummer.	Produksjonsnummer.	Brukes før.	Kun engangsbruk.
Artikelnummer.	Lot nummer.	Te gebruiken voor.	Niet hergebruiken.
Artikelnummer.	Chargenbezeichnug.	Verwendbar bis.	Nicht zur Wiedervewendung.
Número de artigo.	Número de lote.	Utilizar até.	Não reutilizavél.
Artikelnummer.	Batch nummer.	Anvendes før.	Kun til engangs brug.
Tuotenumero.	Eränumero.	Käytä ennen.	Kertakäyttöön.
Katalog numerasi.	Üretim numarasi.	Son kullanma.	Tekrar kullanilmax.
Référence.	No de lot.	Utiliser avant.	Ne pas réutiliser.
Codice articolo.	Numero di lotto.	Prima dell'uso.	Non riutilizzabile.
Numer katalogowy.	Numer serii.	Data waznosci.	Jednorazowego użytku.

STERILIZE Non-pyrogenic

STERILE EO Non- Toxic/No Hazardous Substances DEHP MD CE 0123



Sterile, Method of sterilization: Ethylene Oxide.
Steril. Steriliseringsmetod: Etylenoxid.
Estéril. Método de esterilización: Oxido de etileno.
Steril. Steriliseringsmetode: Ethylen oxid.
Sterile. Sterilisations methode: Ethyleenoxyde.
Sterile, Sterilisation mit Ethylen Oxid.
Estèril, Método de esterilizção: Oxide etileno.
Steril. Steriliseringsmetode: Ethylen oxid.
Steriili. Sterilointimenetelmä: Etyleenioksidi.
Steril. Ethylene oxide.
Stérile. Méthode de steerilisation: oxyde d'étylène.
Sterile. Metodo di sterilizzazione: Ossido dil etilene.
Jawowy. Metoda sterylizacji: Tlenek etylenu.

Made in India

EU REP
Mdi Europa GmbH
Langenhagener Str. 71
30855 Langenhagen - GERMANY
Email: info@mdi-europa.com

Manufactured By:
HARSORIA HEALTHCARE PVT. LTD.
110 -111, Phase-IV, Udyog Vihar,
Gurgaon - 122 015, Haryana, INDIA.
Email: info@harsoria.com
Website: www.harsoria.com
Customer Care No.: +91 124 4523400

PACKED
CHECKED

TO BE PRINT ON BOTTOM FLAP IN BLACK COLOR

CUTT

BACK SIDE

HEALFLON®

INSTRUCTIONS FOR USE

PACKED
CHECKED

TO BE PRINT ON BOTTOM FLAP IN BLACK COLOR

INSTRUCTIONS FOR USE

HEALFLON®
I.V. Cannula with Catheter & Injection Valve

MATERIAL USED :
•PP, POM, HDPE, LDPE, Silicon Rubber & Stainless Steel
•Catheter: PTFE/FEP/PUR/ETFE (see the product ref. code)

COATING MATERIAL :
•Silicone Dispersion

DESCRIPTION :
Catheter gauge size & length are identified on the product packaging. The color of the catheter's plastic hub alsoindicates the gauge size of catheter.

GAUGE:	14G	16G	18G	20G	22G	24G
COLOR:	Orange	Gray	Green	Pink	Blue	Yellow

The materials used to manufacture this device do not contain natural rubber latex or PVC derivatives.

INTENDED PURPOSE:
"Intravenous Cannula is a device, whereby fluids can be introduced into the human circulatory system during infusion therapy."

INDICATIONS :
•Infusion of I.V. Solutions (To maintain hydration and/or correct dehydration if patient is unable to take sufficient volume of oral fluid)
•Intermittent intravenous Drugs administration

CONTRAINDICATIONS:
•Product should not be used in patients with known hypersensitivity to any of the material used including coating materials
•Product should not be used for Administration of high viscous fluids
•Product should not be used for Large volume blood transfusion
•There is no known reactions between the catheter & Magnetic Resonance Imaging (MRI)

INSTRUCTIONS FOR USE:
•The device is manufactured & tested in accordance with the international standard "Over needle Peripheral Catheters EN ISO 10555-1 & 5"
•The connecting port of this device is in compliance to ISO 80369-7
•Carefully select and aseptically prepare the site
•Select suitable size of I.V. Cannula & inspect visually to ascertain that package is intact
•Remove cannula from sterile packing and remove its needle cover
•Grip the cannula from injection port cap & projection provided on hub
•Perform venipuncture & check for flash back of blood in flash back chamber
•Advance the catheter into vein and simultaneously withdraw the needle
•NEVER TRY TO REINSERT THE PARTIALLY OR COMPLETELY WITHDRAWN NEEDLE
•Withdraw the needle completely while pressing the vein just after the tip of catheter into the vein & discard the needle in an appropriate container
•Connect to the I.V. infusion set line
•Cover the puncture site with sterile dressing
•Drugs can be injected with the help of syringe without needle through integrated injection port after removing the port cap. Close the injection port cap after use
•Perform routine monitoring & venipuncture site maintenance according to medical norms
Based on Clinical Evaluation Report following recommendations are made for use of the device:
•Upper limb placements are preferable to lower limbs
•For blood sampling, it is recommended that larger gauge catheters be used than for infusion
•Use of specialized infusion teams for insertion and monitoring of IV catheters has been shown to lead to better patient outcomes
•In case of any adverse events occurred during the usage of device it must be reported to manufacturer and the national competent authority

INTENDED PATIENT OF USE:
The device can be used in all age group patients.
12G ,13G & 14G to 22G for adults, 24G for child, 26G for neonates.
Note: *•Consult doctor before use of this device in pregnant & Breast feeding women.*
•Consult doctor for appropriate size / type of the device or depending upon the patient's condition.

UNDESIRABLE SIDE EFFECTS MAY BE CAUSED BY USED OF THE DEVICE :
• Allergy • Inflammation • Swelling • Infection

TERM OF USE : •Recommended Maximum Duration of Use: 96 hours

WARNINGS:
•The use of this product is restricted to a qualified doctor or a trained paramedic.
•The device is for Single Use Only. Discard after use.
•Read instructions before use. The product should be used according to the instructions for use.
•Do not put the Device to Use after the Use by Date or Date of Expiry
•If there is any change in expected performance of the device or in case of any malfunction the device should be immediately removed & sent back to supplier for analysis
•**For Known/reported adverse events associated with use of this device, refer to the Clinical Evaluation Report .**
•**HARSORIA HEALTHCARE DISCLAIMS ANY RESPONSIBILITY FOR POSSIBLE CONSEQUENCES RESULTING FROM IMPROPER USE**
•Any device that is connected to this product must comply with ISO 80369-7 in order to achieve the intended performance of this product & to avoid leakage in the connection
•The product should not be reprocessed as reprocessing may compromise the integrity of the device and /or lead to device failure.
•Visually inspect and carefully check the product and packaging before use. Improper transport and handling may cause structural and/or functional damage to device or packaging
•Do not clean or re-sterilize the device as this compromise the device performance (functionality), integrity and biological features. This may increase the risk of cross contamination or severe deterioration in health and safety of patients.
•The product is guaranteed sterile & non-pyrogenic if the package has not been damaged or unintentionally opened before use. If package is damaged or unintentionally opened before use, discard as per country's disposal regulation.
•Do not expose to heat or direct sunlight
•The product should be used immediately after opening the packaging.
•Re-use of this device may change its mechanical or biological features and may cause device failure, allergic reactions or infections

Disposal/ Discard:
•Dispose off/ Discard the used device in accordance with your Country's Healthcare and Safety Regulations.
•Used Sharps/ needles should be disposed in sharps disposal container.
•Used product/ catheter should be disposed in sanitary container or as hospital protocol to prevent possible contamination and cross infection.

Product Ref. No.

Batch Number


Date of Manufacture


Use-by-date


Do Not Reuse

Sterilized using Ethylene Oxide and Single Sterile Barrier System


Caution


Consult Instructions for Use


Non-Pyrogenic


Do Not Use if package is Damaged and consult instructions for use


Do not resterilize


Keep away from sunlight

DEHP Free

Latex Free

Temperature limit

Medical Device

Unique Device Identifier


Keep Dry


Non- Toxic/No Hazardous Substances


Importer


Distributer

Harsoria®

Manufactured By:
Harsoria Healthcare Pvt. Ltd.
110 -111, Phase-IV, Udyog Vihar,
Gurgaon - 122 015, Haryana, INDIA.
Email: info@harsoria.com
Website: www.harsoria.com
Customer Care No.: +91 124 4523400

CE 0123

EU REP
Mdi Europa GmbH
Langenhagener Str. 71
30855 Langenhagen - GERMANY
Email: info@mdi-europa.com

Rev. Date :12.09.2025
Ref. Code: AW/IC_01,Rev.19

FILE NAME/TYPE	HEALFLON (TF_01) / FLON TYPE		HARSORIA HEALTHCARE PVT. LTD. ○ CMYK
DESIGNER	BALBEER SINGH		PANTONE COLORS USED: BLUE 2728, BLACK
MINIMUM FONT TYPE AND SIZE	Current Size 4.5	Standard Size 4	
	Arial	Time New Roman/Arial	
DATE STARTED	12/09/2025		
DATE COMPLETE	01/10/2025		
FILE LOCATION	D:\Design-Artwork\CANNULA\FLON TYPE\HEALFLON (TF-01)\OK\HEALFLON SMOLL (TF_01a)\HEALFLON\IC_01a, Rev. 17.cdr		SIZE (LxWxH) -194X128X122mm

FRONT SIDE

CUTT

Only for 22-26 gauge
Colour of print = Blue (Pantone 2728CVC) & Black
Colour Scheme = As per design
Size = 194X128X122 mm
Print Instruction sheet inside top flap
400GSM Duplex Board

Without Varnish This area

HEALFLON®
I.V. CANNULA WITH CATHETER & INJECTION VALVE

USE IS RESTRICTED TO A QUALIFIED DOCTOR OR A PARAMEDIC

Non-pyrogenic · Non-Toxic/No Hazardous Substances · MD · STERILE EO

STRIP STICKER

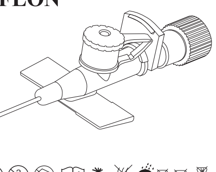
Manufactured By:
HARSORIA HEALTHCARE PVT. LTD.
110 -111, Phase-IV, Udyog Vihar,
Gurgaon - 122 015, Haryana, INDIA.
Email: info@harsoria.com
Website: www.harsoria.com
Customer Care No.: +91 124 4523400
M.L.No.: MFG/MD/2019/000135

EU REP
Mdi Europa GmbH
Langenhagener Str. 71
30855 Langenhagen - GERMANY
Email: info@mdi-europa.com
AW/IC_01a,Rev.17

CE 0123

REF
LOT
Manufacturing Date
Expiry Date
UNITS 100
UDI (01)XXXXXXXXXXXXXX
(17)YYMMDD
(10)XXXXX/XXXX

HEALFLON®



Harsoria®

- I.V. Canula
- I.V. Canule
- IV-kanyyli
- I.V. Kanül
- I.V. Kanyle
- Canula EV
- Infusionskanyl
- Kaniula dozylna
- Cathéter court IV
- Venenverweilkanüle
- Cannula Intravenoso
- Cathéter Intraveineux

100 UNITS

Non-pyrogenic · Non-Toxic/No Hazardous Substances · MD · STERILE EO · CE 0123

Harsoria®

REF	LOT	Use before	Do not reuse
Article number.	Lot number:	Use before:	Do not reuse.
Artikelnummer.	Lot nummer.	Används före.	Fór engångsbruk.
Número de artículo.	Número de Lote.	Utilizar antes.	No reutilizar.
Artikkelnummer.	Produksjonsnummer.	Brukes før.	Kun engangsbruk.
Artikelnummer.	Lot nummer.	Te gebruiken voor.	Niet hergebruiken.
Artikelnummer.	Chargenbezeichnug.	Verwendbar bis.	Nicht zur Wiedervewendung.
Número de artigo.	Número de lote.	Utilizar até.	Não reutilizavél.
Artikelnummer.	Batch nummer.	Anvendes før.	Kun til engangs brug.
Tuotenumero.	Eränumero.	Käytä ennen.	Kertakäyttöön.
Katalog numerasi.	Üretim numarasi.	Son kullanma.	Tekrar kullanilmax.
Référence.	No de lot.	Utiliser avant.	Ne pas réutiliser.
Codice articolo.	Numero di lotto.	Prima dell'uso.	Non riutilizzabile.
Numer katalogowy.	Numer serii.	Data waznosci.	Jednorazowego użytku.

Non-pyrogenic · DEHP

Sterile, Method of sterilization: Ethylene Oxide.
Steril. Steriliseringsmetod: Etylenoxid.
Estéril. Método de esterilización: Oxido de etileno.
Steril. Steriliseringsmetode: Ethylen oxid.
Sterile. Sterilisations methode: Ethyleenoxyde.
Sterile, Sterilisation mit Ethylen Oxid.
Estèril, Método de esterilização: Oxide etileno.
Steril. Steriliseringsmetode: Ethylen oxid.
Steriili. Sterilointimenetelmä: Etyleenioksidi.
Steril. Ethylene oxide.
Stérile. Méthode de steerilisation: oxyde d'étylène.
Sterile. Metodo di sterilizzazione: Ossido dil etilene.
Jawowy. Metoda sterylizacji: Tlenek etylenu.

STERILE EO · Non- Toxic/No Hazardous Substances · MD · CE 0123

Made in India

EU REP
Mdi Europa GmbH
Langenhagener Str. 71
30855 Langenhagen - GERMANY
Email: info@mdi-europa.com

Manufactured By:
HARSORIA HEALTHCARE PVT. LTD.
110 -111, Phase-IV, Udyog Vihar,
Gurgaon - 122 015, Haryana, INDIA.
Email: info@harsoria.com
Website: www.harsoria.com
Customer Care No.: +91 124 4523400

PACKED
CHECKED
TO BE PRINT ON BOTTOM FLAP IN BLACK COLOR

BACK SIDE

CUTT

INSTRUCTIONS FOR USE

PACKED
CHECKED
TO BE PRINT ON BOTTOM FLAP IN BLACK COLOR

INSTRUCTIONS FOR USE

HEALFLON®

I.V. Cannula with Catheter & Injection Valve

MATERIAL USED :
•PP, POM, HDPE, LDPE, Silicon Rubber & Stainless Steel
•Catheter: PTFE/FEP/PUR/ETFE (see the product ref. code)

COATING MATERIAL :
•Silicone Dispersion

DESCRIPTION :
Catheter gauge size & length are identified on the product packaging. The color of the catheter's plastic hub alsoindicates the gauge size of catheter.

GAUGE:	14G	16G	18G	20G	22G	24G
COLOR:	Orange	Gray	Green	Pink	Blue	Yellow

The materials used to manufacture this device do not contain natural rubber latex or PVC derivatives.

INTENDED PURPOSE:
"Intravenous Cannula is a device, whereby fluids can be introduced into the human circulatory system during infusion therapy."

INDICATIONS :
•Infusion of I.V. Solutions (To maintain hydration and/or correct dehydration if patient is unable to take sufficient volume of oral fluid)
•Intermittent intravenous Drugs administration

CONTRAINDICATIONS:
•Product should not be used in patients with known hypersensitivity to any of the material used including coating materials
•Product should not be used for Administration of high viscous fluids
•Product should not be used for Large volume blood transfusion
•There is no known reactions between the catheter & Magnetic Resonance Imaging (MRI)

INSTRUCTIONS FOR USE:
•The device is manufactured & tested in accordance with the international standard "Over needle Peripheral Catheters EN ISO 10555-1 & 5"
•The connecting port of this device is in compliance to ISO 80369-7
•Carefully select and aseptically prepare the site
•Select suitable size of I.V. Cannula & inspect visually to ascertain that package is intact
•Remove cannula from sterile packing and remove its needle cover
•Grip the cannula from injection port cap & projection provided on hub
•Perform venipuncture & check for flash back of blood in flash back chamber
•Advance the catheter into vein and simultaneously withdraw the needle
•NEVER TRY TO REINSERT THE PARTIALLY OR COMPLETELY WITHDRAWN NEEDLE
•Withdraw the needle completely while pressing the vein just after the tip of catheter into the vein & discard the needle in an appropriate container
•Connect to the I.V. infusion set line
•Cover the puncture site with sterile dressing
•Drugs can be injected with the help of syringe without needle through integrated injection port after removing the port cap.
Close the injection port cap after use
•Perform routine monitoring & venipuncture site maintenance according to medical norms
Based on Clinical Evaluation Report following recommendations are made for use of the device:
•Upper limb placements are preferable to lower limbs
•For blood sampling, it is recommended that larger gauge catheters be used than for infusion
•Use of specialized infusion teams for insertion and monitoring of IV catheters has been shown to lead to better patient outcomes
•In case of any adverse events occurred during the usage of device it must be reported to manufacturer and the national competent authority

INTENDED PATIENT OF USE:
The device can be used in all age group patients.
12G ,13G & 14G to 22G for adults, 24G for child, 26G for neonates.
Note: *•Consult doctor before use of this device in pregnant & Breast feeding women.*
•Consult doctor for appropriate size / type of the device or depending upon the patient's condition.

UNDESIRABLE SIDE EFFECTS MAY BE CAUSED BY USED OF THE DEVICE :
• Allergy • Inflammation • Swelling • Infectiion

TERM OF USE : •Recommended Maximum Duration of Use: 96 hours
pital protocol to prevent possible contamination and cross infection.

WARNINGS:
•The use of this product is restricted to a qualified doctor or a trained paramedic.
•The device is for Single Use Only. Discard after use.
•Read instructions before use. The product should be used according to the instructions for use.
•Do not put the Device to Use after the Use by Date or Date of Expiry
•If there is any change in expected performance of the device or in case of any malfunction the device should be immediately removed & sent back to supplier for analysis
•For Known/reported adverse events associated with use of this device, refer to the Clinical Evaluation Report .
•HARSORIA HEALTHCARE DISCLAIMS ANY RESPONSIBILITY FOR POSSIBLE CONSEQUENCES RESULTING FROM IMPROPER USE
•Any device that is connected to this product must comply with ISO 80369-7 in order to achieve the intended performance of this product & to avoid leakage in the connection
•The product should not be reprocessed as reprocessing may compromise the integrity of the device and /or lead to device failure.
•Visually inspect and carefully check the product and packaging before use. Improper transport and handling may cause structural and/or functional damage to device or packaging
•Do not clean or re-sterilize the device as this compromise the device performance (functionality), integrity and biological features. This may increase the risk of cross contamination or severe deterioration in health and safety of patients.
•The product is guaranteed sterile & non-pyrogenic if the package has not been damaged or unintentionally opened before use. If package is damaged or unintentionally opened before use, discard as per country's disposal regulation.
•Do not expose to heat or direct sunlight
•The product should be used immediately after opening the packaging.
•Re-use of this device may change its mechanical or biological features and may cause device failure, allergic reactions or infections

Disposal/ Discard:
•Dispose off/ Discard the used device in accordance with your Country's Healthcare and Safety Regulations.
•Used Sharps/ needles should be disposed in sharps disposal container.
•Used product/ catheter should be disposed in sanitary container or as hos

REF	Product Ref. No.
LOT	Batch Number
	Date of Manufacture
	Use-by-date
	Do Not Reuse
STERILE EO	Sterilized using Ethylene Oxide and Single Sterile Barrier System
	Caution
	Consult Instructions for Use
	Non-Pyrogenic
	Do Not Use if package is Damaged and consult instructions for use
	Do not resterilize
	Keep away from sunlight
PVC DEHP	DEHP Free
LATEX	Latex Free
5°C 30°C	Temperature limit
MD	Medical Device
UDI	Unique Device Identifier
	Keep Dry
	Non- Toxic/No Hazardous Substances
	Importer
	Distributer

Harsoria®

Manufactured By:
Harsoria Healthcare Pvt. Ltd.
110 -111, Phase-IV, Udyog Vihar,
Gurgaon - 122 015, Haryana, INDIA.
Email: info@harsoria.com
Website: www.harsoria.com
Customer Care No.: +91 124 4523400

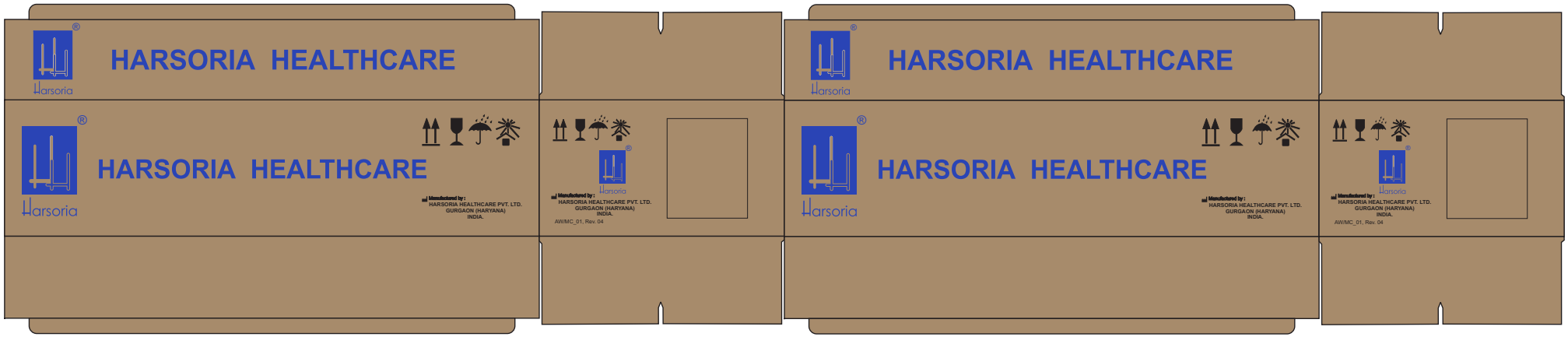
CE 0123
EU REP

Mdi Europa GmbH
Langenhagener Str. 71
30855 Langenhagen - GERMANY
Email: info@mdi-europa.com

Rev. Date :12.06.2025
Ref. Code: AW/IC_01a, Rev. 17

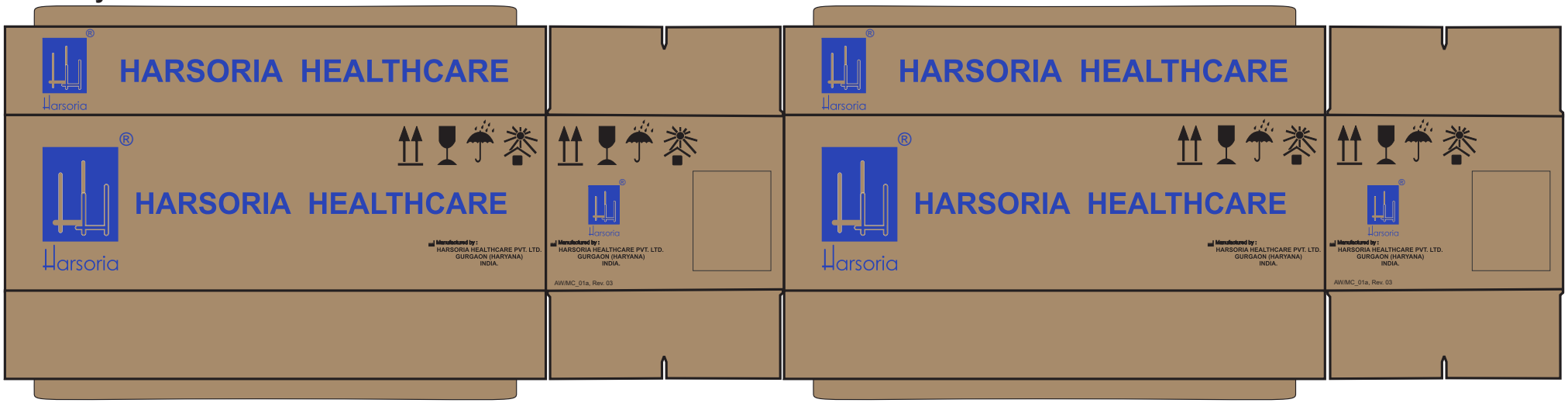
Color of Printing = Blue Pantone 2728cvc & Black
Color scheme = As per design
Base = Brown
Size = 698 mm X 320 X 178 mm
3 PLY, 1 kraft- 180 gsm (top Layer),
2 semi-kraft – 140 gsm

FILE NAME/TYPE	HEALFLON (TF_01) / FLON TYPE		HARSORIA HEALTHCARE PVT. LTD. ○ CMYK
DESIGNER	BALBEER SINGH		
MINIMUM FONT TYPE AND SIZE	Current Size 5.5	Standard Size 5	PANTONE COLORS USED: BLACK, BROWN
	Arial	Time New Roman/Arial	
DATE STARTED	12/09/2025		
DATE COMPLETE	01/10/2025		
FILE LOCATION	Z:\CANNULA\FLON TYPE\HEALFLON (TF-01)\MC_01, Rev. 04.cdr		SIZE (LXWXH) - 698X320X178mm



Color of Printing = Blue Pantone 2728cvc & Black
Color scheme = As per design
Base = Brown
Size = 625 mm X 275 X 204 mm
3 PLY, 1 kraft- 180 gsm (top Layer),
2 semi-kraft – 140 gsm
Only for 22-26 G

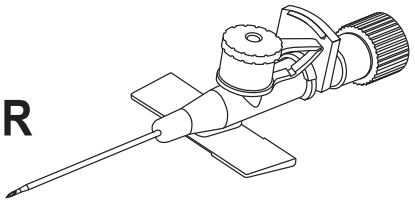
FILE NAME/TYPE	HEALFLON (TF_01) / FLON TYPE		HARSORIA HEALTHCARE PVT. LTD. ○ CMYK
DESIGNER	BALBEER SINGH		
MINIMUM FONT TYPE AND SIZE	Current Size 5.5	Standard Size 5	PANTONE COLORS USED: BLACK, BROWN
	Arial	Time New Roman/Arial	
DATE STARTED	12/09/2025		
DATE COMPLETE	01/10/2025		
FILE LOCATION	Z:\CANNULA\FLON TYPE\HEALFLON (TF-01)\MC_01a, Rev. 03.cdr		SIZE (LXWXH) - 625X275X204mm



Master Carton Sticker
Black print, Size= 100X125 mm

HEALFLON®

I.V. CANNULA WITH CATHETER & INJECTION VALVE



USE IS RESTRICTED TO A QUALIFIED DOCTOR OR A TRAINED PARAMEDIC

Non-pyrogenic | DEHP | LATEX | Non- Toxic/No Hazardous Substances | MD | STERILE EO

STRIP STICKER

Manufactured By:
HARSORIA HEALTHCARE PVT. LTD.
110 -111, Phase-IV, Udyog Vihar,
Gurgaon - 122 015, Haryana, INDIA.
Email: info@harsoria.com
Website: www.harsoria.com
Customer Care No.: +91 124 4523400
M.L.No.: MFG/MD/2019/000135

EU REP
Mdi Europa GmbH
Langenhagener Str. 71
30855 Langenhagen - GERMANY
Email: info@mdi-europa.com
AW/MCS_01, Rev. 16

CE 0123

REF

Manufacturing Date

Expiry Date

LOT

UDI

(01)XXXXXXXXXXXXX
(17)YYMMDD
(10)XXXXX/XXXX

UNITS 1000

Variable Data Change as per Sales Order

FILE NAME/TYPE	HEALFLON (TF_01) / FLON TYPE	HARSORIA HEALTHCARE PVT. LTD.	○ CMYK
DESIGNER	BALBEER SINGH		
MINIMUM FONT TYPE AND SIZE	**Current Size** 5.5 / Arial	**Standard Size** 5 / Time New Roman/Arial	**PANTONE COLORS USED** BLACK
DATE STARTED	12/09/2025		
DATE COMPLETE	01/10/2025		
FILE LOCATION	Z:\CANNULA\FLON TYPE\HEALFLON (TF-01)\MCS_01, Rev. 16.cdr		**SIZE (LXH) -100X125mm**

Actual Product Image