

























### Load syringe and begin:

1. With syringe gradations facing up, push the syringe flange against the orange syringe locator.
2. Make sure the syringe flange is seated within the front part of the orange syringe locator. Place the luer disc inside the syringe driver nose so the syringe is firmly seated inside the syringe driver.
3. Close the top lid to start the infusion. Periodically monitor by checking the progress window, until syringe is empty. **To pause infusion:** Simply open the syringe driver. To continue, re-close the top lid.

### NOTE:

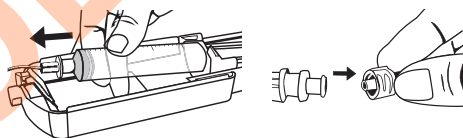
- You should not need to use significant force to load or remove the syringe. You can test proper fit by gently tugging on the syringe. It will stay in place if properly attached.
- When closing the syringe driver, make sure that the top lid is fully extended and aligns with the bottom portion.

**If using multiple syringes:** Once the first syringe is empty, open the FreedomEdge. If instructed, close the clamp on the vascular access device. Remove the syringe from the syringe driver and disconnect from tubing. With aseptic technique, connect the additional syringe to the luer disc end of the Precision tubing set and load back into the syringe driver. If closed, open the clamp on the vascular access device. Close the top lid to continue infusion. Repeat until total dosage is complete.

## 7. End of Infusion

When the syringe is completely empty and the total dosage is infused, open the FreedomEdge and remove the empty syringe.

If instructed, close the clamp on the vascular access device. Disconnect Precision tubing from the on the vascular access device or needle-free connector.



## 8. Flush

Always follow the healthcare provider's instructions on flushing the vascular access device. Refer to the **SASH** technique below.\*

**S** **Saline Flush:** Ensure the vascular access device is open and unobstructed.

**A** **Administer:** Administer the medicinal product.

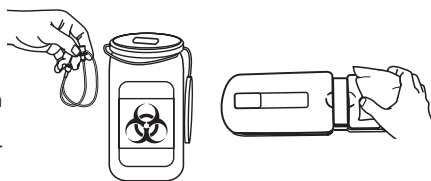
**S** **Saline Flush:** Clear the residual medicinal product from the vascular access device and ensure the vascular access device is open and unobstructed.

**H** **Heparin** (If required for patency): Minimize the potential of a blood clot forming inside the vascular access device.

## 9. Discard Supplies & Clean

Discard all supplies as instructed by your healthcare provider.

Remove visible soil as soon as possible after use of the device. Cleaning should be initiated as soon as possible after use of the device and delays between steps should be avoided. See **page 11** for full cleaning instructions.



## Troubleshooting

If the suggestions in this section do not solve your problem, or if problems persist, discontinue use and consult your healthcare provider.

**NOTE:** Any serious incident should be reported to the local healthcare provider and KORU Medical Systems. Please contact KORU Medical systems at **+1 845-469-2042**.

### Syringe will not load or remove from syringe driver:

- You should not need to use significant force to load or remove a syringe.
- Make sure the syringe driver is fully open, and that nothing is blocking the orange syringe locator.
- Confirm you are not overfilling the syringe (filling a 20 ml syringe with more than 20 ml of solution or a 30 ml syringe with more than 30 ml of solution), or using a syringe larger than 30 ml.
- If you still have difficulty, use one hand to slide the syringe locator all the way back, then place the syringe.

### Syringe will not stay inside in the syringe driver:

- Make sure you are using the proprietary Precision Flow Rate Tubing™ sets and that the luer disc end of the tubing has been connected to a BD® Plastipak™ 20 or 30 ml syringe or a Hizentra® 20 ml prefilled syringe.
- Make sure the luer disc is seated properly in the nose of the syringe driver.
  - **For subcutaneous use:** make sure you have not attached the syringe directly to the HIGh-Flo subcutaneous needle set.
- Make sure the flange shape of the syringe is correctly seated into the shape of the orange syringe locator.

### No flow:

- Open and close the lid to ensure the syringe pusher slides freely and does not bind.
- Make sure all the slide clamps are unclamped. If vascular access device is being used, make sure its clamps, if any, are open.
- Use aseptic technique as recommended by the healthcare provider; disconnect the tubing set from the needle set, vascular access device or needle-free connector, and check for medication drip. If the medication does not drip:
  - **Subcutaneous administration:** replace the tubing as it may be damaged.
  - **Intravenous administration:** check that the catheter is open and unobstructed.

### Slow flow:

- If the slide clamp has been used, the tubing may be damaged.
- Verify that you are using the proper syringe. 30 ml syringes will flow approx. 73% of the rate of a 20 ml syringe.
- Ensure the syringe driver is level with the infusion sites. If the syringe driver is positioned lower than the sites, the flow rate may be slower than expected.
- **Subcutaneous administration:**
  - Administration may be slow based on how well the medicinal product is absorbed through the tissue. Some infusions may be faster than others. The first infusions may take longer than expected because the body may need to adapt.
  - Avoid placing needles on top of scar tissue or muscle.
  - It is possible you may need more sites, longer needles or a faster flow rate tubing set.

### Stopping the flow quickly:

- The syringe driver is designed to maintain pressure during and after the infusion to prevent blood/medicinal product return.

- To stop the flow, fully open the lid to relieve pressure from the syringe plunger.
- The slide clamp can be used in the case of an emergency.

### **Medication (5ml or less) left in the syringe:**

- Verify that you are using a proper recommended syringe: BD® Plastipak™ 20 ml, BD® Plastipak™ 30 ml, or Hizentra® 20 ml Prefilled syringe.
- If the syringe does not completely empty, contact the healthcare provider.

### **Subcutaneous swelling, pain or redness at the site:**

- It is recommended to insert subcutaneous needles dry as the medicinal product may irritate the skin.
- Assure that the needles are long enough to reach the subcutaneous layer. If the selected needle is too short, leaking at the site may occur.
- Assure that the needles are not too long, as they may hit muscle.
- Try a slower flow rate tubing set as the rate may be too fast.
- Rotate infusion sites if recommended by your healthcare provider. Periodically returning to sites that worked well in the past may provide best results.

## **Care, Maintenance and Reprocessing**

The FreedomEdge® syringe driver does not require any preventative maintenance or calibration. The flow rate tubing set determines the flow rate, not the syringe driver; therefore the syringe driver does not need calibration. If you choose the correct tubing set, the proper flow rate will be achieved.

Between uses, the FreedomEdge syringe driver requires to be first thoroughly cleaned, and then disinfected.

After cleaning and disinfection, inspect the device for unacceptable deterioration such as corrosion, discoloration, pitting, and cracked seals and properly dispose any devices that fail the inspection.

### **Cleaning Procedure:**

1. The FreedomEdge syringe driver may be cleaned with a soft cloth dampened with a weak mixture of mild detergent and warm water (minimum ratio of 1 part detergent to 50 parts water by volume).
2. Using the prepared detergent solution and a clean non-linting wipe or soft cloth, wipe all the external surfaces of the syringe driver, including the driver nose and syringe tray up to the syringe shield for at least one (1) minute. During the one (1) minute wipe, pay special attention to the ridges, crevices, raised lettering during wiping. Replace soiled cloths or wipes as needed, changing wipes when necessary to ensure that all surfaces are cleaned.  
**Caution:** Clean only those areas that are exposed and external. No attempt should be made to clean any part of the syringe driver that is not easily accessible.
3. Using a clean non-linting wipe or soft cloth wetted with room temperature tap water (wet but not dripping), wipe all the external surfaces of the syringe driver, including the driver nose and syringe tray up to the syringe shield. Pay special attention to the ridges, crevices, raised lettering during wiping. Continue wiping until all residue is removed to ensure the syringe driver is thoroughly clean. Replace or re-wet cloth or wipes as needed, changing wipes when necessary to ensure that all surfaces are rinsed.
4. Dry device using a clean non-linting wipe or soft cloth.
5. Inspect the device for any visible soil after the cleaning steps (but before the disinfection steps) to ensure that the device is thoroughly cleaned between uses prior to disinfection. If the device has remaining visible soil following cleaning, repeat the cleaning steps (1 through 4).

## Disinfection Procedure:

1. Wipe the outside surfaces of the FreedomEdge syringe driver with 70% Isopropyl Alcohol (IPA) and a non-linting cloth or wipe, or pre-saturated IPA wipe.
2. Use pre-saturated IPA wipes, or non-linting wipes saturated with 70% Isopropyl Alcohol (IPA) (wetted but not dripping) to thoroughly wipe all exterior surfaces of the device. Ensure all external surfaces of the syringe driver, including the driver nose, syringe tray and top lid are wiped. Pay special attention to the ridges and crevices during wiping. Allow all surfaces to remain visibly wet for a minimum of five (5) minutes.

**Caution:** Clean only those areas that are exposed and external. No attempt should be made to clean any part of the syringe driver that is not easily accessible.

3. During the five (5) minute contact time, use additional wipes to ensure all contacted surfaces remain wet for the full contact duration time.
4. Thoroughly dry the device using non-linting wipe(s) or allow to air dry.
5. Visually inspect devices for signs of damage or wear.

## Storage:

The FreedomEdge syringe driver and its components (Precision tubing sets and HlgH-Flo needle sets) are recommended to be stored in a cool, dry place at room temperature (approximately 20–25°C or 68–77°F).

## Technical Specifications

**NOTE:** This section is intended for healthcare providers only.

Testing was performed in a controlled test lab environment and as a result infusions should be administered within the same environmental conditions of 20–25°C (68–77°F) and atmospheric pressure of 1.01 bar ( $\pm 0,09$ ).

### Syringe Driver:

Weight: 0,34 kg (12 oz)  
Length: Closed: 229 mm (9")  
Extended: 299 mm (11,75")  
Width: 83 mm (3,25")  
Height: 38 mm (1,5")

**Syringe:** Reservoir volume: 20/30 ml  
(BD® Plastipak™ 20/30 ml syringes  
or Hizentra® prefilled 20 ml syringes)

**Target Operating Temperature:**  
20–25°C (68–77°F)

### Height Sensitivity:

Vertical Height (cm)	% Variation From Target Flow Rate
$\pm 7,62$ cm from infusion site	Equivalent to Level
$\pm 15,24$ cm from infusion site	up to $\pm 1,2\%$ from target flow rate
$\pm 30,48$ cm from infusion site	up to $\pm 2,4\%$ from target flow rate
$\pm 60,96$ cm from infusion site	up to $\pm 4,8\%$ from target flow rate

### System Max Operating Pressure:

Tubing/Needle Combo	Pressure at the Beginning of Needle Set (psi)	Measured Pressure at End of Needle Set (psi)
<b>F60 + 24G</b>	0,3 psi	0 psi
<b>F2400 + 24G</b>	7,7 psi	0 psi

Data represents pressure changes through the Freedom System (Freedom syringe driver, Precision Flow Rate Tubing™, and HlgH-Flo Subcutaneous Safety Needle Sets™) with the slowest flow rate parameter (F60) and the fastest flow rate parameter (F2400). The net effect: the pressure at the needle is significantly reduced from the initial head pressure.

## Factors that Affect Flow Rate:

It is important to understand that flow rates of infused medicinal products can be affected by multiple factors such as ambient temperature, patient conditions, height differences between the system and infusion site, and variations in solution viscosity.

Using a combination of HlGH-Flo Subcutaneous Safety Needle Sets™ and Precision Flow Rate Tubing™ not specified in the tables on the following pages may result in a flow rate outside of what has been approved for a specific medicinal product.

The total flow rate values presented in the following tables for subcutaneous administration are based on bench testing of combinations of either a 24G or 26G HlGH-Flo needle set connected to a Precision Flow Rate Tubing set. Testing was performed in a controlled test lab with temperatures ranging between 20–25°C (68–77°F).

The infusion times presented in the following table for intravenous administration are approximate. The flow rates shown in the table resulted from testing of distilled water performed in a controlled test lab with temperatures ranging between 20–25°C (68–77°F).

### Testing Flow Accuracy (if required by your local protocol)

1. Remove all air from a new BD® 20 ml syringe with sterile water. Do NOT use a 30 ml syringe for this test.
2. Completely fill with 20 ml of sterile water.
3. Attach a sterile F120 Precision Flow Rate Tubing set to the syringe.
4. Remove all air from the tubing set.
5. Load the syringe into the syringe driver.
6. Using a stop watch or similar time tracking device, start the timer when the top lid of the syringe driver is fully closed (flow will begin).
7. Monitor and stop the timer when 10 ml of water has left the syringe.
8. The elapsed time should fall between 3:50–5:11 minutes.

**NOTE:** If the test results fall outside the range indicated in Step 8, factory refurbishment and testing are available. Please contact your local KORU Medical Systems distributor.

## Ancillary Supply Product Information

### Precision Flow Rate Tubing™ Sets:

Description	Item #	Residual Vol.	p/Box
Very Low Flow	F0.5	0,09 ml	50
Very Low Flow	F1	0,08 ml	50
Very Low Flow	F2	0,10 ml	50
Very Low Flow	F3	0,09 ml	50
Very Low Flow	F3.8	0,09 ml	50
Very Low Flow	F5	0,08 ml	50
Very Low Flow	F8	0,08 ml	50
Very Low Flow	F10	0,14 ml	50
Very Low Flow	F15	0,11 ml	50
Low Flow	F30	0,13 ml	50
Low Flow	F45	0,11 ml	50

Description	Item #	Residual Vol.	p/Box
Low Flow	F60	0,14 ml	50
Low Flow	F120	0,16 ml	50
Low Flow	F180	0,13 ml	50
High Flow	F275	0,11 ml	50
High Flow	F420	0,10 ml	50
High Flow	F500	0,09 ml	50
High Flow	F600	0,09 ml	50
High Flow	F900	0,08 ml	50
High Flow	F1200	0,13 ml	50
High Flow	F2400	0,15 ml	50

## Flow Rate Starter Kits:

Item Number	Description	Contents per Box
H20KT	High Flow Starter Kit	(2) F275, (5) F600, (5) F900, (4) F1200, (4) F2400
L20KT	Low Flow Starter Kit	(2) F30, (5) F45, (5) F60, (4) F120, (4) F180

## KORU Related Accessories:

Item #	Description	Residual Vol.
LRVY	Low Residual Volume Y-Connector	0,14 ml
FEXT	24" Extension Set	0,4 ml

## 26G HlgH-Flo Subcutaneous Safety Needle Sets™:

Single-Needle Sets			
Length	Item #	Residual Vol.	p/ Box
4 mm	RMS12604	0,1 ml	20
6 mm	RMS12606	0,1 ml	20
9 mm	RMS12609	0,1 ml	20
12 mm	RMS12612	0,1 ml	20
14 mm	RMS12614	0,1 ml	20
Three-Needle Sets			
Length	Item #	Residual Vol.	p/ Box
4 mm	RMS32604	0,3 ml	10
6 mm	RMS32606	0,3 ml	10
9 mm	RMS32609	0,3 ml	10
12 mm	RMS32612	0,3 ml	10
14 mm	RMS32614	0,3 ml	10
Five-Needle Sets			
Length	Item #	Residual Vol.	p/ Box
4 mm	RMS52604	0,5 ml	10
6 mm	RMS52606	0,5 ml	10
9 mm	RMS52609	0,5 ml	10
12 mm	RMS52612	0,5 ml	10
14 mm	RMS52614	0,5 ml	10

Two-Needle Sets			
Length	Item #	Residual Vol.	p/ Box
4 mm	RMS22604	0,2 ml	10
6 mm	RMS22606	0,2 ml	10
9 mm	RMS22609	0,2 ml	10
12 mm	RMS22612	0,2 ml	10
14 mm	RMS22614	0,2 ml	10
Four-Needle Sets			
Length	Item #	Residual Vol.	p/ Box
4 mm	RMS42604	0,4 ml	10
6 mm	RMS42606	0,4 ml	10
9 mm	RMS42609	0,4 ml	10
12 mm	RMS42612	0,4 ml	10
14 mm	RMS42614	0,4 ml	10
Six-Needle Sets			
Length	Item #	Residual Vol.	p/ Box
4 mm	RMS62604	0,6 ml	10
6 mm	RMS62606	0,6 ml	10
9 mm	RMS62609	0,6 ml	10
12 mm	RMS62612	0,6 ml	10
14 mm	RMS62614	0,6 ml	10

## 24G HlgH-Flo Subcutaneous Safety Needle Sets™:

Single-Needle Sets			
Length	Item #	Residual Vol.	p/ Box
6 mm	RMS12406	0,4 ml	20
9 mm	RMS12409	0,4 ml	20
12 mm	RMS12412	0,4 ml	20
14 mm	RMS12414	0,4 ml	20
Three-Needle Sets			
Length	Item #	Residual Vol.	p/ Box
6 mm	RMS32406	1,1 ml	10
9 mm	RMS32409	1,1 ml	10
12 mm	RMS32412	1,1 ml	10
14 mm	RMS32414	1,1 ml	10

Two-Needle Sets			
Length	Item #	Residual Vol.	p/ Box
6 mm	RMS22406	0,7 ml	10
9 mm	RMS22409	0,7 ml	10
12 mm	RMS22412	0,7 ml	10
14 mm	RMS22414	0,7 ml	10
Four-Needle Sets			
Length	Item #	Residual Vol.	p/ Box
6 mm	RMS42406	1,4 ml	10
9 mm	RMS42409	1,4 ml	10
12 mm	RMS42412	1,4 ml	10

## Selected Flow Rate Tables

This section is to guide healthcare providers in selecting the Precision Flow Rate Tubing and HlgH-Flo Subcutaneous Safety Needle Sets\* to achieve the desired flow rate based on the selected medicinal product and number of infusion sites.

Infusion parameters (flow rate and volume) are determined based on the medicinal product's prescribing information and the prescriber. The decision on the optimal flow rate tubing and subcutaneous needle configuration (if used) is solely made by the healthcare provider. Patient training by the qualified healthcare provider needs to be completed before starting the self-administration of the prescribed medicinal product.

When using HyQvia®, please refer to prescribing information of the medicinal product for recommended flow rates and to the KORU Precision Flow Rate Controller Instructions for Use.

Note that if using the same tubing set, BD 30 ml syringes will have slightly slower flow rates and slightly increased delivery times versus BD 20 ml syringes (30 ml syringes will flow approx. 73% of the rate of a 20 ml syringe).

Contact the local distributor of KORU Medical Systems for any questions or for further assistance in determining which flow rate tubing and subcutaneous needle set to use.

**NOTE:** All flow rate tables are based on bench top testing which was performed with 0 psi of back pressure.

\*HlgH-Flo Subcutaneous Safety Needle sets are only to be used for subcutaneous administration.

### How to Use Flow Rate Tables for Subcutaneous Administration:

- Select prescribed medicinal product and refer to its prescribing information for recommended infusion flow rate and infusion time.
- Select the subcutaneous needle type – 26G or 24G needle. Verify the correct flow rate table.
- Evaluate and select flow rate tubing and number of needles based on the infusion phase and flow rate.

### Subcutaneous Flow Rate Table Contents:

Aspaveli®/Empaveli® (pegcetacoplan) . . . . .	19
Desferal® (desferrioxamine mesilate) . . . . .	19
Cuvitru® (Immune Globulin Subcutaneous (Human), 20% Solution) . . . .	20
Gammanorm® (Human Normal Immunoglobulin, 165 mg/ml Solution) . .	21
Hizentra® (Immune Globulin Subcutaneous (Human), 20% Liquid) . . . .	22
Hizentra® 20 ml Prefilled Syringe (Immune Globulin Subcutaneous (Human), 20% Liquid) . . . . .	23

## Aspaveli®/Empaveli® (pegcetacoplan) Flow Rate Combinations:

The following table indicates the average, minimum, and maximum predicted infusion times with 26G HlgH-Flo Subcutaneous Safety Needle Sets™ when used in combination with KORU Precision Flow Rate Tubing™ and FreedomEdge® Syringe Infusion System with a 20 ml BD syringe.

Clinical studies were performed using F2400 Precision Flow Rate Tubing with either a 1-leg or 2-leg 26G HlgH-Flo needle set. Typical infusion times were approximately 60 minutes with one infusion site and 30 minutes with two infusion sites. Alternatives may be selected if a longer infusion time is desired.

**Please refer to medicinal product labeling for infusion time and number of needles.**

**HlgH-Flo 26G with Precision Tubing**  
Average (Min-Max) Infusion Time for 20 ml Syringe (Minutes)

	F2400	F1200	F900	F600	F500
<b>1 needle</b>	30 (16-43)	37 (21-52)	38 (22-54)	44 (26-62)	47 (28-65)
<b>2 needles</b>	18 (10-26)	25 (14-36)	27 (16-37)	33 (20-45)	35 (22-48)

Configuration used in clinical studies

## Desferal® (desferrioxamine mesilate) Flow Rate Combinations:

The following tables indicate the nominal predicted infusion times with one (1) 26G HlgH-Flo Subcutaneous Safety Needle Set™ when used in combination with KORU Precision Flow Rate Tubing™ and FreedomEdge® Syringe Infusion System with a 20 ml and 30 ml syringe for the subcutaneous use of Desferal (±15%).

Infusion times are based on the standard subcutaneous concentration per medicinal product labeling. Higher concentrations may result in slower infusions, whereas lower concentrations may result in faster infusions.

**Please refer to medicinal product labeling for maximum indicated flow rate, volume, and infusion time.**

**HlgH-Flo 26G with Precision Tubing - Nominal Infusion Time for 20 ml BD Syringe**

Tubing Set (Rate ml/h)	Syringe Volume (ml)		
	Time for 5 ml	Time for 10 ml	Time for 20 ml
F0.5 (0,60 ml/h)	8 h 18 min	16 h 42 min	33 h 18 min
F1 (1,10 ml/h)	4 h 30 min	9 h 06 min	18 h 12 min
F2 (2,20 ml/h)	2 h 18 min	4 h 30 min	9 h 06 min
F3 (3,20 ml/h)	1 h 36 min	3 h 06 min	6 h 18 min
F3.8 (3,80 ml/h)	1h 18 min	2 h 36 min	5 h 18 min
F5 (5,40 ml/h)	0 h 54 min	1 h 54 min	3 h 42 min

**HlgH-Flo 26G with Precision Tubing - Nominal Infusion Time for 30 ml BD Syringe**

Tubing Set (Rate ml/h)	Syringe Volume (ml)			
	Time for 5 ml	Time for 10 ml	Time for 20 ml	Time for 30 ml
F0.5 (0,50 ml/h)	1 h 00 min	20 h 00 min	40 h 00 min	60 h 00 min
F1 (0,90 ml/h)	5 h 36 min	1 h 06 min	22 h 12 min	33 h 18 min
F2 (1,90 ml/h)	2 h 36 min	5 h 18 min	10 h 30 min	15 h 48 min
F3 (2,70 ml/h)	1 h 54 min	3 h 42 min	7 h 24 min	11 h 06 min
F3.8 (3,20 ml/h)	1 h 36 min	3 h 06 min	6 h 18 min	9 h 24 min
F5 (4,60 ml/h)	1 h 2 min	2 h 12 min	4 h 18 min	6 h 30 min

## Cuvitru® (Immune Globulin Subcutaneous (Human), 20% Solution)

### Flow Rate Combinations:

The following tables indicate the nominal predicted flow rates per site with HlgH-Flo Subcutaneous Safety Needle Sets™ (26G and 24G) when used in combination with KORU Precision Flow Rate Tubing™ and FreedomEdge® Syringe Infusion System with a 20 ml syringe for the subcutaneous use of Cuvitru (±15%).

To determine the appropriate flow rate tubing and subcutaneous needle configuration, please refer to medicinal product labeling for the maximum indicated flow rate and volume per infusion site for initial and subsequent infusions.

To calculate estimated infusion time, please use the formulas below		
Step 1	Flow rate per site, ml/h x Number of needles	Total flow rate, ml/h
Step 2	(Total medicinal product volume, ml / Total flow rate, ml/h) x 60 min = Total infusion time, min	Total infusion time, min

**NOTE:** The infusion is expected to last a maximum of two hours.

### HlgH-Flo **26G** with Precision Tubing - Nominal Flow Rate Per Site (ml/hr/site)

	F120	F180	F275	F420	F500	F600	F900	F1200	F2400
1 needle	8,4	10,4	14,0	18,5	21,0	22,7	27,3	28,6	35,3
2 needles	4,7	6,0	8,5	12,0	14,1	15,7	20,4	21,9	31,0
3 needles	3,2	4,2	6,1	8,8	10,6	12,0	16,3	17,7	27,6
4 needles	2,5	3,2	4,7	7,0	8,5	9,7	13,6	14,9	24,8
5 needles	2,0	2,6	3,9	5,8	7,1	8,1	11,6	12,9	22,6
6 needles	1,7	2,2	3,3	4,9	6,1	7,0	10,2	11,3	20,7

- Flow rates for initial infusion (≤10 ml/hr/site)
- Flow rates for second infusion (≤20 ml/hr/site)
- As per patient's tolerability

### HlgH-Flo **24G** with Precision Tubing - Nominal Flow Rate Per Site (ml/hr/site)

	F120	F180	F275	F420	F500	F600	F900	F1200	F2400
1 needle	10,0	13,1	19,3	28,9	35,5	40,5	57,8	64,1	112,2
2 needles	5,1	6,8	10,1	15,6	19,5	22,5	33,7	38,1	77,6
3 needles	3,4	4,6	6,9	10,7	13,4	15,6	23,8	27,1	59,3
4 needles	2,6	3,4	5,2	8,1	10,2	11,9	18,4	21,0	48,0

- Flow rates for initial infusion (≤10 ml/hr/site)
- Flow rates for second infusion (≤20 ml/hr/site)
- As per patient's tolerability

## Gammanorm® (Human Normal Immunoglobulin, 165 mg/ml Solution)

### Flow Rate Combinations:

The following tables indicate the nominal predicted flow rates per site with HlgH-Flo Subcutaneous Safety Needle Sets™ (26G and 24G) when used in combination with KORU Precision Flow Rate Tubing™ and FreedomEdge® Syringe Infusion System with a 20 ml syringe for the subcutaneous use of Gammanorm (±15%).

To determine the appropriate flow rate tubing and subcutaneous needle configuration, please refer to medicinal product labeling for the maximum indicated flow rate and volume per infusion site for initial and subsequent infusions.

#### HlgH-Flo **26G** with Precision Tubing - Nominal Flow Rate Per Site (ml/hr/site)

	F120	F180	F275	F420	F500	F600	F900	F1200	F2400
1 needle	12,6	16,1	21,5	27,6	30,2	32,9	38,6	42,3	49,4
2 needles	7,1	9,3	13,1	18,0	20,2	22,7	28,6	32,9	42,3
3 needles	4,9	6,5	9,4	13,3	15,2	17,4	22,7	26,9	37,0
4 needles	3,8	5,1	7,4	10,6	12,2	14,0	18,8	22,7	32,9
5 needles	3,1	4,1	6,1	8,8	10,2	11,8	16,1	19,7	29,6
6 needles	2,6	3,5	5,1	7,5	8,7	10,2	14,0	17,4	26,9

- Flow rates for initial infusion (≤15 ml/hr/site)
- Flow rates for second and subsequent infusions (≤25 ml/hr/site)
- Maximum for all sites combined (≤100 ml/hr total)
- Exceeds medicinal product manufacturer's maximum indicated flow rate

#### HlgH-Flo **24G** with Precision Tubing - Nominal Flow Rate Per Site (ml/hr/site)

	F120	F180	F275	F420	F500	F600	F900	F1200	F2400
1 needle	15,2	20,5	30,2	43,8	50,7	58,8	80,2	98,0	146,9
2 needles	7,8	10,6	15,9	23,7	27,7	32,7	46,4	58,8	97,9
3 needles	5,3	7,2	10,8	16,2	19,1	22,6	32,7	42,0	73,4
4 needles	4,0	5,4	8,2	12,3	14,6	17,3	25,2	32,7	58,8

- Flow rates for initial infusion (≤15 ml/hr/site)
- Flow rates for second and subsequent infusions (≤25 ml/hr/site)
- Maximum for all sites combined (≤100 ml/hr total)
- Exceeds medicinal product manufacturer's maximum indicated flow rate

## Hizentra® (Immune Globulin Subcutaneous (Human) 20% Liquid) Flow Rate Combinations:

The following tables indicate the nominal predicted flow rates per site with HiGH-Flo Subcutaneous Safety Needle Sets™ (26G and 24G) when used in combination with KORU Precision Flow Rate Tubing™ and FreedomEdge® Syringe Infusion System with a 20 ml syringe for the subcutaneous use of Hizentra (±15%).

To determine the appropriate flow rate tubing and subcutaneous needle configuration, please refer to medicinal product labeling for the maximum indicated flow rate and volume per infusion site for initial and subsequent infusions.

### HiGH-Flo **26G** with Precision Tubing - Nominal Flow Rate Per Site (ml/hr/site)

	F120	F180	F275	F420	F500	F600	F900	F1200	F2400
1 needle	8,2	10,2	13,7	18,1	20,6	22,2	26,7	28,0	34,6
2 needles	4,6	5,8	8,3	11,7	13,8	15,3	20,0	21,4	30,3
3 needles	3,2	4,1	5,9	8,6	10,4	11,7	16,0	17,4	27,0
4 needles	2,4	3,1	4,6	6,9	8,4	9,5	13,3	14,6	24,3
5 needles	2,0	2,6	3,8	5,7	7,0	8,0	11,4	12,6	22,2
6 needles	1,6	2,2	3,2	4,8	6,0	6,9	9,9	11,1	20,3

Flow rates for initial infusion (≤20 ml/hr/site)

Flow rates for second and third infusions (≤35 ml/hr/site)

### HiGH-Flo **24G** with Precision Tubing - Nominal Flow Rate Per Site (ml/hr/site)

	F120	F180	F275	F420	F500	F600	F900	F1200	F2400
1 needle	9,8	12,8	18,9	28,3	34,8	39,7	56,7	62,8	109,9
2 needles	5,0	6,6	9,9	15,3	19,1	22,0	33,0	37,3	76,0
3 needles	3,4	4,5	6,7	10,4	13,1	15,3	23,3	26,5	58,1
4 needles	2,5	3,4	5,1	7,9	10,0	11,7	18,0	20,6	47,0

Flow rates for initial infusion (≤20 ml/hr/site)

Flow rates for second and third infusions (≤35 ml/hr/site)

Flow rates for fourth and subsequent infusions (per patient's tolerability)

## Hizentra® (Immune Globulin Subcutaneous (Human) 20% Liquid) 20 ml Prefilled Syringe Flow Rate Combinations:

The following tables indicate the average, minimum, and maximum predicted flow rates per site with HiGH-Flo Subcutaneous Safety Needle Sets™ (26G and 24G) when used in combination with KORU Precision Flow Rate Tubing™ and FreedomEdge® Syringe Infusion System with a 20 ml syringe for the subcutaneous use of Hizentra.

**To determine the appropriate flow rate tubing and subcutaneous needle configuration, please refer to medicinal product labeling for the maximum indicated flow rate and volume per infusion site for initial and subsequent infusions.**

### HiGH-Flo 26G with Precision Tubing - Average (Min-Max) Flow Rate Per Site (ml/hr/site)

	F120	F180	F275	F420	F500	F600	F900	F1200	F2400
1 needle	9,9 (7,0-12,9)	12,5 (9,7-15,4)	17,2 (13,1-21,2)	22,7 (16,7-28,7)	25,2 (19,9-30,5)	27,7 (21,6-33,9)	34,6 (27,3-42)	37,3 (28,8-45,9)	45,7 (35,7-55,6)
2 needles	5,4 (3,8-7,1)	7,0 (5,4-8,7)	10,1 (7,6-12,6)	14,2 (10,1-18,2)	16,1 (12,6-19,6)	18,3 (14,0-22,5)	24,8 (19,2-30,3)	27,6 (20,7-34,6)	37,8 (28,7-46,9)
3 needles	3,7 (2,6-4,9)	4,9 (3,7-6,0)	7,2 (5,4-8,9)	10,3 (7,3-13,3)	11,9 (9,3-14,5)	13,6 (10,3-16,9)	19,3 (14,8-23,7)	22,0 (16,2-27,7)	32,3 (24-40,6)
4 needles	2,9 (2,0-3,7)	3,7 (2,9-4,6)	5,5 (4,2-6,9)	8,1 (5,7-10,5)	9,4 (7,3-11,5)	10,9 (8,2-13,5)	15,8 (12,1-19,5)	18,2 (13,3-23,2)	28,2 (20,6-35,8)
5 needles	2,3 (1,6-3,0)	3,0 (2,3-3,8)	4,5 (3,4-5,7)	6,7 (4,7-8,7)	7,8 (6,0-9,5)	9,0 (6,8-11,2)	13,3 (10,2-16,5)	15,6 (11,2-19,9)	25 (18,1-32,0)
6 needles	1,9 (1,3-2,5)	2,5 (1,9-3,2)	3,8 (2,9-4,8)	5,7 (3,9-7,4)	6,6 (5,1-8,1)	7,7 (5,8-9,6)	11,6 (8,8-14,4)	13,6 (9,8-17,4)	22,5 (16,1-28,9)

- Flow rates for initial infusion ( $\leq 20$  ml/hr/site)
- Flow rates for second and third infusions ( $\leq 35$  ml/hr/site)
- Flow rates for fourth and subsequent infusions

### HiGH-Flo 24G with Precision Tubing - Average (Min-Max) Flow Rate Per Site (ml/hr/site)

	F120	F180	F275	F420	F500	F600	F900	F1200	F2400
1 needle	11,5 (8,0-15,1)	15,1 (11,5-18,7)	22,5 (16,8-28,2)	33,1 (23,2-43,0)	38,5 (29,9-47,1)	44,8 (33,8-55,8)	66,0 (50,3-81,7)	76,8 (55,6-98,0)	122,5 (88,8-156,3)
2 needles	5,9 (4,0-7,7)	7,8 (5,9-9,6)	11,7 (8,7-14,7)	17,6 (12,2-23,0)	20,7 (16,0-25,4)	24,4 (18,2-30,5)	37,5 (28,3-46,7)	44,7 (31,7-57,7)	79,1 (55,3-102,8)
3 needles	4,0 (2,7-5,2)	5,2 (4,0-6,5)	7,9 (5,9-10,0)	12,0 (8,3-15,7)	14,2 (11,0-17,4)	16,7 (12,5-21,0)	26,2 (19,7-32,7)	31,5 (22,2-40,9)	58,4 (40,2-76,6)
4 needles	3,0 (2,0-3,9)	3,9 (3,0-4,9)	6,0 (4,5-7,5)	9,1 (6,3-11,9)	10,8 (8,3-13,2)	12,8 (9,5-16,0)	20,1 (15,1-25,2)	24,3 (17-31,7)	46,3 (31,5-61,1)

- Flow rates for initial infusion ( $\leq 20$  ml/hr/site)
- Flow rates for second and third infusions ( $\leq 35$  ml/hr/site)
- Flow rates for fourth and subsequent infusions

## How to Use Flow Rate Table for Intravenous Antibiotic Administration:

- Select prescribed medicinal product and refer to its prescribing information for recommended infusion flow rate and infusion time.
- Verify the expected infusion time and the syringe volume.
- Evaluate and select flow rate tubing based on the expected infusion time and the syringe volume.

## Selected Flow Rates for Intravenous Administration:

The following tables indicate the nominal predicted infusion times when used in combination with KORU Precision Flow Rate Tubing™ and the FreedomEdge® Syringe Driver with a 20 ml and 30 ml syringe for the intravenous use of meropenem, ertapenem, oxacillin, and tobramycin (±15%).

Please refer to medicinal product labeling for recommended infusion rates and times.

20 ml BD Syringe

Tubing Set (Rate ml/h)	Syringe Volume (ml)	
	Infusion time for 10ml	Infusion time for 20 ml
F2 (2,37 ml/h)	4 h 12 min	8 h 24 min
F3 (3,40 ml/h)	2 h 54 min	5 h 54 min
F3.8 (4,10 ml/h)	2 h 24 min	4 h 54 min
F5 (5,83 ml/h)	1 h 42 min	3 h 24 min
F8 (8,63 ml/h)	1 h 12 min	2 h 18 min
F10 (10,79 ml/h)	0 h 54 min	1 h 54 min
F15 (16,19 ml/h)	0 h 36 min	1 h 12 min
F30 (37,77 ml/h)	0 h 18 min	0 h 30 min
F45 (59,25 ml/h)	0 h 12 min	0 h 18 min
F60 (77,71 ml/h)	0 h 6 min	0 h 18 min
F120 (144,14 ml/h)	0 h 6 min	0 h 6 min
F180 (194,27 ml/h)	0 h 6 min	0 h 6 min
F275 (296,90 ml/h)	0 h 0 min	0 h 6 min

30 ml BD Syringe

Tubing Set (Rate ml/h)	Syringe Volume (ml)		
	Infusion time for 10ml	Infusion time for 20 ml	Infusion time for 30 ml
F2 (1,88 ml/h)	5 h 18 min	10 h 36 min	15 h 54 min
F3 (2,70 ml/h)	3 h 42 min	7 h 24 min	11 h 06 min
F3.8 (3,25 ml/h)	3 h 06 min	6 h 06 min	9 h 12 min
F5 (4,62 ml/h)	2 h 12 min	4 h 18 min	6 h 30 min
F8 (6,85 ml/h)	1 h 30 min	2 hr 54 min	4 hr 24 min
F10 (8,56 ml/h)	1 h 12 min	2 hr 18 min	3 hr 30 min
F15 (12,84 ml/h)	0 h 48 min	1 h 36 min	2 h 18 min
F30 (29,97 ml/h)	0 h 18 min	0 h 42 min	1 h 00 min
F45 (47,01 ml/h)	0 h 12 min	0 h 24 min	0 h 36 min
F60 (61,65 ml/h)	0 h 12 min	0 h 18 min	0 h 30 min
F120 (114,36 ml/h)	0 h 6 min	0 h 12 min	0 h 18 min
F180 (154,14 ml/h)	0 h 6 min	0 h 6 min	0 h 12 min
F275 (235,49 ml/h)	0 h 0 min	0 h 6 min	0 h 6 min

## Warranty Information

This warranty and the rights and obligations hereunder, shall be construed under and governed by the laws of the State of New York, USA.

**Limited Warranty:** KORU Medical Systems (“Manufacturer”) warrants the FreedomEdge® syringe driver to be free from defects in materials and workmanship under normal use. Warranty is limited to Original Purchaser and covers the FreedomEdge for a period of two years from the purchase date. This warranty is not valid for any damage caused by the use of non-KORU products. The “Original Purchaser” is the person purchasing the syringe driver from the Manufacturer or Manufacturer’s Representative. Warranty does not extend to subsequent purchasers. Subject to the conditions of and upon compliance with the procedures set forth in this limited warranty, the Manufacturer will repair or replace, at its option, any syringe driver, or part thereof, which has been actually received by the Manufacturer or Manufacturer’s Representative within the two-year warranty period, and which examination discloses, to the Manufacturer’s satisfaction, that the product is defective. Replacement product and parts are warranted only for the remaining portion of the original two-year warranty period.

KORU tests the FreedomEdge syringe driver using KORU accessories to ensure that the FreedomEdge operates in accordance with published specification standards. If non-KORU accessories are used in conjunction with the FreedomEdge, KORU does not represent that the FreedomEdge will operate in accordance with published specification standards. The FreedomEdge warranty does not cover third-party products or accessories.











### The following conditions, procedures, and limitations apply to the Manufacturer’s obligations under this warranty:

- **Parties Covered by this Warranty:** This warranty extends only to the Original Purchaser of the infusion syringe driver. This warranty does not extend to subsequent purchasers.
- **Warranty Performance Procedure:** Notice of the defect must be made in writing to Customer Support Department, KORU Medical Systems, 24 Carpenter Road, Chester, NY 10918, USA. Notice to KORU Medical Systems must include the model and serial number, date of purchase, and description of the defect in sufficient detail to facilitate repairs. Authorization must be obtained by the Original Purchaser from the Manufacturer or Manufacturer’s Representative prior to returning the product to the Manufacturer. The defective syringe driver must be properly packaged and returned to the Manufacturer, postage-prepaid. Any loss or damage during shipment is at the risk of the Original Purchaser.
- **Conditions of Warranty:** This warranty does not apply to any product, or part thereof, which has been repaired or altered outside of the Manufacturer’s facility in a way so as, in Manufacturer’s judgment, to affect its stability or reliability, or which has been subjected to misuse, negligence or accident.

- **Limitations and Exclusions:** Repair or replacement of a syringe driver or component part is the EXCLUSIVE remedy offered by the Manufacturer. The following exclusions and limitations shall apply:
  - No agent, representative, or employee of the Manufacturer has authority to bind the Manufacturer to any representation or warranty, expressed or implied, or to change this limited warranty in any way.
  - THIS LIMITED WARRANTY IS IN LIEU OF ALL OTHER WARRANTIES, EXPRESS OR IMPLIED, INCLUDING, BUT NOT LIMITED TO, WARRANTIES OF MERCHANTABILITY AND FITNESS FOR A PARTICULAR PURPOSE. THERE ARE NO WARRANTIES THAT EXTEND BEYOND THE DESCRIPTION ON THE FACE HEREOF.
  - Manufacturer's liability under this Limited Warranty Agreement shall not extend to special, indirect, or consequential damages.
  - The syringe driver can only be used under the supervision of medical personnel whose skill and judgment determine the suitability of the syringe driver for a particular medical treatment.
  - All recommendations, information, and descriptive literature supplied by the Manufacturer or its agents are believed to be accurate and reliable, but do not constitute warranties.

OBSOLETE

## Definition of Symbols

	Caution		Use by YYYY-MM-DD or YYYY-MM
	Consult Instructions For Use		Manufacturer
<b>EC</b> <b>REP</b>	Authorized Representative in the European Community		Do Not Reuse
<b>CH</b> <b>REP</b>	Swiss Authorized Representative		Do Not Resterilize
<b>LOT</b>	Batch Code		Not Made with Natural Rubber Latex
<b>QTY</b>	Quantity		Do Not Use if Package is Damaged
<b>REF</b>	Catalog Number		MR Unsafe
<b>SN</b>	Serial Number	<b>Rx</b> <b>ONLY</b>	Prescription Only
<b>STERILE</b> <b>R</b>	Sterilized Using Irradiation	<b>CE</b>	European Conformity
<b>MD</b>	Medical Device		Importer



+1 845-469-2042 | [korumedical.com](http://korumedical.com) | [@korumedical](https://twitter.com/korumedical)



Manufacturer  
**KORU Medical Systems**  
24 Carpenter Road  
Chester, NY 10918 USA  
+1 (845) 469 2042



**EC REP** European Representative

**ICON plc**  
95 bis Boulevard Pereire  
75017 Paris, France  
+33 (0)1 42 12 83 30

**CH REP** Swiss Representative

**MedEnvoy Switzerland**  
Gotthardstrasse 28  
6302 Zug  
Switzerland



**MedEnvoy**  
Prinses Margrietplantsoen 33  
Suite 123  
2595 AM The Hague  
The Netherlands



**MedEnvoy Switzerland**  
Gotthardstrasse 28  
6302 Zug  
Switzerland

The FreedomEdge® Syringe Infusion System, Precision Flow Rate Tubing™, HiGH-Flo Subcutaneous Safety Needle Sets™, and DynEQ® are trademarks of KORU Medical Systems and are compliant with Medical Device Directive 93/42/EEC. All other trademarks are the property of their respective owners. KORU Medical Systems is ISO 13485 certified. ©2022 KORU Medical Systems; All Rights Reserved.