

Baxter

ПОДТВЕРЖДАЮ

Руководитель отдела регистрации,
Россия и СНГ
АО Компания «Бакстер»
Гусев М.В.



2021г.

**ФОТОГРАФИЧЕСКИЕ ИЗОБРАЖЕНИЯ МЕДИЦИНСКОГО
ИЗДЕЛИЯ**

**«Помпа медицинская инфузионная эластомерная
одноразовая стерильная Folfusor в вариантах исполнения»**

производства компании
«Бакстер Хелскеа СА» (Baxter Healthcare SA), Швейцария

**1. Помпа медицинская инфузионная эластомерная одноразовая стерильная
Foliusor SV 0,5 мл/ч**



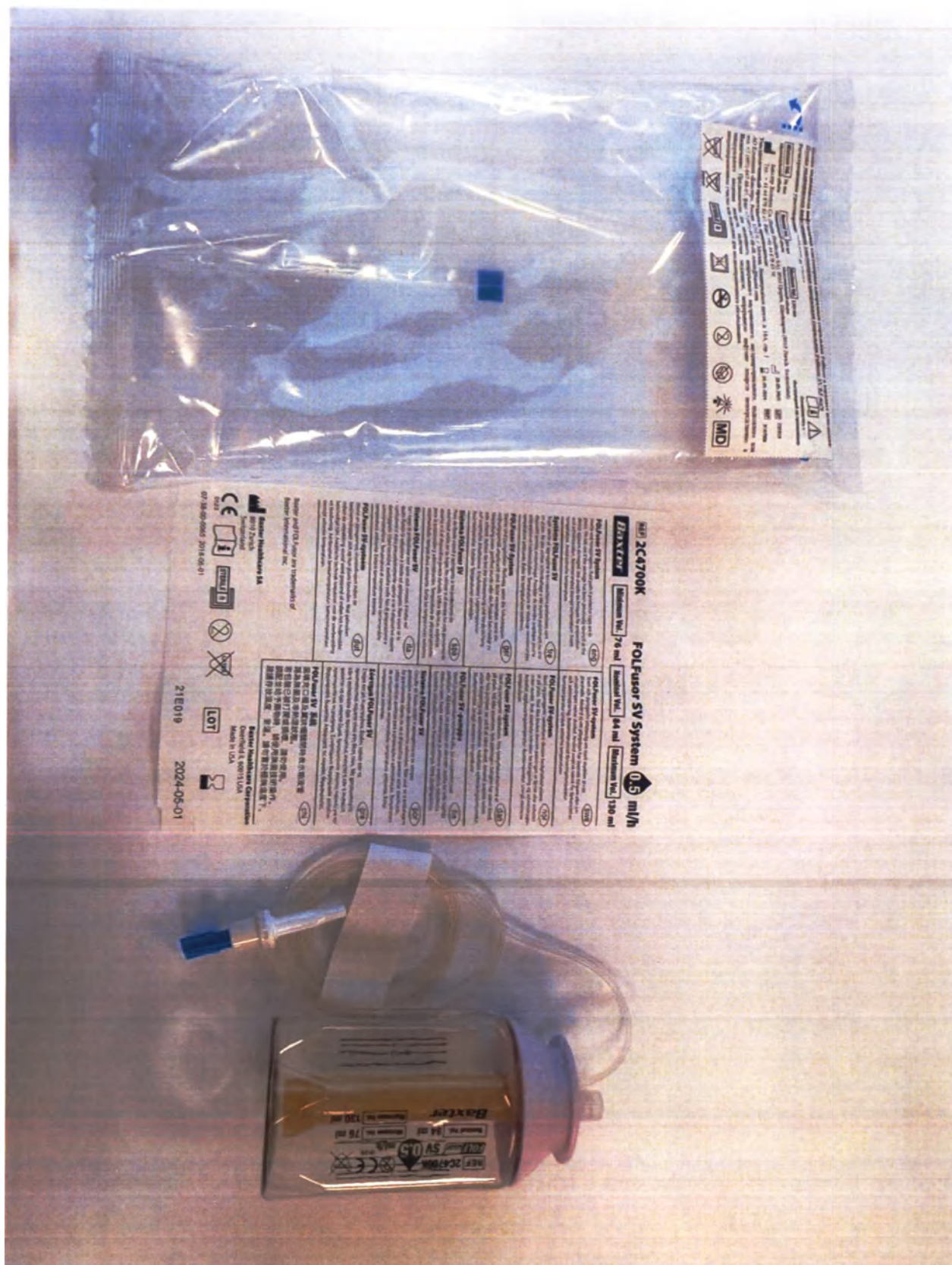
1.1. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor SV 0,5 мл/ч; Индивидуальная упаковка, лицевой сторона, с оригинальной маркировкой



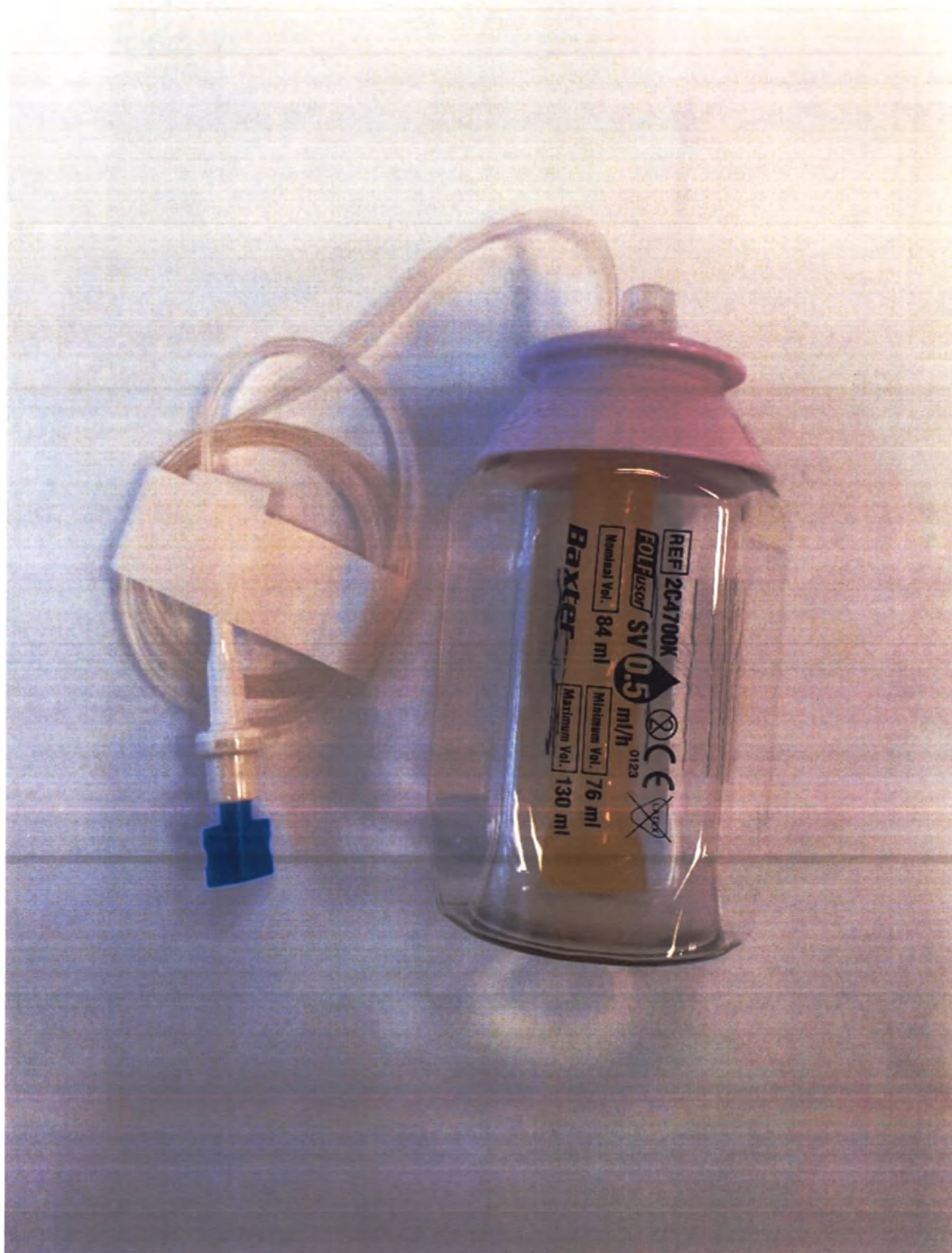
1.2. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor SV 0,5 мл/ч; Индивидуальная упаковка, оборотная сторона



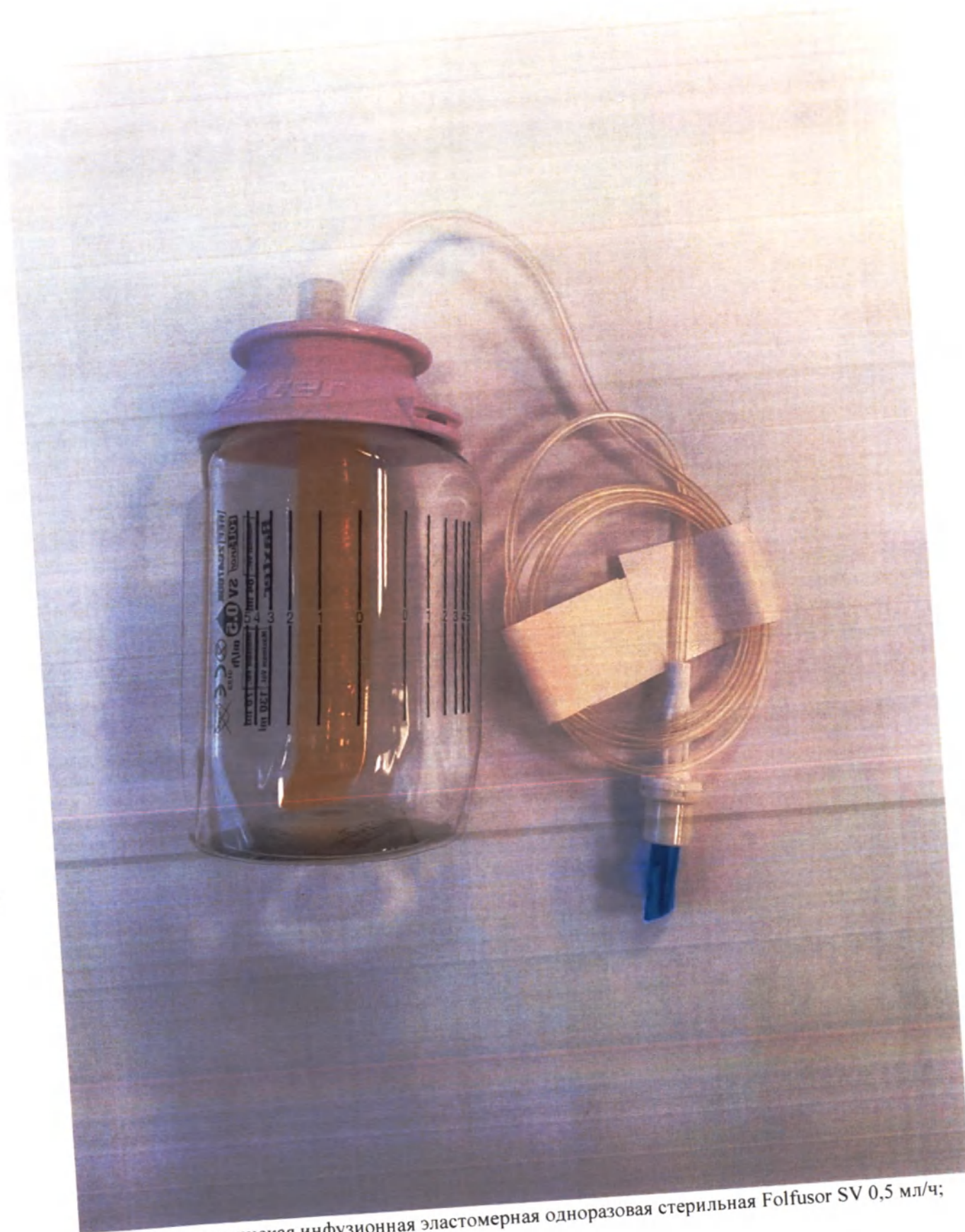
1.3. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor SV 0,5 мл/ч; Индивидуальная упаковка, лицевая сторона, с нанесенной маркировкой на русском языке



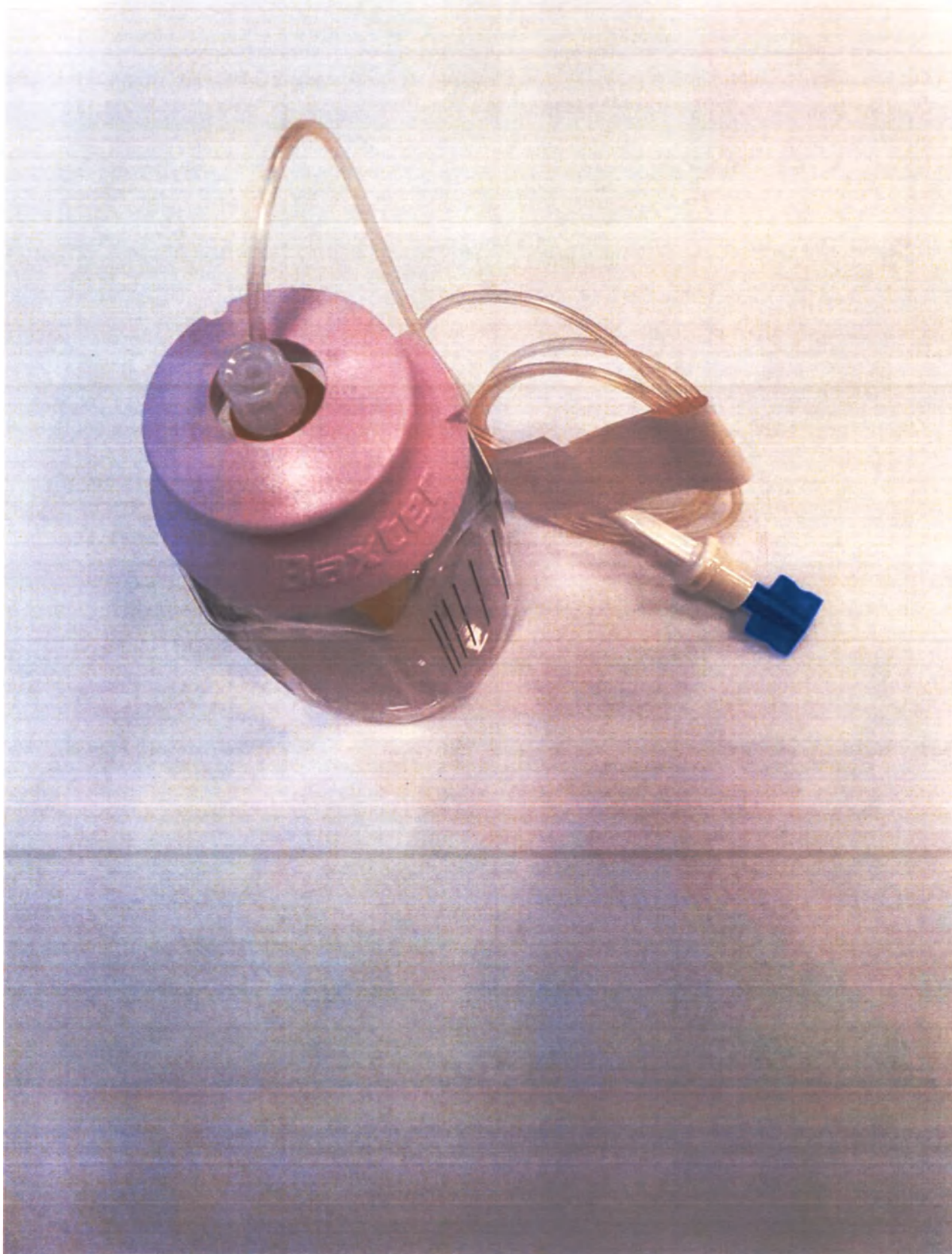
1.4. Помпа медицинская инфузионная эластомерная одноразовая стерильная Foliusor SV 0,5 мл/ч; Комплект поставки



1.5. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor SV 0,5 мл/ч;
Помпа, лицевая сторона



1.6. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor SV 0,5 мл/ч;
Помпа, сторона со шкалой инфузии



1.7. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor SV 0,5 мл/ч;
Помпа, вид сверху

Baxter

FOLFUSOR SV Portable Elastomeric Infusion System

REF	Description	Flow Rate	Flow Rate	Infused Volume	Flow Rate	Flow Rate	Approximate Infused Volume
10000	FOLFUSOR SV 0.5	0.5 ml/h	0.5 ml/h	100 ml	0.5 ml/h	0.5 ml/h	100 ml
10001	FOLFUSOR SV 1	1 ml/h	1 ml/h	100 ml	1 ml/h	1 ml/h	100 ml
10002	FOLFUSOR SV 2	2 ml/h	2 ml/h	100 ml	2 ml/h	2 ml/h	100 ml
10003	FOLFUSOR SV 3	3 ml/h	3 ml/h	100 ml	3 ml/h	3 ml/h	100 ml
10004	FOLFUSOR SV 4	4 ml/h	4 ml/h	100 ml	4 ml/h	4 ml/h	100 ml

*The device is designed to operate at maximum flow rate only (0.5 ml/h). Use of flow rates other than 0.5 ml/h will affect the flow rate of the device in Operating Conditions.

Description

The FOLFUSOR SV system is a portable, disposable device that uses an elastomeric reservoir to infuse medication, saline, or other fluids. The device is designed to provide a continuous flow of medication over the course of infusion of the medication. The flow rate is adjustable and can be set to 0.5, 1, 2, 3, 4, or 5 ml/h.

The device is designed to be used with a standard 100 ml infusion set. The device is designed to be used with a standard 100 ml infusion set.

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Operating Conditions

Caution: Please read the following instructions carefully before using the device.

1. **Impact of Volume**
The device is designed to operate at a flow rate of 0.5 ml/h. The flow rate will increase if the volume of the reservoir is increased. The flow rate will decrease if the volume of the reservoir is decreased.

2. **Impact of Temperature**
The device is designed to operate at a temperature of 20°C to 30°C. The flow rate will increase if the temperature is increased. The flow rate will decrease if the temperature is decreased.

3. **Impact of Viscosity of the Medication**
The device is designed to operate with a viscosity of 1 cP to 10 cP. The flow rate will increase if the viscosity is increased. The flow rate will decrease if the viscosity is decreased.

4. **Impact of Head Height**
The device is designed to operate at a head height of 0 to 100 cm. The flow rate will increase if the head height is increased. The flow rate will decrease if the head height is decreased.

5. **Impact of Catheter Size**
The device is designed to operate with a catheter size of 18 to 24 gauge. The flow rate will increase if the catheter size is increased. The flow rate will decrease if the catheter size is decreased.

6. **Impact of Catheter Length**
The device is designed to operate with a catheter length of 100 to 200 cm. The flow rate will increase if the catheter length is increased. The flow rate will decrease if the catheter length is decreased.

7. **Impact of Catheter Material**
The device is designed to operate with a catheter material of polyurethane or polyethylene. The flow rate will increase if the catheter material is polyurethane. The flow rate will decrease if the catheter material is polyethylene.

8. **Impact of Catheter Configuration**
The device is designed to operate with a catheter configuration of single lumen or double lumen. The flow rate will increase if the catheter configuration is single lumen. The flow rate will decrease if the catheter configuration is double lumen.

9. **Impact of Catheter Location**
The device is designed to operate with a catheter location of peripheral or central. The flow rate will increase if the catheter location is peripheral. The flow rate will decrease if the catheter location is central.

10. **Impact of Catheter Condition**
The device is designed to operate with a catheter condition of new or used. The flow rate will increase if the catheter condition is new. The flow rate will decrease if the catheter condition is used.

11. **Impact of Catheter Maintenance**
The device is designed to operate with a catheter maintenance of daily or weekly. The flow rate will increase if the catheter maintenance is daily. The flow rate will decrease if the catheter maintenance is weekly.

12. **Impact of Catheter Replacement**
The device is designed to operate with a catheter replacement of daily or weekly. The flow rate will increase if the catheter replacement is daily. The flow rate will decrease if the catheter replacement is weekly.

13. **Impact of Catheter Disposal**
The device is designed to operate with a catheter disposal of daily or weekly. The flow rate will increase if the catheter disposal is daily. The flow rate will decrease if the catheter disposal is weekly.

14. **Impact of Catheter Storage**
The device is designed to operate with a catheter storage of daily or weekly. The flow rate will increase if the catheter storage is daily. The flow rate will decrease if the catheter storage is weekly.

15. **Impact of Catheter Transport**
The device is designed to operate with a catheter transport of daily or weekly. The flow rate will increase if the catheter transport is daily. The flow rate will decrease if the catheter transport is weekly.

16. **Impact of Catheter Use**
The device is designed to operate with a catheter use of daily or weekly. The flow rate will increase if the catheter use is daily. The flow rate will decrease if the catheter use is weekly.

17. **Impact of Catheter Care**
The device is designed to operate with a catheter care of daily or weekly. The flow rate will increase if the catheter care is daily. The flow rate will decrease if the catheter care is weekly.

18. **Impact of Catheter Education**
The device is designed to operate with a catheter education of daily or weekly. The flow rate will increase if the catheter education is daily. The flow rate will decrease if the catheter education is weekly.

19. **Impact of Catheter Research**
The device is designed to operate with a catheter research of daily or weekly. The flow rate will increase if the catheter research is daily. The flow rate will decrease if the catheter research is weekly.

20. **Impact of Catheter Development**
The device is designed to operate with a catheter development of daily or weekly. The flow rate will increase if the catheter development is daily. The flow rate will decrease if the catheter development is weekly.

21. **Impact of Catheter Innovation**
The device is designed to operate with a catheter innovation of daily or weekly. The flow rate will increase if the catheter innovation is daily. The flow rate will decrease if the catheter innovation is weekly.

22. **Impact of Catheter Improvement**
The device is designed to operate with a catheter improvement of daily or weekly. The flow rate will increase if the catheter improvement is daily. The flow rate will decrease if the catheter improvement is weekly.

23. **Impact of Catheter Advancement**
The device is designed to operate with a catheter advancement of daily or weekly. The flow rate will increase if the catheter advancement is daily. The flow rate will decrease if the catheter advancement is weekly.

24. **Impact of Catheter Progression**
The device is designed to operate with a catheter progression of daily or weekly. The flow rate will increase if the catheter progression is daily. The flow rate will decrease if the catheter progression is weekly.

25. **Impact of Catheter Evolution**
The device is designed to operate with a catheter evolution of daily or weekly. The flow rate will increase if the catheter evolution is daily. The flow rate will decrease if the catheter evolution is weekly.

26. **Impact of Catheter Revolution**
The device is designed to operate with a catheter revolution of daily or weekly. The flow rate will increase if the catheter revolution is daily. The flow rate will decrease if the catheter revolution is weekly.

27. **Impact of Catheter Transformation**
The device is designed to operate with a catheter transformation of daily or weekly. The flow rate will increase if the catheter transformation is daily. The flow rate will decrease if the catheter transformation is weekly.

28. **Impact of Catheter Change**
The device is designed to operate with a catheter change of daily or weekly. The flow rate will increase if the catheter change is daily. The flow rate will decrease if the catheter change is weekly.

29. **Impact of Catheter Conversion**
The device is designed to operate with a catheter conversion of daily or weekly. The flow rate will increase if the catheter conversion is daily. The flow rate will decrease if the catheter conversion is weekly.

30. **Impact of Catheter Modification**
The device is designed to operate with a catheter modification of daily or weekly. The flow rate will increase if the catheter modification is daily. The flow rate will decrease if the catheter modification is weekly.

31. **Impact of Catheter Adjustment**
The device is designed to operate with a catheter adjustment of daily or weekly. The flow rate will increase if the catheter adjustment is daily. The flow rate will decrease if the catheter adjustment is weekly.

32. **Impact of Catheter Alteration**
The device is designed to operate with a catheter alteration of daily or weekly. The flow rate will increase if the catheter alteration is daily. The flow rate will decrease if the catheter alteration is weekly.

33. **Impact of Catheter Amendment**
The device is designed to operate with a catheter amendment of daily or weekly. The flow rate will increase if the catheter amendment is daily. The flow rate will decrease if the catheter amendment is weekly.

34. **Impact of Catheter Addition**
The device is designed to operate with a catheter addition of daily or weekly. The flow rate will increase if the catheter addition is daily. The flow rate will decrease if the catheter addition is weekly.

35. **Impact of Catheter Subtraction**
The device is designed to operate with a catheter subtraction of daily or weekly. The flow rate will increase if the catheter subtraction is daily. The flow rate will decrease if the catheter subtraction is weekly.

36. **Impact of Catheter Multiplication**
The device is designed to operate with a catheter multiplication of daily or weekly. The flow rate will increase if the catheter multiplication is daily. The flow rate will decrease if the catheter multiplication is weekly.

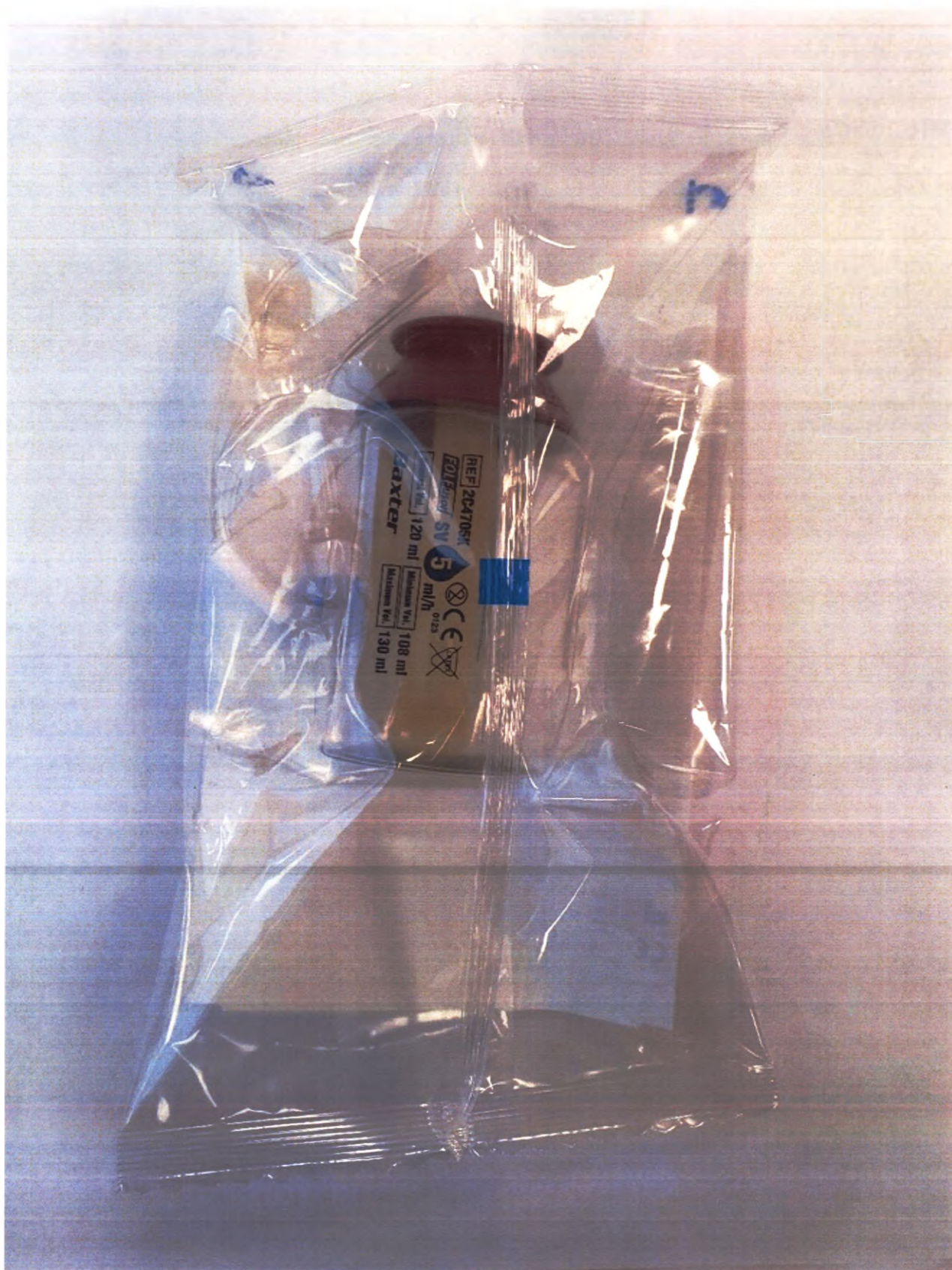
1.8. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor SV 0,5 мл/ч; Оригинальная инструкция

2. Помпа медицинская инфузионная эластомерная одноразовая стерильная

Folfusor SV 5 мл/ч

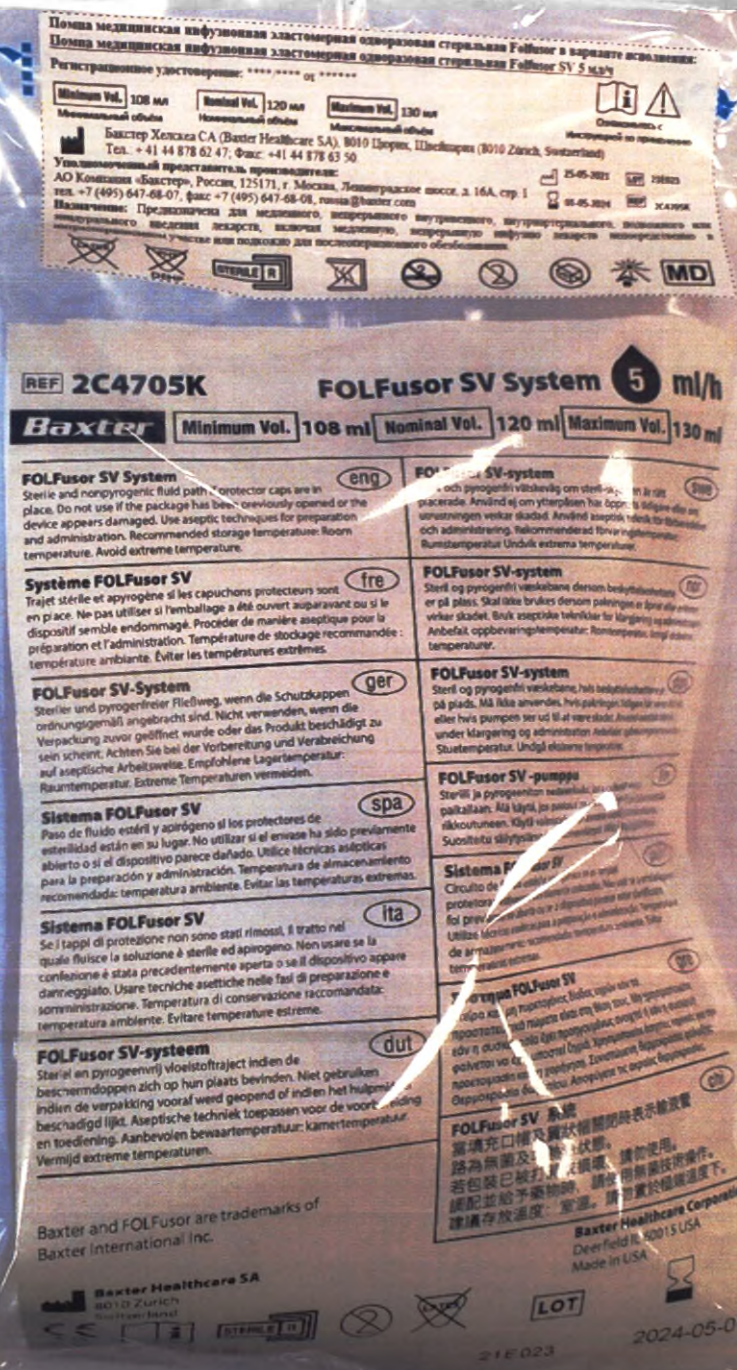


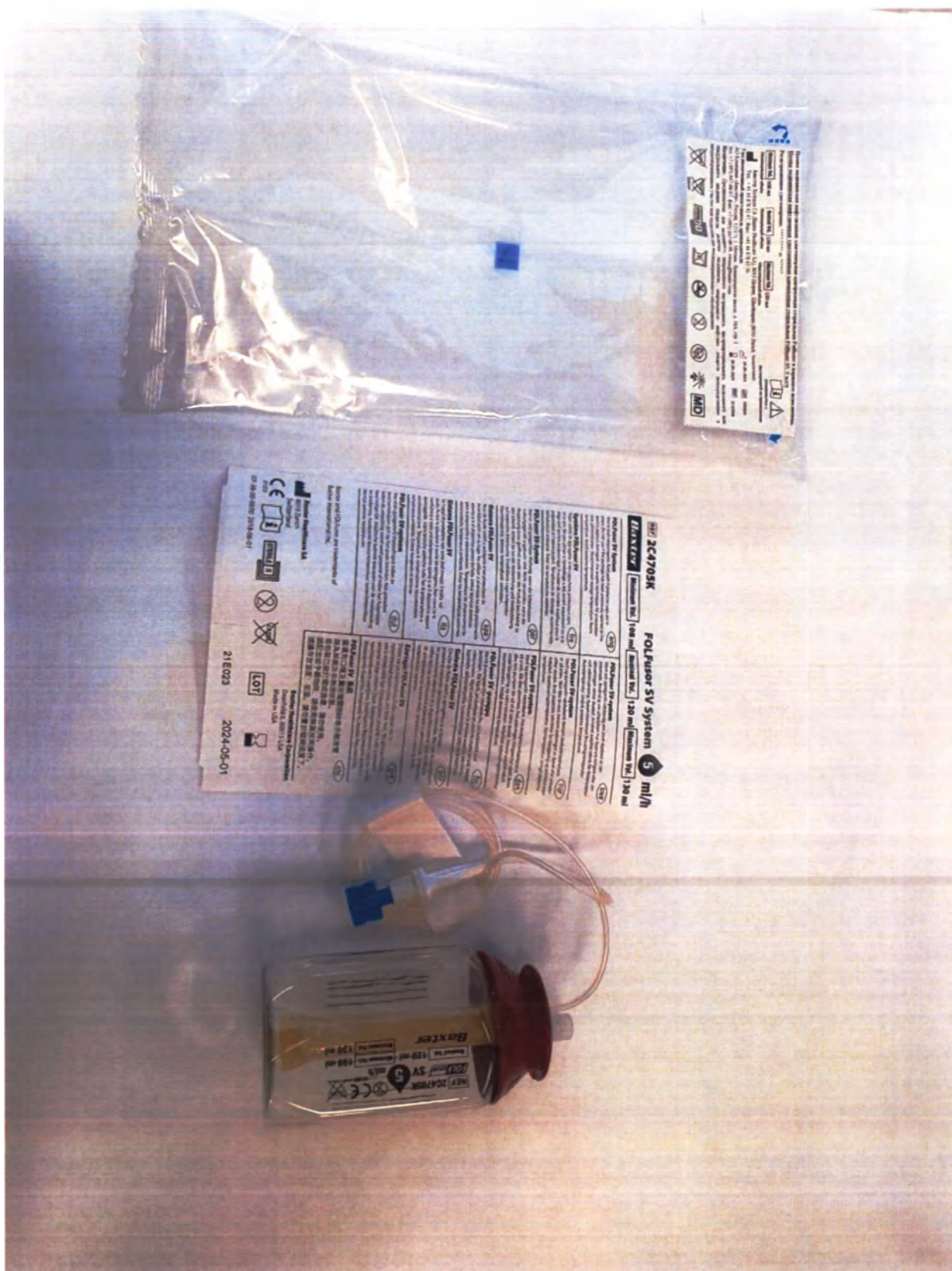
2.1. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor SV 5 мл/ч; Индивидуальная упаковка, лицевой сторона, с оригинальной маркировкой



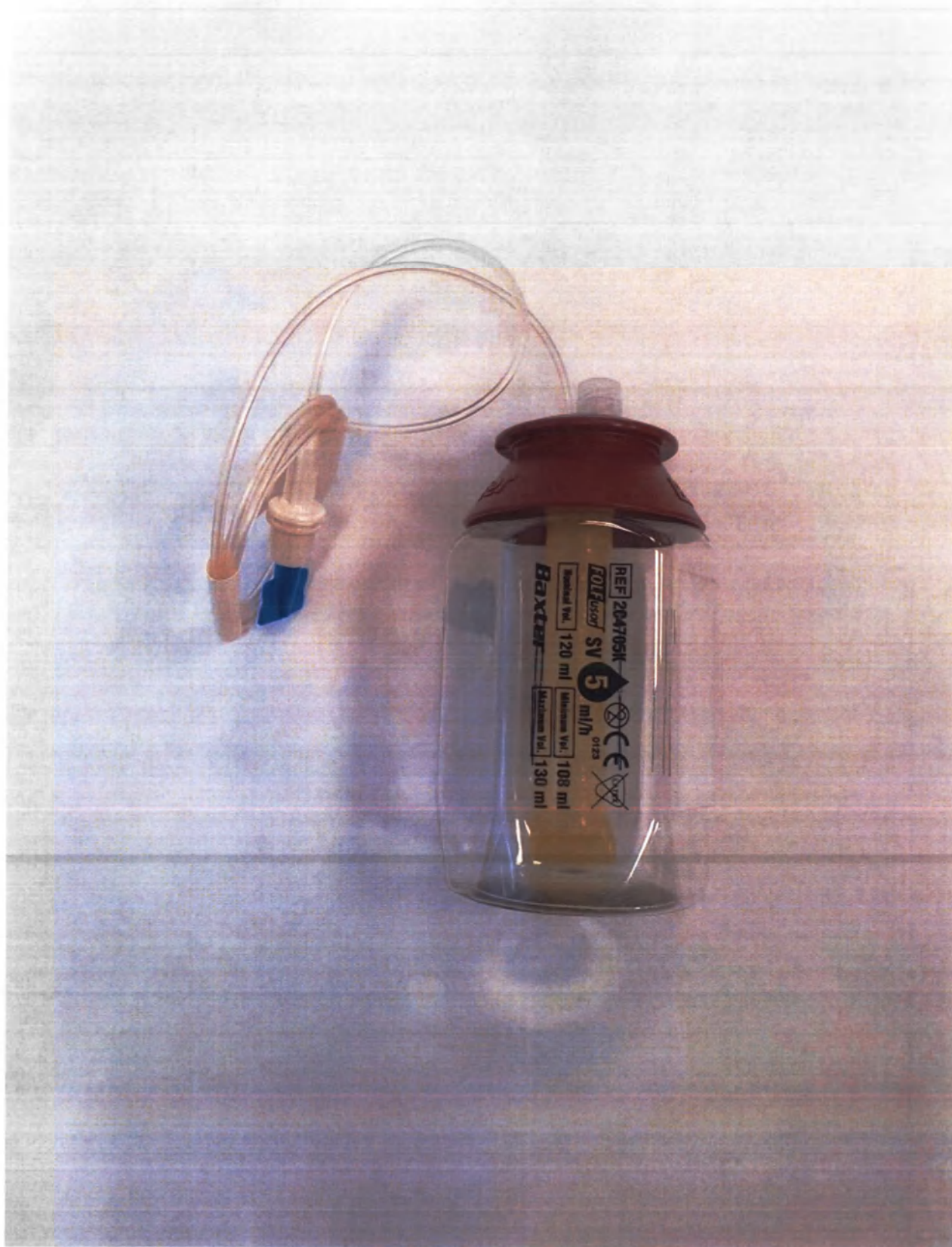
2.2. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor SV 5 мл/ч;
Индивидуальная упаковка, оборотная сторона

2.3. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor SV 5 мл/ч; Индивидуальная упаковка, лицевая сторона, с нанесенной маркировкой на русском языке





2.4. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor SV 5 мл/ч;
Комплект поставки



2.5. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor SV 5 мл/ч;
Помпа, лицевая сторона



2.6. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor SV 5 мл/ч;
Помпа, сторона со шкалой инфузии



**2.7. Помпа медицинская инфузионная эластомерная одноразовая стерильная Foliusor SV 5 мл/ч;
Помпа, вид сверху**

REF	Description	Barcode Id.	Barcode File Size	Barcode Delivery Size	Barcode Id.	Barcode File Size	Aggregated Estimated Volume
A-0000	CD-ROM 10-18	0000	32 MB	320	0000	32000	1 MB
A-0001	CD-ROM 10-19	0001	1 MB	100	0001	10000	1 MB
V-0000	CD-ROM 10-20	0000	1 MB	100	0000	10000	1 MB
V-0001	CD-ROM 10-21	0001	1 MB	100	0001	10000	1 MB
V-0002	CD-ROM 10-22	0002	1 MB	100	0002	10000	1 MB
V-0003	CD-ROM 10-23	0003	1 MB	100	0003	10000	1 MB
V-0004	CD-ROM 10-24	0004	1 MB	100	0004	10000	1 MB

*The engine is designed to operate at nominal fire rating of 1A, however, if the use of elements other than DOW will affect the fire rating of the device, the Operating Conditions must be modified.

The shipping carton may not affect the flow data accuracy. (1) Fill solution, (2) Temperature, (3) Viscosity of the solution, (4) The difference in height between the 700 Port and the Digital End User-Level-Flow Indicator, and (5) Calculator may result in Sparging Conditions.

The 100-psi or greater reducing capacity, ultrasonically-qualified and tested to ASME BPE Class 3B, VCR, and most VCRs, etc.

The product is not manufactured using GMP practices.

The product is not manufactured with a lot of natural value.

The intended use of the "C" User ID option includes the slow, continuous intravenous infusion of a solution or solution of medications, to try and reduce the slow, continuous infusion of medications directly into an intravenous site or subcutaneous or percutaneous skin placement.

It is a change recommendation that a "full drug argument" be updated drug safety to clearly demonstrate that the drug is safe for use in the context of administration.

It is the responsibility of the health-care provider to ensure the use of an appropriate device when administering injections of critical medications (such as morphine and the calcium preparations) when

- Integrated education, cross-generational education would stay the same in terms of health.**
 1. **Integrated education:** The relationship between the two is that the students will remain just present throughout the whole school.
 2. **Cross-generational education:** The relationship between the two is that the students will remain just present throughout the whole school.

The POLY user's guide must be filled in accordance with the procedure described under **Reactions for Filling and Printing**.

is the responsibility of the healthcare provider to instruct the patient and caregivers of the following

- [illegible]

Caution: Please note that a single or combination of the following factors may impact the fire risk.

- The 100-Pass 100-Volume is designed to increase the overall flow rate of water filled with a specific "gelling" from the maximum 100 volume to the maximum 100 volume using more product information table above.
- Do not fill effluents from the maximum 100 volume or more than the maximum 100 volume. It was greater information table above. Filling the device with less than the maximum 100 volume will result in increased flow rate and turbulence in filling time.

- Extended storage of the fluid disks at a temperature environment may result in a decrease in flow rate and increase in delivery time.

• The PCC/Juice 20 system is designed to operate at the hottest flow rate using 70 Gallons (265 L) per day. Flow will be an approximately 12% increase in the normal flow rate of 20% Sodium Chloride (NaCl) a week.

The PLR-participation is assigned to each of the national flow between the FR Part and the total of the Low/Low/Flow between is positioned at approximately the same height. Flow will decrease approximately 1% for every 254 cm (1 inch) the FR Part is positioned below the threshold and Low/Flow between will increase approximately 0.5% for every 254 cm (1 inch) the FR Part is positioned above the threshold. Low/Low/Flow between is positioned at approximately the same height.

• The POLYMER is extremely easy used & identical to the normal flow rate when used with 22 gauge (3/16 inch) or less of a cannula smaller than 22 gauge (3/16 inch) with clearance. **Flow rate:** Also refer to the directions for use provided by the cannula manufacturer.

Directions for Use
Mixing and Use Information:

Directions for Filling and Priming

Warning: Use aseptic technique throughout the procedure.

When withdrawing medication from a glass ampule, use a Star needle and remove the needle before filling the

- [illegible]

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1. To start the session, review the *Wingspan* Card that the students have. Lay the *Wingspan* Matrix that each student has, placing it on top of the *Wingspan* Card that they have.
2. Read the *Wingspan* Matrix aloud.
3. Read each address aloud and have students correct the card held by the student who is on that line again.
4. To adjust the game even more, have students write the *Wingspan* Card that they have on a piece of paper and place it on the table.

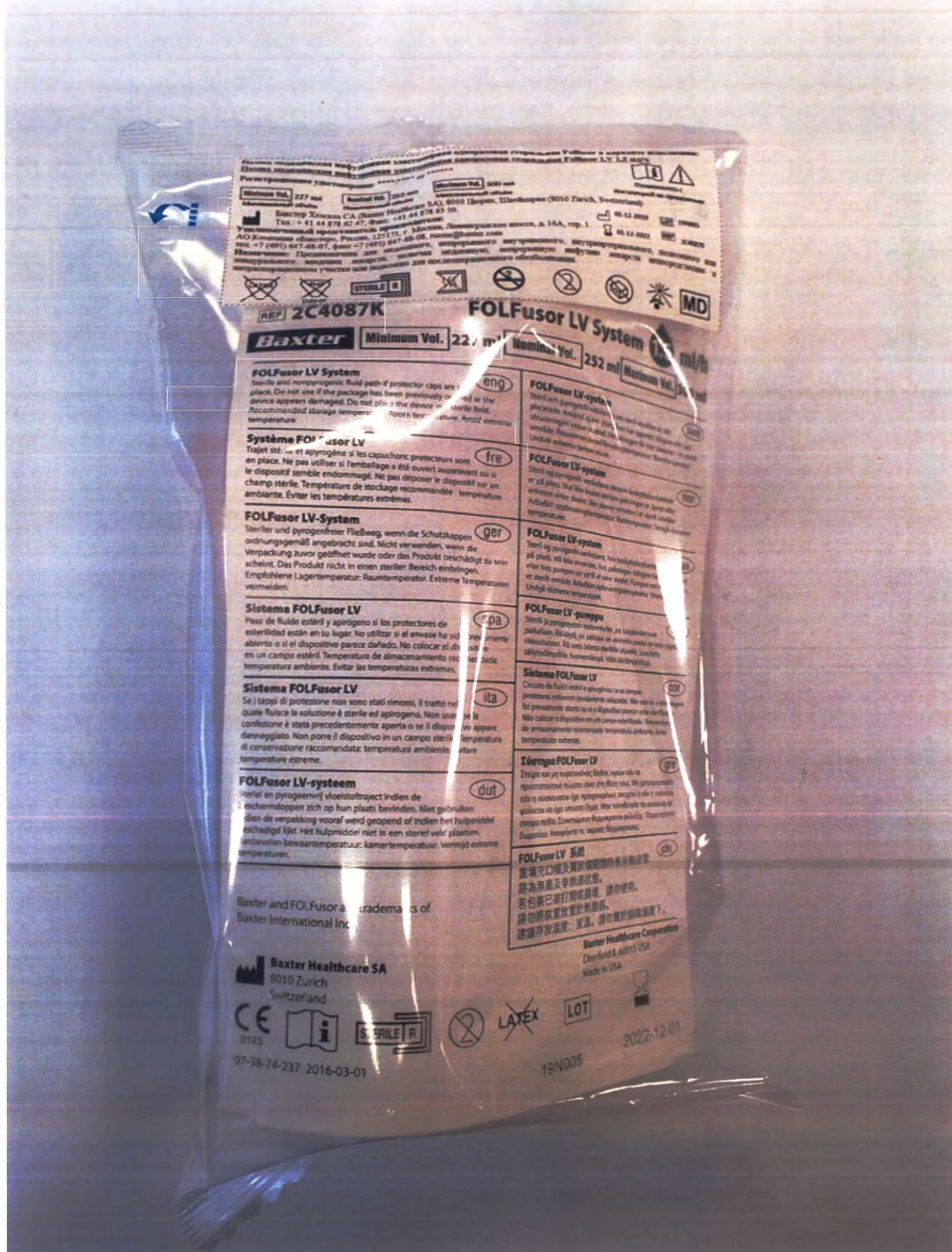
3. Помпа медицинская инфузионная эластомерная одноразовая стерильная
Foliusor LV 1,5 мл/ч



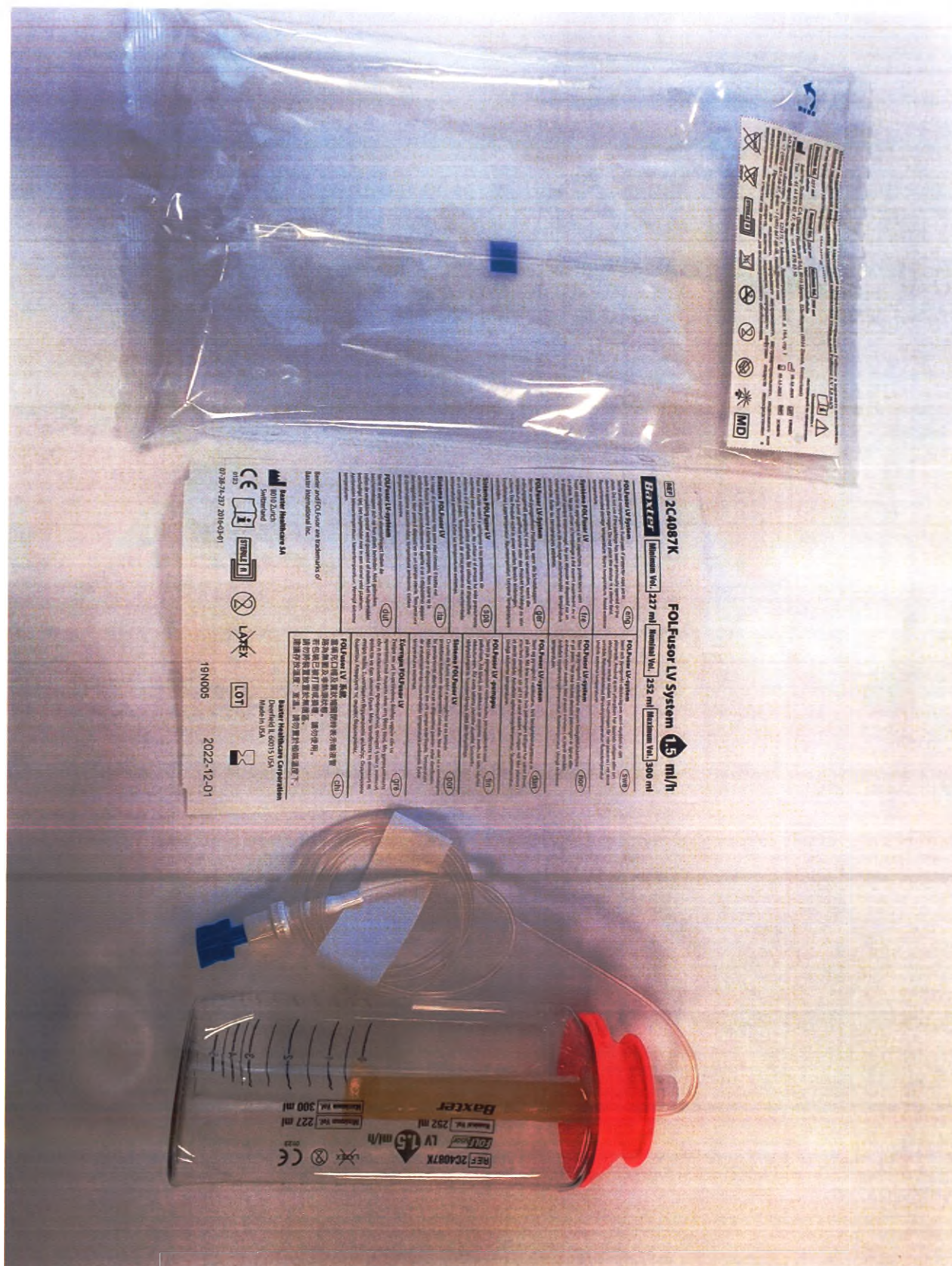
3.1. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 1,5 мл/ч; Индивидуальная упаковка, лицевой сторона, с оригинальной маркировкой



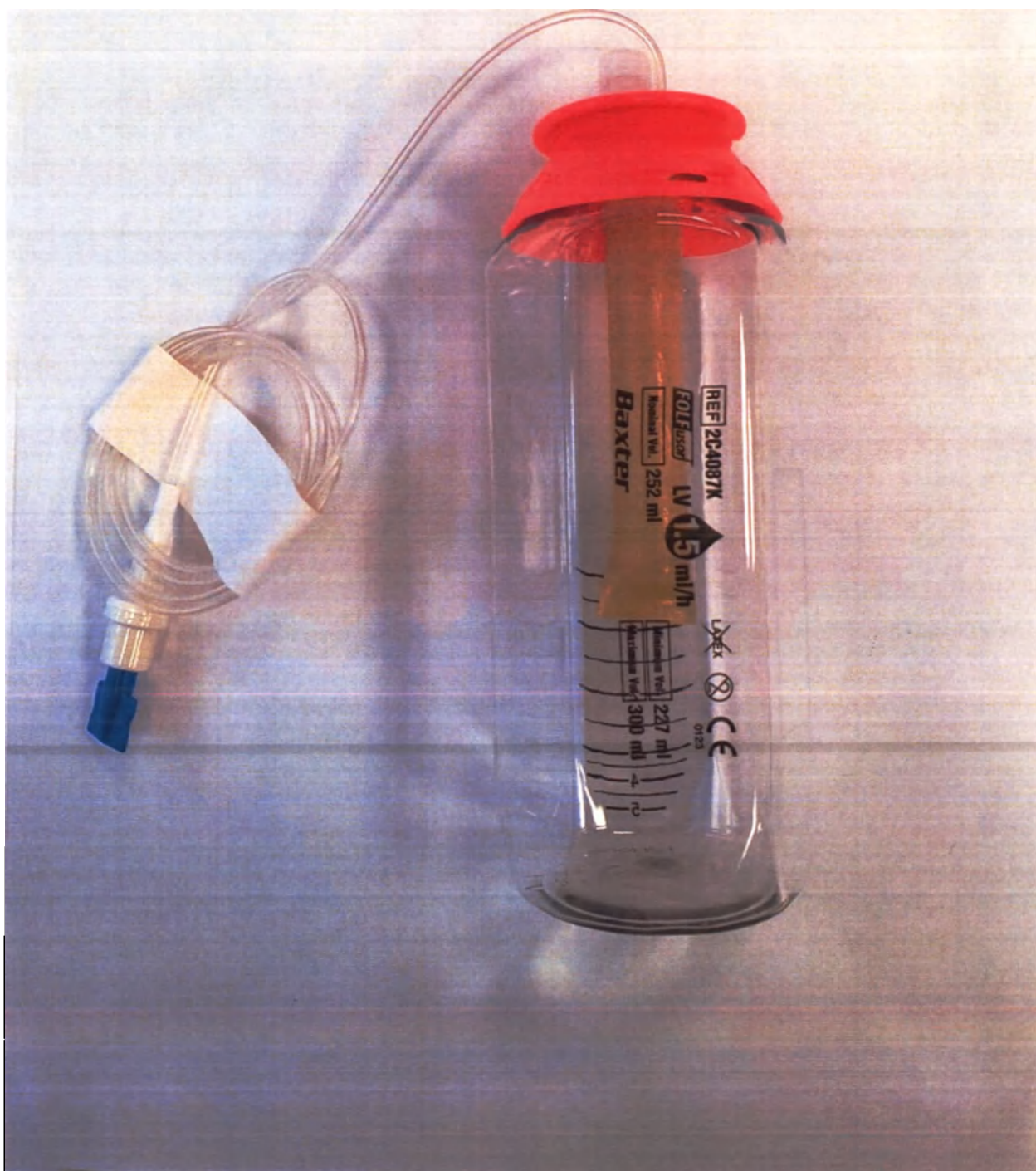
3.2. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 1,5 мл/ч; Индивидуальная упаковка, обратная сторона



3.3. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 1,5 мл/ч; Индивидуальная упаковка, лицевая сторона, с нанесенной маркировкой на русском языке



3.4. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 1,5 мл/ч; Комплект поставки



3.5. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 1,5 мл/ч; Помпа, лицевая сторона



3.6. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 1,5 мл/ч;
Помпа, сторона со шкалой инфузии



**3.7. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 1,5 мл/ч;
Помпа. вид сверху**

FOLFusor LV Portable Elastomeric Infusion System

REF	Designation	Revised In	Revised Date/Rev	Revised Delivery Time	Revised In	Revised In	Revised Delivery Time
2-020	100,000 1.1	2000	1.1.00	1000	1000	1000	1.1.00
2-020	100,000 1.2	2000	1.1.00	1000	1000	1000	1.1.00
2-020	100,000 1.3	2000	1.1.00	1000	1000	1000	1.1.00
2-020	100,000 1.4	2000	1.1.00	1000	1000	1000	1.1.00
2-020	100,000 1.5	2000	1.1.00	1000	1000	1000	1.1.00
2-020	100,000 1.6	2000	1.1.00	1000	1000	1000	1.1.00
2-020	100,000 1.7	2000	1.1.00	1000	1000	1000	1.1.00
2-020	100,000 1.8	2000	1.1.00	1000	1000	1000	1.1.00
2-020	100,000 1.9	2000	1.1.00	1000	1000	1000	1.1.00
2-020	100,000 2.0	2000	1.1.00	1000	1000	1000	1.1.00

*The device is designed to operate at nominal flow rate using 1% Dextrose (D5W). Use of diluents other than D5W will affect the flow rate of the device (see Operating Conditions).

The FQ4000 IV system is a laboratory, disposable device that acts as a dynamic model of a renal radiator. When fluid, the FQ4000 IV system can be used with a constant flow pump. Conversely, the FQ4000 IV system can be used with a peristaltic pump and a flow restrictor.

The FQ4000 IV system is designed to provide a continuous flow of medication over a period of minutes and the narrow bore with an accuracy of $\pm 1\%$ after fluid in the volume chamber has reached the minimum fill volume to the volume of volume plus the residual volume.

Our drawing editors may be used to draw 2D and 3D objects. [1] File editors, [2] Parameters, [3] Assembly of the visualization, [4] The differences in layout between the PDS part and the Design and Low Level Part Editors are, and 5. Cellular case prior to digitizing constraints.

For a 3D part system, drawing consists of drawing 3D objects in 2D on which both JPL, LVC, and most others.

The product is not manufactured using 3D printers.

The product is not manufactured with natural rubber latex.

The intended use of the POU Water V-System includes the slow, continuous infusion, intravenous administration, or epidural administration of medication. It may also include the slow, continuous infusion of medications directly into an intrathecal site or subcutaneously for postoperative pain management.

Special Administration

Guidelines administration of analgesics is limited to use with intravenous catheters specifically indicated by other directions in drug product literature for intravenous administration.

Important information: drugs that are not indicated for epidural use, or that are contraindicated for this route, are contraindicated.

This company recommends that a PO/IV use label and for injectable drugs/delivery be clearly differentiated from a PO/IV use label. Please contact your local regulatory agency for more information.

It is the responsibility of the healthcare provider to ensure the use of an appropriate device when administering treatment of critical medications (such as Morphine) and life sustaining medications when unplanned cessation, interruption or overdose would likely lead to serious injury or death.

- [illegible]

The FCC's final rule states that, in accordance with the procedure described in the Commission's existing

- Allow the fluid TCU cool/UP system to reach room temperature before mixing. Do not apply a heating source to bring the disks to room temperature.

is the responsibility of the health-care provider to inspect the patient and determine if the following

To achieve the desired results, you must ensure that the data is accurate and that the system is properly configured.

- [illegible]

Answer: Please note that a single or combination of the following factors may impact the flow rate:

- The PGL2 gene LV system is designed to operate at the nominal flow rate when it is backfilled with a viscous slurry with the same viscosity as the in-process slurry (e.g. 1000 cP).

Do not fill with less than the minimum fill volume or more than the maximum fill volume; jaw product information cards show. Filling the device with less than the minimum fill volume will result in decreased flow rates and reduction in delivery time.

- **Impact of Temperature**
 - The CO₂ Arrhenius Temperature is generally at the normal form where the maximum CO₂ is approximately 1000°C, but can be as low as 700°C and as high as 2000°C, 2600°C, and 3000°C. To remove the optimal temperature for the CO₂ Arrhenius Temperature, the Arrhenius Temperature can be used to find the CO₂ rate of decomposition approximately 2.7 x 10⁻¹⁰ s⁻¹ at 1000°C. The Arrhenius Temperature can be used to find the CO₂ rate of decomposition approximately 2.7 x 10⁻¹⁰ s⁻¹ at 1000°C.
 - Extended periods of the CO₂ rate of decomposition can be used to find the CO₂ rate of decomposition at 1000°C.
- **Impact of the Number of the CO₂ Rate**
 - The CO₂ rate of decomposition is dependent on the CO₂ rate of decomposition at 1000°C. The CO₂ rate of decomposition is dependent on the CO₂ rate of decomposition at 1000°C.
 - The CO₂ rate of decomposition is dependent on the CO₂ rate of decomposition at 1000°C.
- **Impact of the Heat Flow**
 - The CO₂ rate of decomposition is dependent on the CO₂ rate of decomposition at 1000°C. The CO₂ rate of decomposition is dependent on the CO₂ rate of decomposition at 1000°C.
 - The CO₂ rate of decomposition is dependent on the CO₂ rate of decomposition at 1000°C.
- **Impact of the CO₂ Rate**
 - The CO₂ rate of decomposition is dependent on the CO₂ rate of decomposition at 1000°C. The CO₂ rate of decomposition is dependent on the CO₂ rate of decomposition at 1000°C.
 - The CO₂ rate of decomposition is dependent on the CO₂ rate of decomposition at 1000°C.

Mixing and the Information

data is the only information storage device for long-term storage and storage procedures.

not use if the package has been previously opened or the device appears damaged.

Warning: the deeper techniques throughout each procedure.
 Item failure to accurately reflect the items was made in follow-up research

Don't forget to read what they say about your work.

7. Remove the Shreddy Protection Cap from the FOLFuser. If present, install the Shreddy Protection Cap to replace

- [illegible]

Refer to the Information for Patients and Healthcare Provider section.

1. To place the network, including the gateway, on the same host as the Data Center LinkFlow Controller, modify the

1. Measure the distance from the center of the wheel to the center of the axle. This is the radius of the wheel.
2. Measure the distance from the center of the wheel to the center of the axle. This is the radius of the wheel.
3. Measure the distance from the center of the wheel to the center of the axle. This is the radius of the wheel.
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8. Measure the distance from the center of the wheel to the center of the axle. This is the radius of the wheel.
9. Measure the distance from the center of the wheel to the center of the axle. This is the radius of the wheel.
10. Measure the distance from the center of the wheel to the center of the axle. This is the radius of the wheel.

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4. Помпа медицинская инфузионная эластомерная одноразовая стерильная
Folfusor LV 2 мл/ч



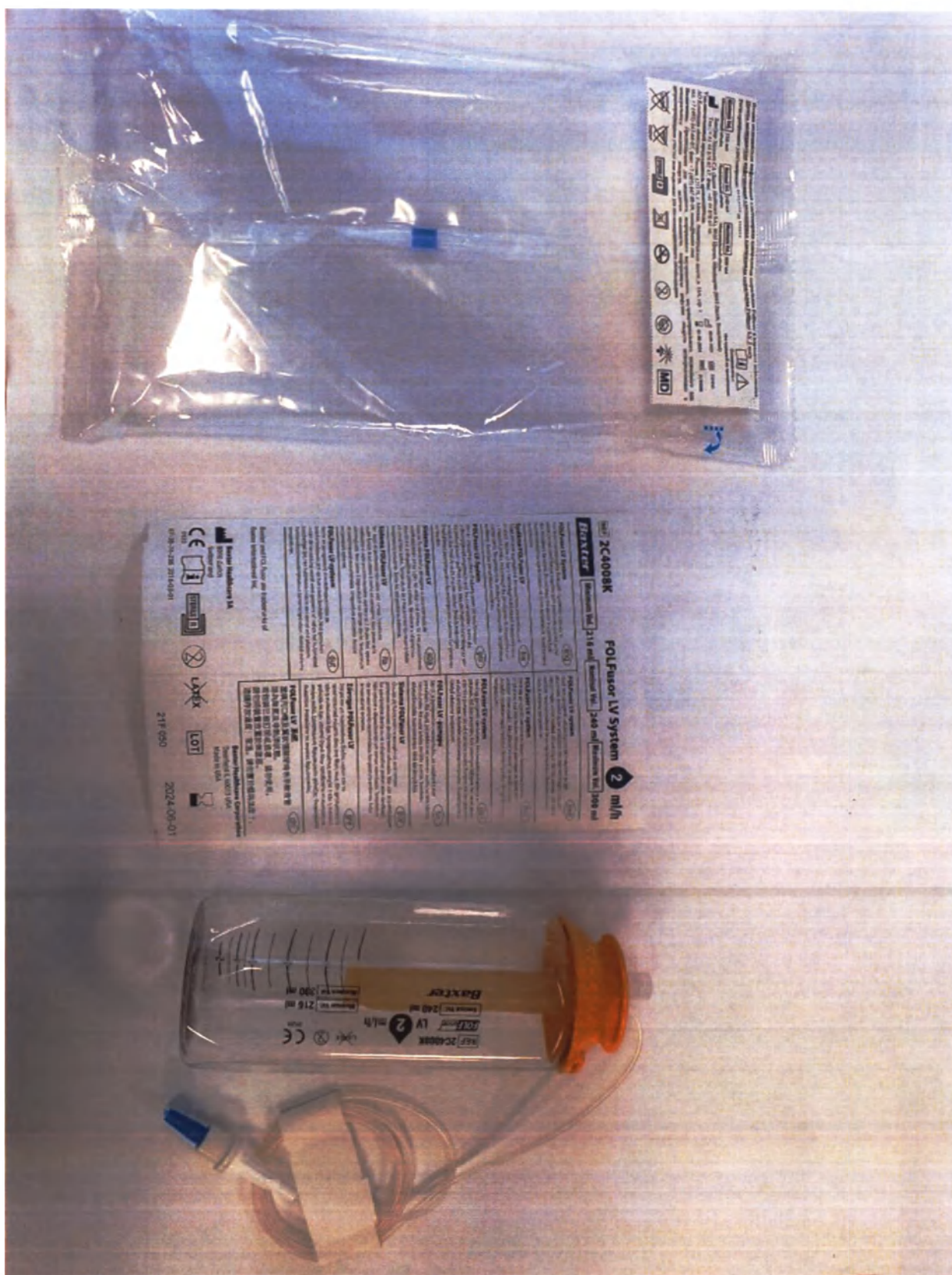
4.1. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 2 мл/ч; Индивидуальная упаковка, лицевой сторона, с оригинальной маркировкой



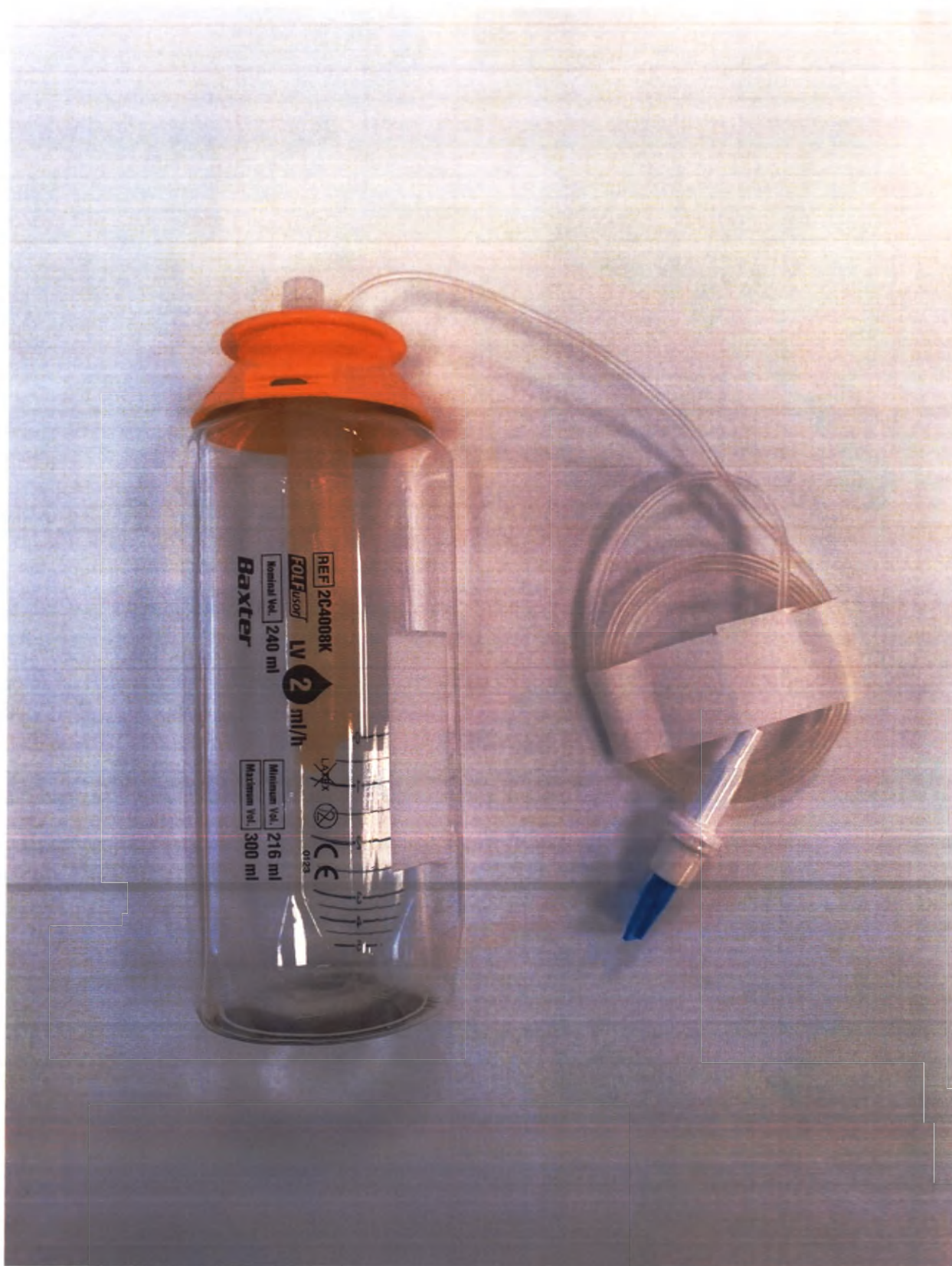
4.2. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 2 мл/ч; Индивидуальная упаковка, обратная сторона



4.3. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 2 мл/ч; Индивидуальная упаковка, лицевая сторона, с нанесенной маркировкой на русском языке



4.4. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 2 мл/ч;
Комплект поставки



4.5. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 2 мл/ч; Помпа, лицевая сторона



4.6. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 2 мл/ч;
Помпа, сторона со шкалой инфузии



4.7. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 2 мл/ч; Помпа, вид сверху

5. Помпа медицинская инфузионная эластомерная одноразовая стерильная
Folfusor LV 5 мл/ч



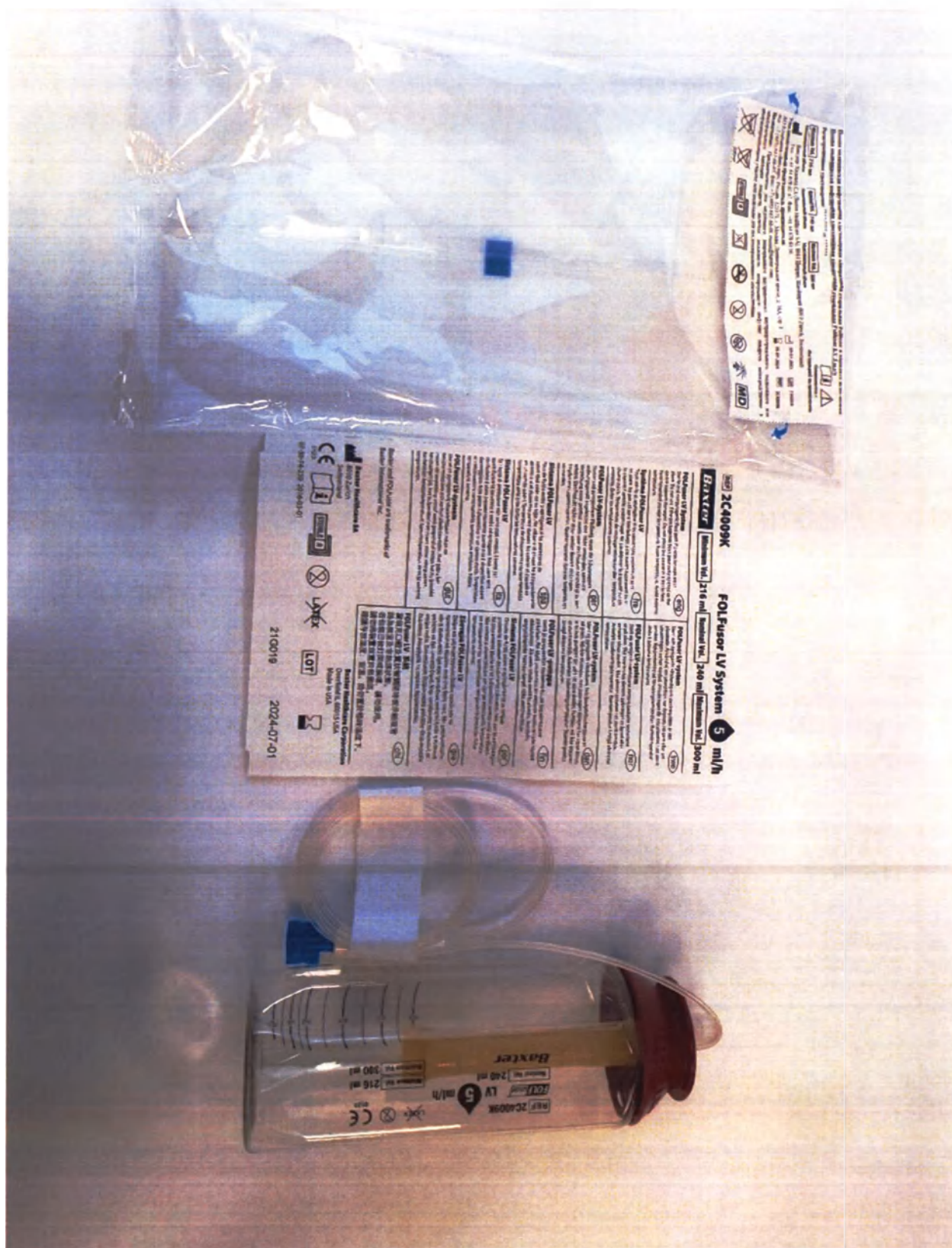
5.1. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 5 мл/ч
Индивидуальная упаковка, лицевой сторона, с оригинальной маркировкой



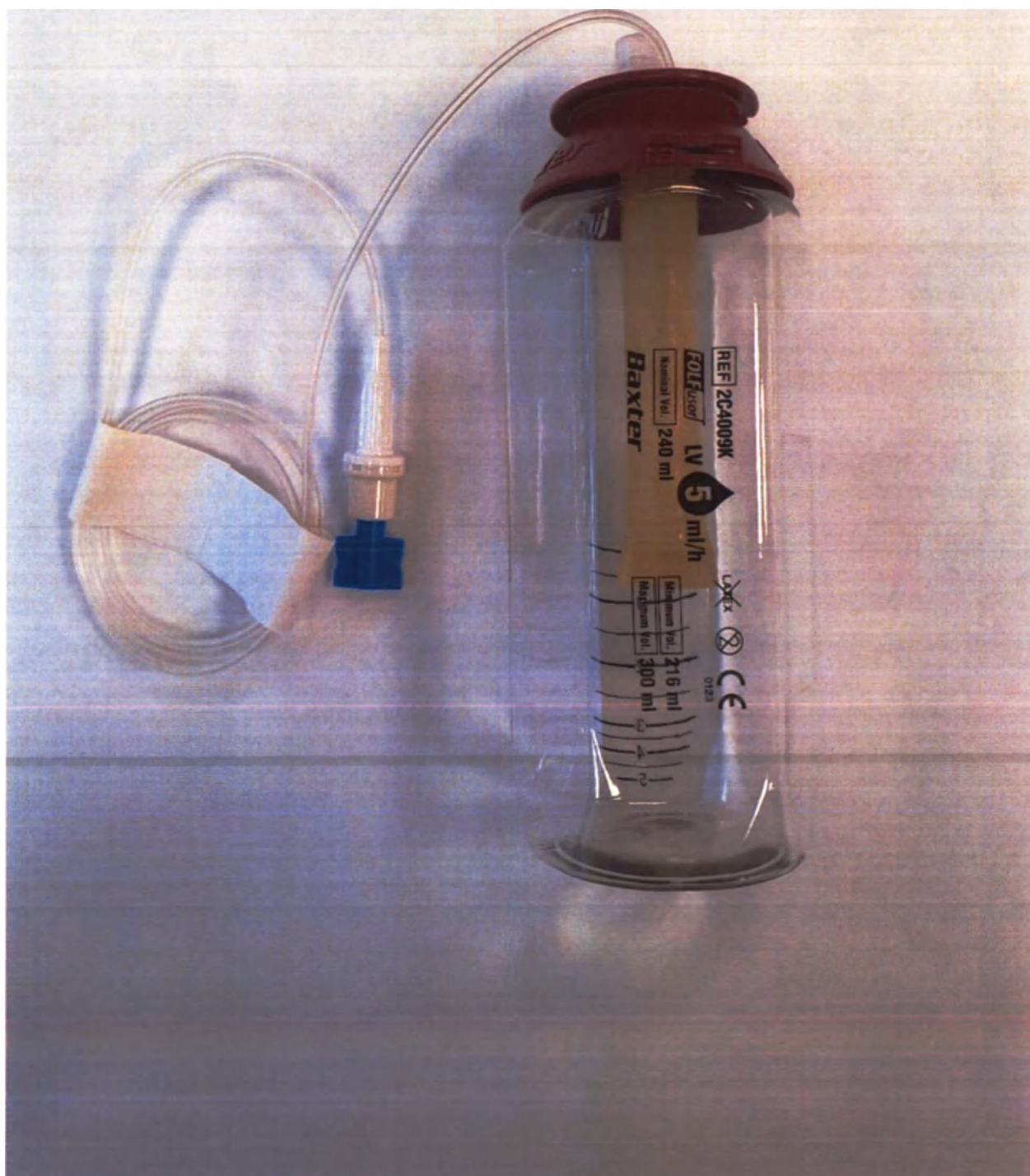
5.2. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 5 мл/ч
Индивидуальная упаковка, оборотная сторона



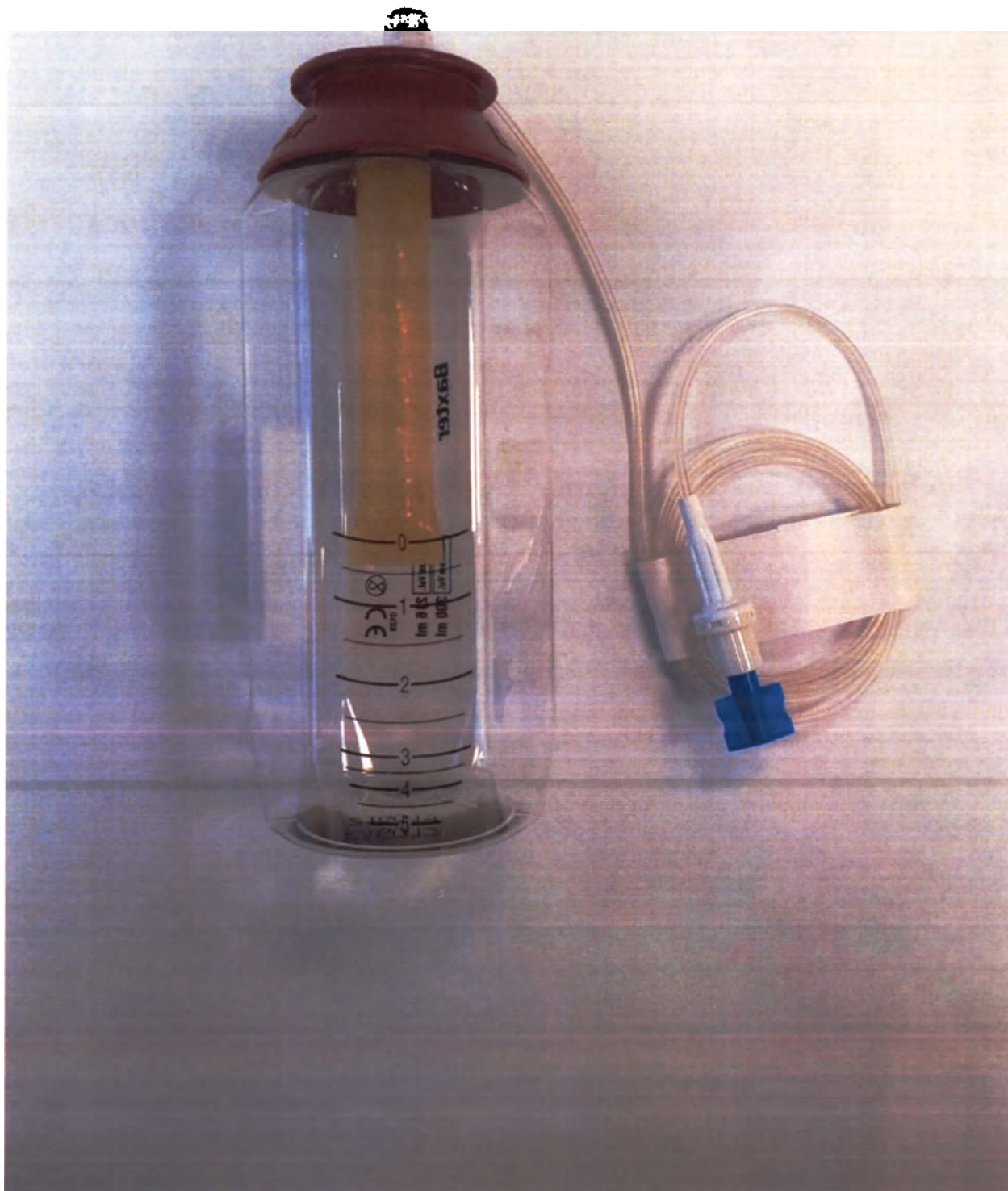
5.3. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 5 мл/ч
Индивидуальная упаковка, лицевая сторона, с нанесенной маркировкой на русском языке



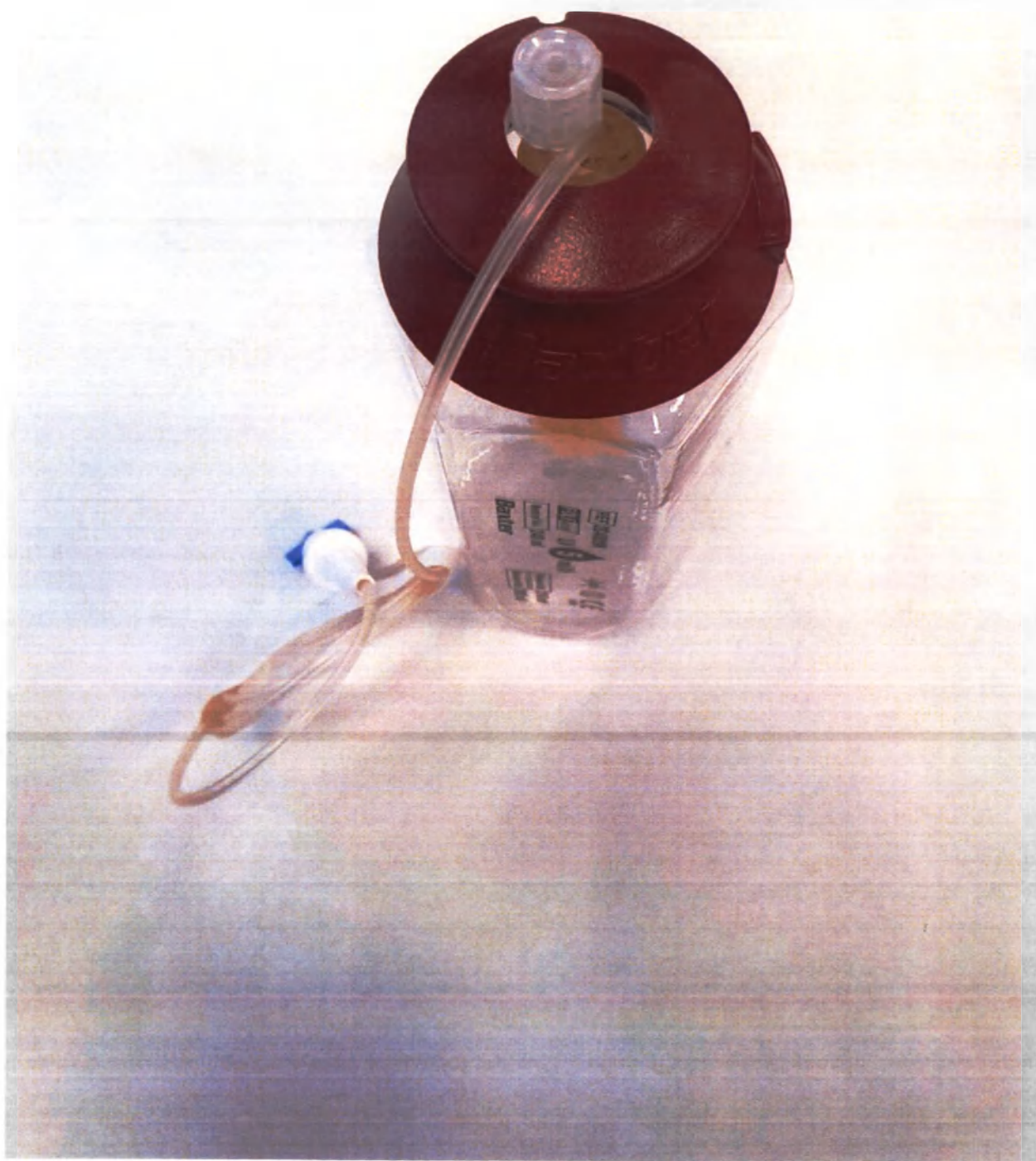
5.4. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 5 мл/ч
Комплект поставки



5.5. Помпа медицинская инфузионная эластомерная одноразовая стерильная FolFusor LV 5 мл/ч
Помпа. лицевая сторона

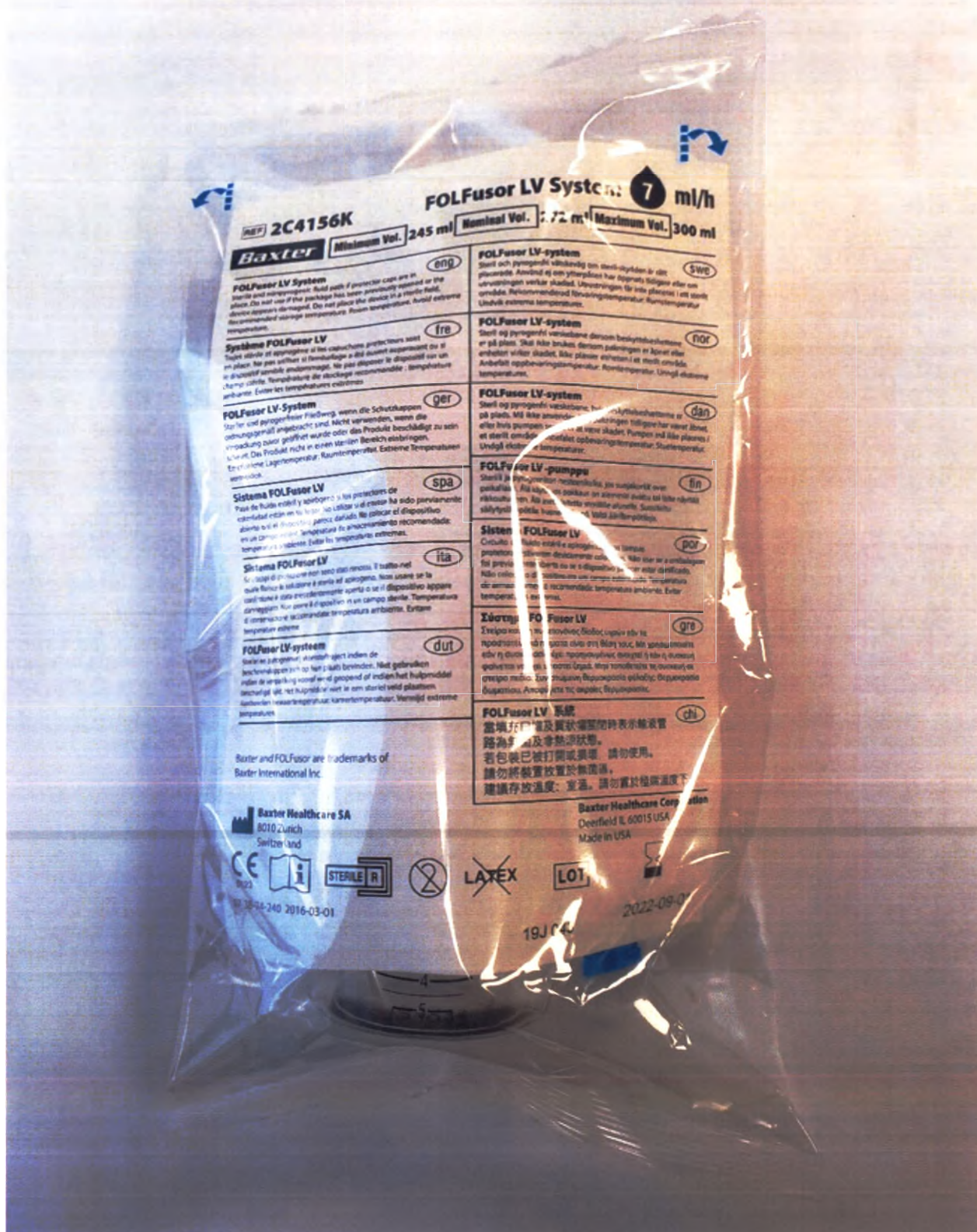


5.6. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 5 мл/ч
Помпа, сторона со шкалой инфузии



5.7. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 5 мл/ч
Помпа, вид сверху

6. Помпа медицинская инфузионная эластомерная одноразовая стерильная
Folfusor LV 7 мл/ч



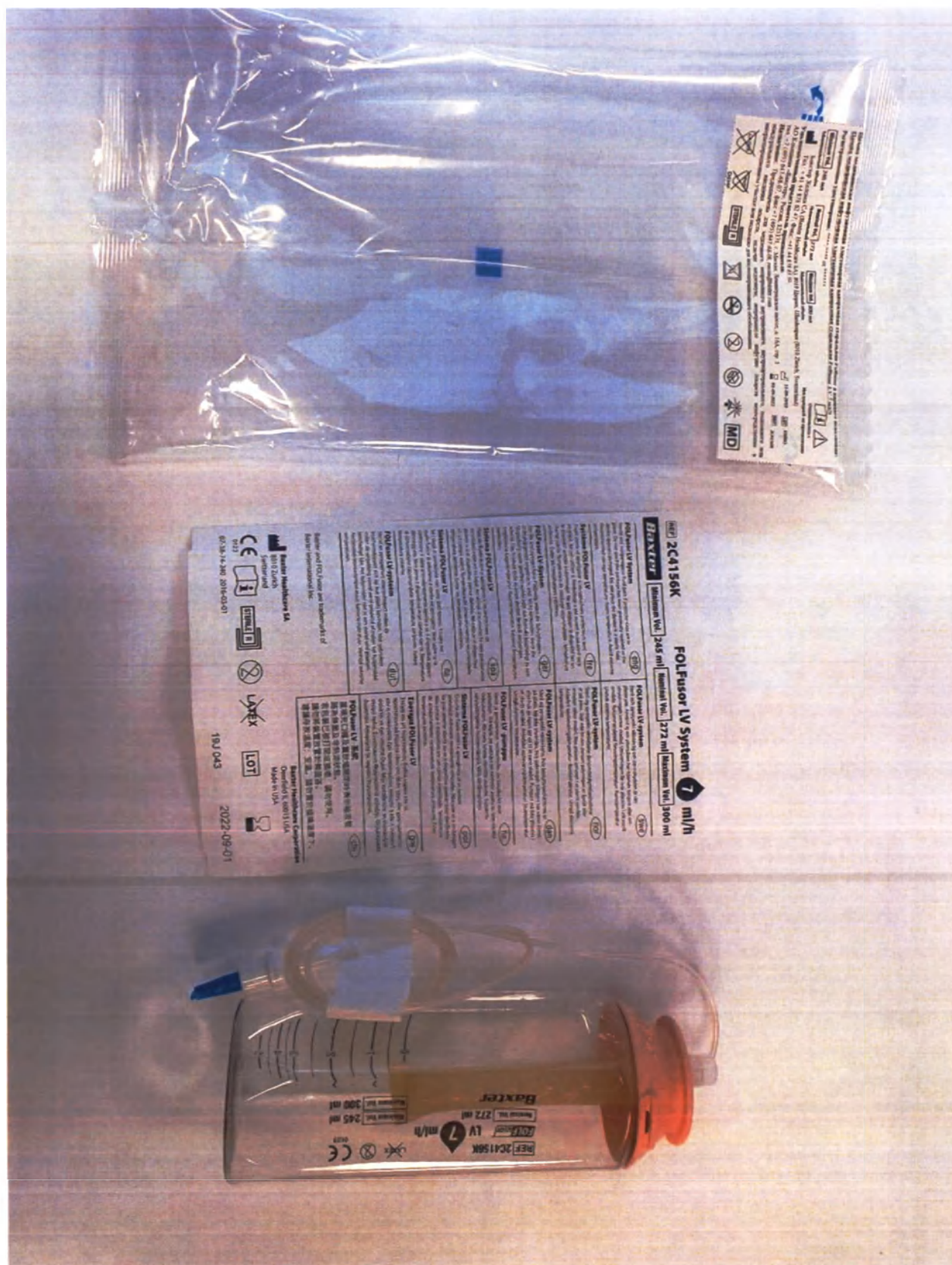
6.1. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 7 мл/ч; Индивидуальная упаковка, лицевой сторона, с оригинальной маркировкой



6.2. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 7 мл/ч; Индивидуальная упаковка, обратная сторона



6.3. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 7 мл/ч; Индивидуальная упаковка, лицевая сторона, с нанесенной маркировкой на русском языке



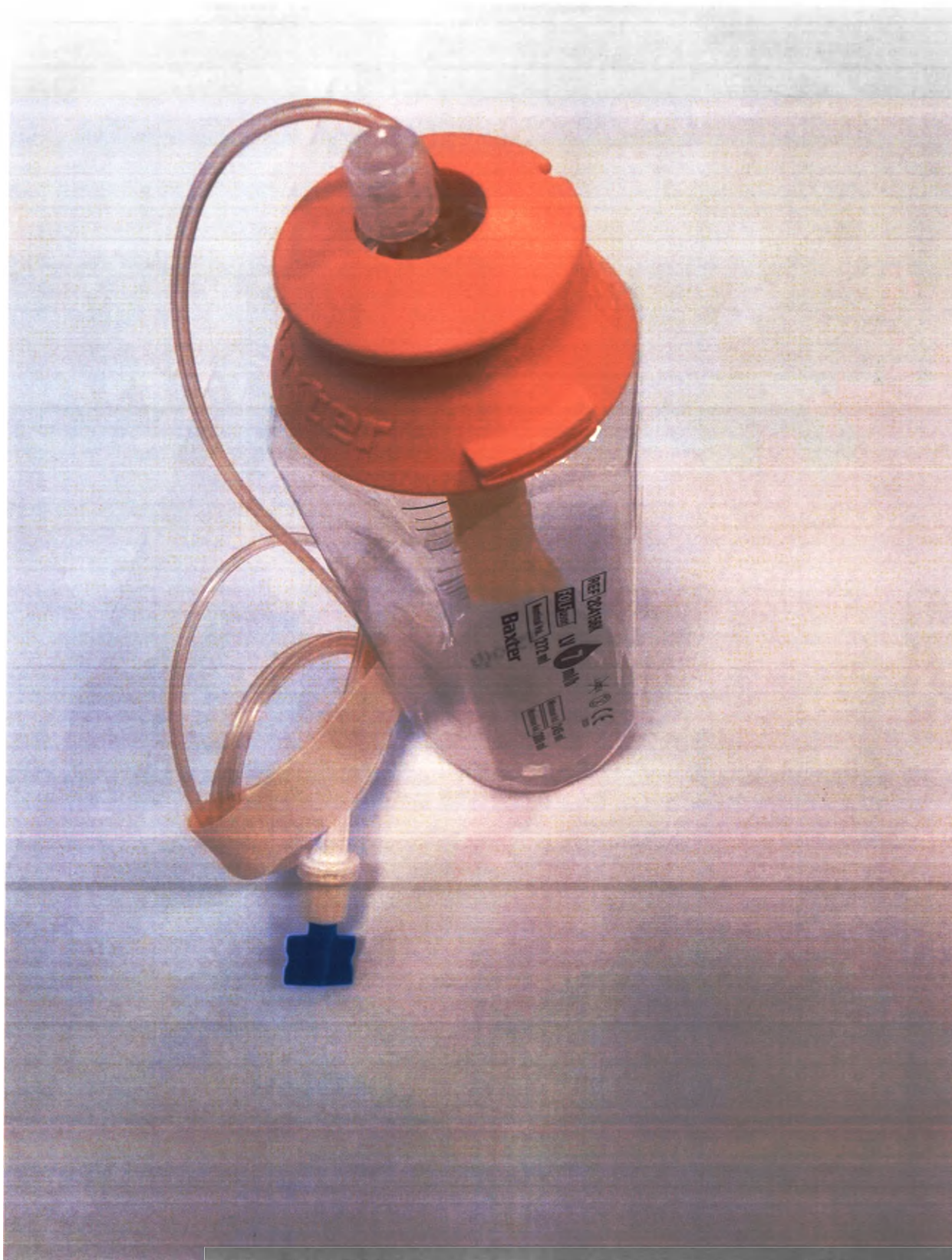
6.4. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 7 мл/ч; Комплект поставки



**6.5. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 7 мл/ч;
Помпа, лицевая сторона**



6.6. Помпа медицинская инфузионная эластомерная одноразовая стерильная Foliusor LV 7 мл/ч; Помпа, сторона со шкалой инфузии



6.7. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 7 мл/ч; Помпа, вид сверху

FOLFusor LV Portable Elastomeric Infusion System

[illegible]

*The device is designed to operate at nominal flow rate using 5% Dextrose (D5W). Use of diluents other than D5W will affect the flow rate of the device (see Operating Conditions).

The PCD user (5) system is an introductory, disposable device that uses an elastomer reservoir to store medication. When filled, the PCD flow rate can be adjusted with a contained internal pressure. Connections consist of two 3-particulate, 1/8-in. ID and 1/4-in. OD fittings.

The FOLIO and its system is designed to provide a continuous flow of medication over the course of 24 hours at the target flow rate with an accuracy of $\pm 1\%$ flow. The flow is continuously monitored and adjusted $\pm 1\%$ several times a minute to achieve just the right flow rate.

The following display may appear during the flow rate adjustment:

- (1) Flow rate, (2) Temperature, (3) Density after modification, (4) The difference in height between the FOLIO Port and the Client End Low Level Flow Sensor for use X, (5) Current set point for Operating Constraints.

The FOLIO system allows changing between two different calibration schemes in 20 sec. which include JGL, JVC, and most JALs.

This product is not manufactured using OTC plasticizer.
This product is not manufactured with natural rubber latex.

The Handed use of the FOLIO user system includes the following continuous functions: intra-portal administration, or up-down administration of Handedness. It may also include the following continuous functions of modification: directly first in intra-portal site or sub-portal use for postoperative pain management.

Epidural Administration
Epidural administration of analgesics is limited to use with infusing catheters specifically indicated for either short-term or long-term analgesic epidural drug delivery.

Its strongly recommended that a POL/PLU be created and to ensure regularity in their office duties, to carry out special duties and other administrative.

Warnings:

1. Do not fill with less than the minimum fill volume or more than the maximum fill volume (see product literature for details).

- [illegible]

The FCA Factor (Weight) used by the FCA in accordance with the procedure described under provisions for Rating and Selection

- Allow the fixed TQW assembly system to reach room temperature before using. Do not apply a heating source to bring the device to room temperature.

- It is the responsibility of the healthcare provider to instruct the patient and caregivers of the following:
- To achieve the best of medication management, ensure that the child always takes the medicine as directed by the doctor.

height during the infusion. Use of a coasting leg or coe stay with positioning of the wheel.
Check the PO/heel pressure is directed by your health care provider. If there is any problem with the PO/heel
or leg, contact your healthcare provider.

- [illegible]

Impact of PM Volume

- * The PCL-1000 is designed to operate at the nominal line rate with a maximum average current of 100mA. The PCL-1000 is not recommended for use in applications where the current is expected to exceed 100mA.

© 2000 Blackwell Science Ltd, *Journal of Internal Medicine* 247: 399–406

Do not fill with less than three inches of crushed gravel. The minimum fill volume (per gravel information table above, table 2) depends on the size of the gravel. The minimum fill volume will increase in volume and reduce its effectiveness.

The "2" from "2-phenyl" assigned is identical to the nominal flow rate when the molecular weight is 200,000. At 21.2°C (80°F) for 1.0 min and 21.2°C, and 21.2°C (80°F) for 20 min, 20 min, and 20 min. To achieve a desired molecular weight, the GPC SEC (all) (all) flow rate should be varied forward to the 21.2°C flow rate.

- Extended storage of the Web server in a refrigerative environment may result in unknown performance degradation.

• The TOL (or LV) system is compared to operate at the nominal rate using the DOL (or LV) as the flow. There is an acceptable 10% increase in the nominal flow rate if DOL (or LV) is not used as the flow.

- * The FCM-Late LV system is designed to operate at the maximal flow rate where the FCM Port and the Delay 2nd Line Lock/Time Restrictor are positioned at approximately the same height. Flow will decrease approximately 0.5% for every 1.5 mm (1 inch) the FCM Port is positioned below the cross 2nd Line Lock/Flow

Import or Collector's Use

* The FCM Fuel Air System is designed to operate at the nominal flow rate which occurs with a 22 gauge (3 mm) orifice. However, collector's use flow 22 mm (3/4 inch) orifice will produce the flow rate shown, table below.

- As a result of this attack, the use of the data in the system will lead to a reduction in the data.

ing and the information
to all our customers, storage users for they need to know and storage products.

Instructions for Filling and Priming

...failures to properly prime the basics may result in delayed change.

- Remove the paper and begin the loading.
 Remove the Starter/Producer Cap from the F20 Four V system. Replace the Starter/Producer Cap to replace when filling & complete.
 Remove the air from the air separator tank by draining it. Fill the separator tank with water. When possible, introduce the gas directly into the center of the water stream coming from the top.
 Load a small amount of catalyst or filling media into the F20 Port and measure. Do not over-charge to avoid damaging the F20 Port.
 Fill the F20 until a vacuum.
 Fill the F20 until a vacuum.
 Fill the F20 until a vacuum.

- [illegible]

Replace the Winged Low Cap with the Dye-Infused Low Cap. Always use the Winged Low Cap when the dye is not in use.

- Label the 1st & 2nd year of growth in accordance with appropriate vegetation and cover modifications only.
Place the final "Classification" growth in its position for transport to the pulpit.

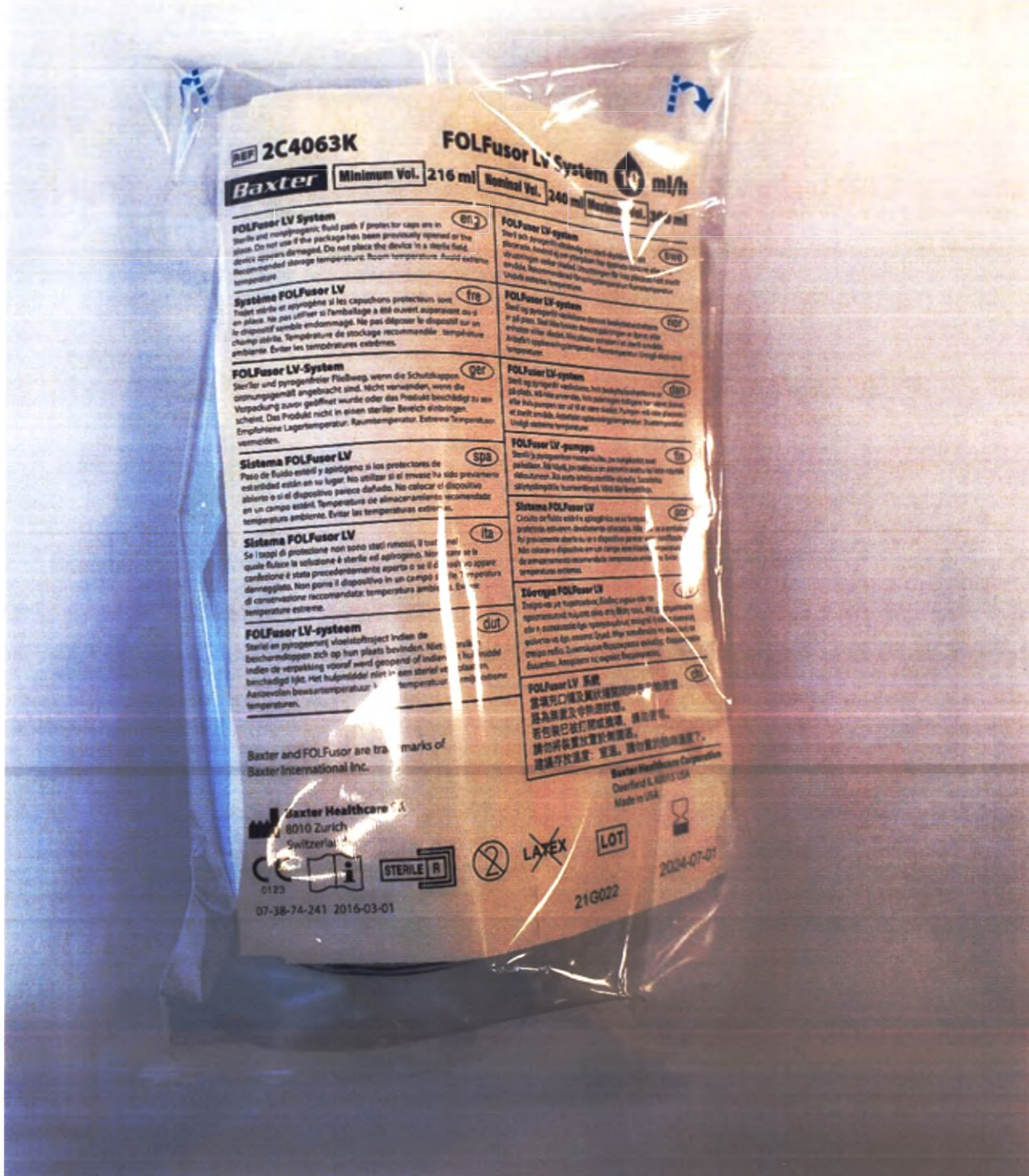
- To start the reaction, remove the stirrer/Lap Cap from the Media Reservoir/LapFlow Reservoir. Manually degas the media by slowly bubbling the culture media down the media reservoir. To degas the media, turn the Media Reservoir/LapFlow Reservoir before use.

To achieve the desired medication temperature, ensure that the Clearing Jug is at room temperature before use.

- Source: *Journal of the American Medical Association*, 2000; 284: 1033-1038.

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7. Помпа медицинская инфузионная эластомерная одноразовая стерильная
Folfusor LV 10 мл/ч



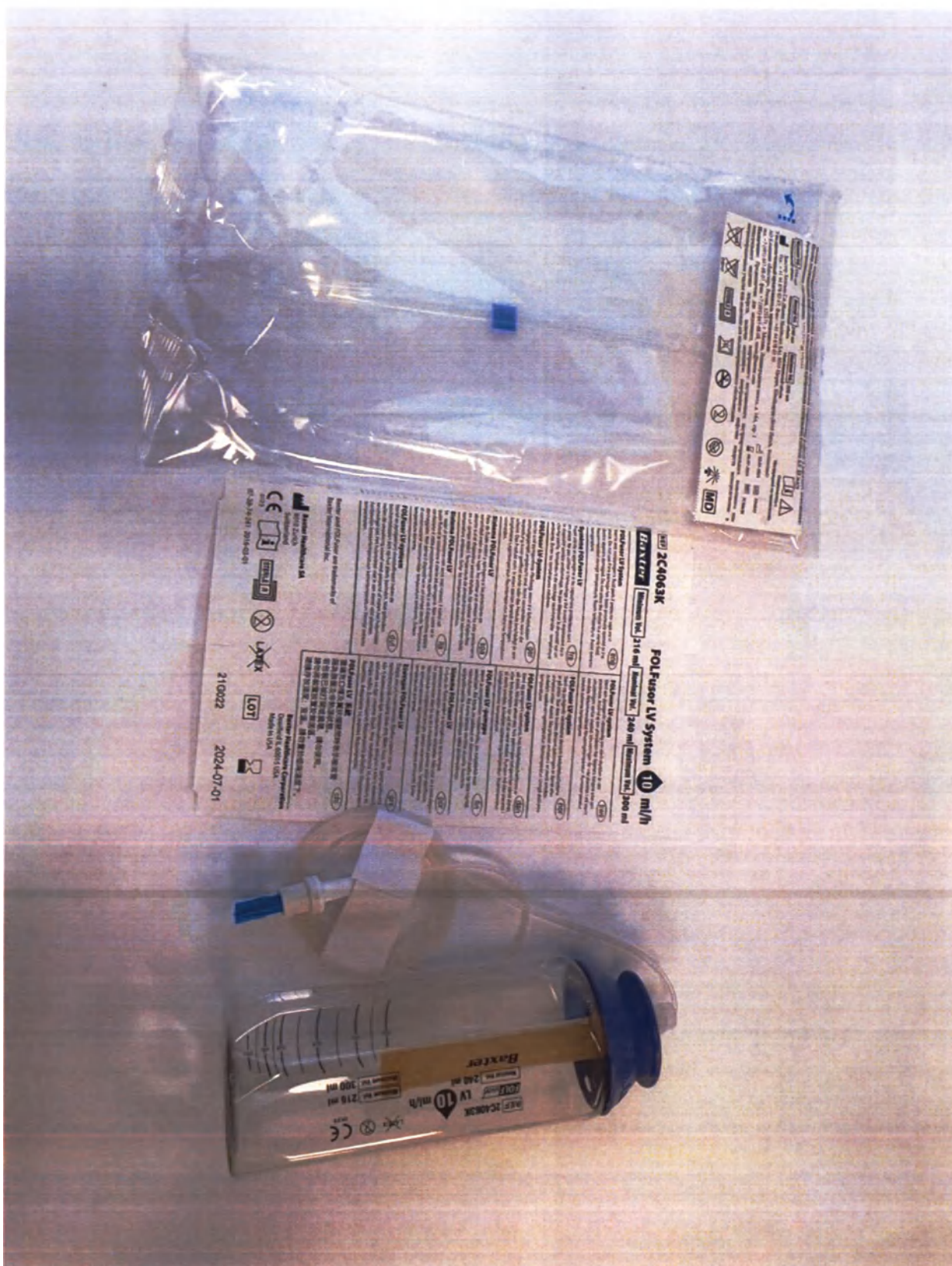
7.1. Помпа медицинская инфузионная эластомерная одноразовая стерильная FolFusor LV 10 мл/ч; Индивидуальная упаковка, лицевой сторона, с оригинальной маркировкой



**7.2. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 10 мл/ч;
Индивидуальная упаковка, оборотная сторона**



7.3. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 10 мл/ч; Индивидуальная упаковка, лицевая сторона, с нанесенной маркировкой на русском языке



7.4. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 10 мл/ч; Комплект поставки



7.5. Помпа медицинская инфузионная эластомерная одноразовая стерильная Folfusor LV 10 мл/ч; Помпа, лицевая сторона



**7.6. Помпа медицинская инфузионная эластомерная одноразовая стерильная Foliusor LV 10 мл/ч;
Помпа, сторона со шкалой инфузии**



7.7. Помпа медицинская инфузионная эластомерная одноразовая стерильная Foliusor LV 10 мл/ч;
Помпа, вид сверху

Принадлежности:

8. Сумка поясная для помпы



8.1. Сумка поясная для помпы

Индивидуальная упаковка, с нанесенной маркировкой на русском языке



8.2. Сумка поясная для помпы
Лицевая сторона



8.3. Сумка поясная для помпы
Оборотная сторона с крючком крепления к ремню



8.4. Сумка поясная для помпы
Вид сбоку



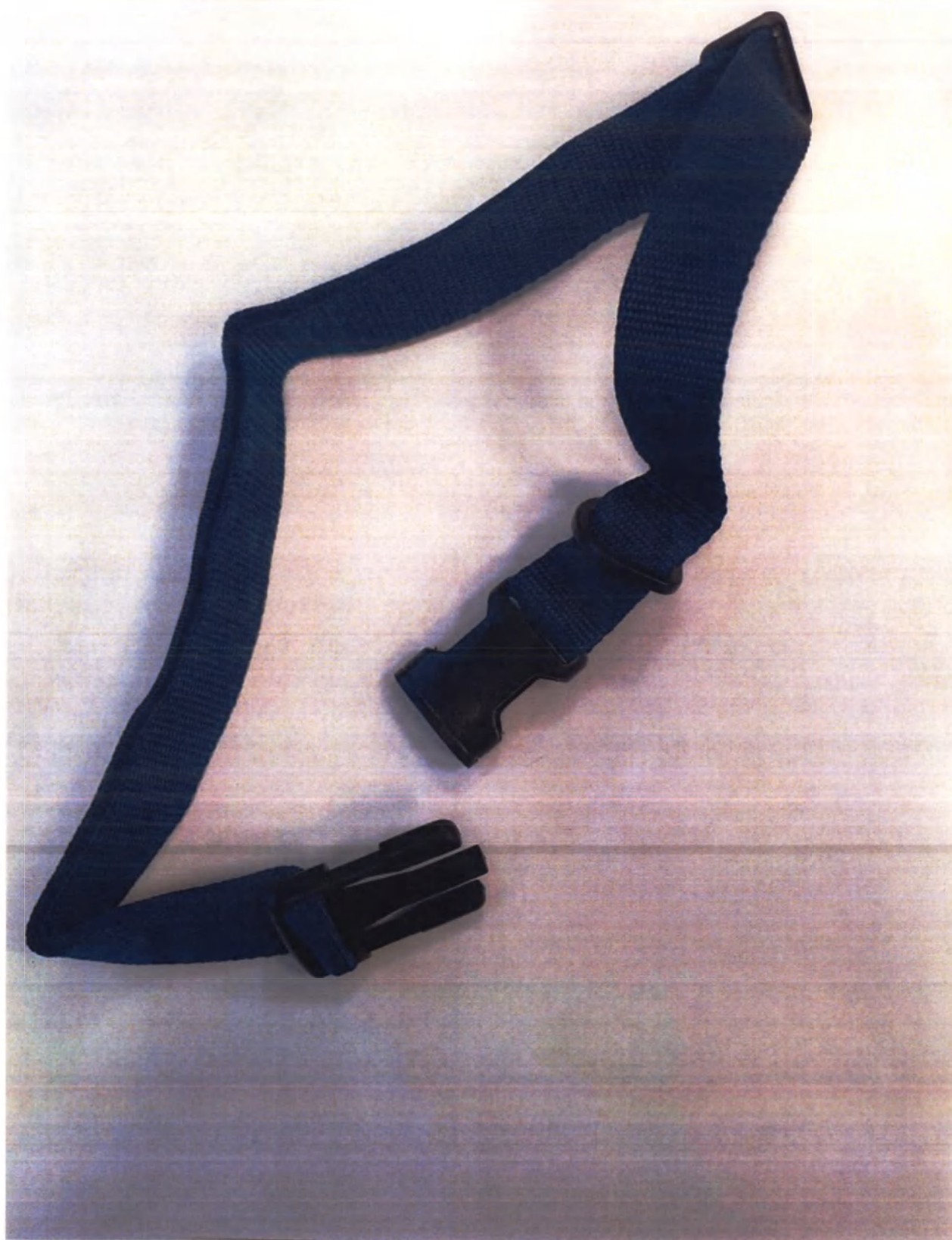
8.5. Сумка поясная для помпы
Крепление сумки



8.6. Сумка поясная для помпы
Внутри сумки хранится ремень для сумки



8.7. Сумка поясная для помпы
Сумка и ремень



8.8. Сумка поясная для помпы
Ремень



8.9. Сумка поясная для помпы
Вид сверху, сумка раскрыта



8.10. Сумка поясная для помпы
Сумка, пристегнутая к ремню, помпа внутри



8.11. Сумка поясная для помпы
Сумка с помпой внутри, вид сверху

Пример маркировки на русском языке:

Помпа медицинская инфузионная эластомерная одноканальная стерильная Falfasor в варианте исполнения:
Помпа медицинская инфузионная эластомерная одноканальная стерильная Falfasor LV 10 м.г/ч

Регистрационное удостоверение: **** от *****

Nominal Vol. 216 мл	Nominal Vol. 240 мл	Nominal Vol. 300 мл
Номинальный объем	Номинальный объем	Максимальный объем

Бакстер Хелскав СА (Baxter Healthcare SA), 8010 Zürich, Switzerland (Швейцария)
 Тел.: +41 44 878 62 47; Факс: +41 44 878 63 50.

Уполномоченный представитель производителя:
 Акционерное Общество Компания «Бакстер» / АО Компания «Бакстер»,
 125171, г. Москва, Ленинградское шоссе, д. 16А, стр. 1
 тел. +7 (495) 647-68-07, факс +7 (495) 647-68-08, russia@baxter.com

Назначение: Предназначена для медленного, непрерывного внутривенного, внутримышечного, подкожного или эпидурального введения лекарств, включая медленную, непрерывную инфузию лекарств непосредственно в интраоперационном участке или подкожно для постоперационного обезболивания

19-07-2021 LOT 210822
 01-07-2024 RDP 2C4063K

Сопроводитель с инструкцией по применению

Принадлежности:
Оушка полноразмерные для помпы

Регистрационное удостоверение: ****/***** от *****

Бакстер Хелскав СА (Baxter Healthcare SA), 8010 Zürich, Switzerland (Швейцария)
 Тел.: +41 44 878 62 47; Факс: +41 44 878 63 50.

Уполномоченный представитель производителя:
 АО Компания «Бакстер», Россия, 125171, г. Москва, Ленинградское шоссе, д. 16А, стр. 1
 тел. +7 (495) 647-68-07, факс +7 (495) 647-68-08, russia@baxter.com

25-02-2020 LOT 300046
 Срок годности - бесконечно RDP 2C1100

Сопроводитель с инструкцией по применению

прошито, пронумеровано и
скреплено печатью 77 листа(ов).

Руководитель отдела регистрации,
Россия и СНГ
АО Компания «Бакстер»

Гусев М.В.

« 17 » октября 2021 г.

