



VYJUVEK[®] Gel Preparation

Step-by-Step Instructions for VYJUVEK Gel Dose Preparation and Disposal

INDICATION AND USAGE

VYJUVEK is a topical gel indicated for the treatment of wounds in patients 6 months of age and older with dystrophic epidermolysis bullosa (DEB) with mutation(s) in the *collagen type VII (COL7A1)* gene.

Please see the **Important Safety Information** at the end of this document and <u>click here for **Full Prescribing Information**.</u>

Prepare the Administration Syringes

Disposal of Preparation Materials

Overview and Supplies

VYJUVEK® (beremagene geperpavec-svdt) gel should be prepared at a pharmacy by mixing VYJUVEK biological suspension into the excipient gel.

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uals who are pregnant should epare or apply VYJUVEK gel.

Preparation of VYJUVEK gel will take about 35 minutes



AT LEAST 20 minutes to thaw the VYJUVEK biological suspension and excipient gel vials **APPROXIMATELY 15 minutes to:**

- Mix the VYJUVEK biological suspension with the excipient gel to create the VYJUVEK gel
- · Prepare the administration syringes
- · Properly dispose of all preparation materials

VYJUVEK Gel Preparation Supplies

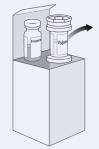
- One (1) carton containing (1) VYJUVEK biological suspension vial and (1) excipient gel vial (See Figure 1)
- Two (2) 18-gauge needles
- Two (2) to four (4) 1-mL administration syringes*
- One (1) 3-mL preparation syringe
- Two (2) to four (4) syringe caps*
- Protective gloves

- 70% isopropyl alcohol pads
- · Biohazard waste container
- Labels for administration syringes
- Virucidal agent—such as 70% isopropyl alcohol, 6% hydrogen peroxide, or <0.4% ammonium chloride-for clean-up
- Absorbent Materials (eq, paper towels)

*Based on the age of the patient. Six months to <3 years of age (up to 2 syringes each containing 0.4 mL VYJUVEK gel); ≥3 years of age (up to 4 syringes each containing 0.4 mL of VYJUVEK gel).

Figure 1.

Carton containing the VYJUVEK biological suspension vial and excipient gel vial





VYJUVEK Shelf-carton

VYJUVEK Biological Suspension Vial



Excipient Gel Vial

Shelf-cartons may be stored:

- In a freezer at -15°C to -25°C (5°F to -13°F)
- In a refrigerator for up to 1 month at 2°C to 8°C (35.6°F to 46.4°F), if a freezer is not available

VYJUVEK biological suspension is combined with the excipient gel to create VYJUVEK gel.

Prepare the Preparation Syringe

Step 1: Wash your hands and put on protective gloves.

Step 2: Thaw the vials.

Remove the VYJUVEK[®] (beremagene geperpavec-svdt) biological suspension and excipient gel vials from the shelf-carton and thaw at room temperature (20°C to 25°C [68°F to 77°F]) for **AT LEAST 20 minutes**.

Visually inspect both vials to ensure they are in liquid form and completely thawed. Excipient gel is more viscous and will take longer to thaw than the VYJUVEK biological suspension. Figure 2 shows a timelapse of the thawing of the excipient gel.

Once either the VYJUVEK biological suspension or the excipient gel is thawed, do not refreeze.

Step 3: Invert the VYJUVEK biological suspension vial 4–5 times.

DO NOT INVERT the excipient gel vial.

Step 4: Remove caps from vials.

Clean each vial stopper with a fresh 70% isopropyl alcohol pad. Allow them to dry.

Step 5: Connect preparation syringe to needle.

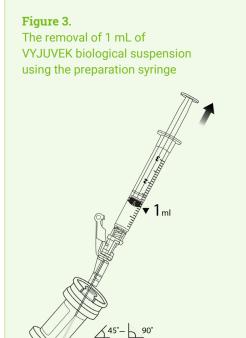
Aseptically connect an 18-gauge needle to the 3-mL preparation syringe.

Step 6: Remove the needle cap.

Puncture the VYJUVEK biological suspension vial stopper.

Step 7: Withdraw 1 mL of VYJUVEK biological suspension.

Hold the VYJUVEK biological suspension vial at a **45° to 90° angle**, as pictured in Figure 3, while pulling the plunger away from the vial.





10 Min

0 Min

Figure 2.

20 Min

Steps 8–15 continue on the next page.

Step 8: Remove the preparation syringe from the vial.

Preparation syringe (still connected to needle) contains 1 mL of VYJUVEK[®] (beremagene geperpavec-svdt) biological suspension from the vial. **DO NOT engage the safety lock**.

Step 9: Discard the VYJUVEK biological suspension vial in the biohazard waste container.

Step 10: Puncture the excipient gel stopper.

Transfer 1 mL of VYJUVEK biological suspension into the excipient gel vial, as illustrated in **Figure 4**.

Step 11: Remove air.

Without removing the needle from the excipient gel vial, lift the bevel of the needle above the liquid, as highlighted in Figure 5.

Do not remove any liquid.

Pull the plunger back to the 1 mL mark to remove air, as shown in Figure 6.

Step 12: Remove the preparation syringe from the excipient gel vial.

Remove the preparation syringe with the 1 mL of air and engage the safety lock.

Step 13: Discard the preparation syringe and needle into the biohazard waste container.

Step 14: Cover excipient gel stopper with an alcohol pad.

Place a 70% isopropyl alcohol pad on top of the excipient gel stopper and hold it tightly in place.

Step 15: Shake VYJUVEK gel mixture.

Hold excipient gel vial with 70% isopropyl alcohol pad between index finger and thumb, as shown in Figure 7, and manually shake VIGOROUSLY for 10 seconds.



Figure 5. The needle abo

The needle above the liquid without removal of any material Figure 6. Pulling the plunger back to the 1 mL mark to remove air





Figure 7. Manually mixing the excipient gel vial



Mixture of VYJUVEK biological suspension and excipient gel is referred to as VYJUVEK gel.

Please click here to access the Important Safety Information and Full Prescribing Information.

Prepare the Administration Syringes

Step 1: Connect needle to administration syringe.

Aseptically connect an 18-gauge needle to the first 1-mL administration syringe and remove the needle cap.

Step 2: Insert needle into excipient vial.

Insert the 18-gauge needle into the excipient vial containing VYJUVEK[®] (beremagene geperpavec-svdt) gel. Figure 8 shows the inserted needle prior to withdrawal.

Step 3: Withdraw 0.4 mL of VYJUVEK gel.

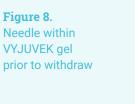
Tilt the excipient vial to a 45° to 90° angle as pictured in **Figure 9**.

An air pocket may form near the plunger when extracting VYJUVEK gel.

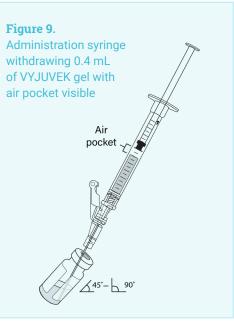
Step 4: Disconnect administration syringe containing 0.4 mL of mixed VYJUVEK gel.

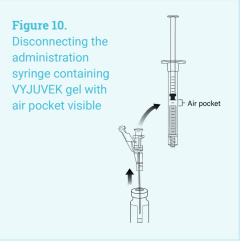
DO NOT REMOVE THE NEEDLE FROM the excipient gel vial stopper. Lift the bevel of the needle above the VYJUVEK gel and disconnect the administration syringe, leaving the needle in the excipient gel vial stopper, as illustrated in **Figure 10**.

Steps 5–7 continue on the next page.









Prepare the Preparation Syringe

Step 5: Remove air pockets from administration syringe.

DO NOT FLICK the syringe to remove the air pocket.

Manipulate the plunger up and down, until all air pockets have been removed, as highlighted in **Figure 11**.

Step 6: Cap the administration syringe.

Once capped, as shown in **Figure 12**, put an appropriate label on the administration syringe and set aside.

Note: Administration syringes are labeled as #1, #2, #3, and #4.

Step 7: Connect a new administration syringe to needle remaining in the excipient gel vial containing VYJUVEK[®] (beremagene geperpavecsvdt) gel.

Connect with the bevel of the needle **ABOVE** the VYJUVEK gel, as pictured in Figure 13.

Repeat Steps 3–7 until you have prepared the appropriate number of administration syringes based on age of patient.



Prepare up to 2 syringes if the patient is aged 6 months to <3 years (total volume = 0.8 mL)

Prepare up to 4 syringes if the patient is ≥3 years of age (total volume = 1.6 mL)

Place the capped administration syringes containing the VYJUVEK gel in a sealable plastic bag. Place the sealable plastic bag with the administration syringes into an appropriate insulated secondary container at 2°C to 8°C (35.6°F to 46.4°F) for transport from the preparation site to the administration site.

For questions around storage and handling of prepared VYJVUEK gel syringes, contact Krystal Biotech at 1-844-5-KRYSTAL. Discard any used or unused administration syringes in a biohazard waste container.

Figure 11. Manipulation of administration syringe plunger to remove air pockets

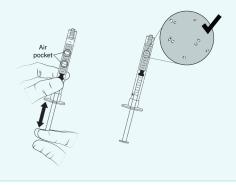
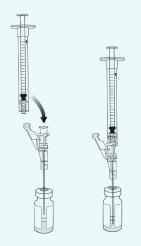




Figure 13.

Connecting an administration syringe to the needle within the excipient gel vial stopper



Disposal of Preparation Materials

Step 1: Dispose of all materials used to prepare administration syringes.

Discard the excipient gel vial (with the needle within the vial stopper) into the biohazard waste container.

Step 2: Clean all surfaces and treat all VYJUVEK® (beremagene geperpavec-svdt) spills.

Any surface that may have come in contact with the VYJUVEK biological suspension or gel should be cleaned. Treat all VYJUVEK spills with a virucidal agent such as 70% isopropyl alcohol, 6% hydrogen peroxide, or <0.4% ammonium chloride. Blot using absorbent materials.

Step 3: Dispose of used absorbent materials and protective gloves.

The used absorbent materials and protective gloves should be discarded in a biohazard waste container.

In the event of an accidental exposure to VYJUVEK (eg, through a splash to the eyes or mucous membranes), flush the area with clean water for at least 15 minutes.

IMPORTANT SAFETY INFORMATION

WARNINGS AND PRECAUTIONS

Accidental Exposure to VYJUVEK gel: VYJUVEK will not replicate in the subject's cells and does not integrate into the subject cells' native genetic material. For precautions, avoid direct contact with treated wounds (eg, touching or scratching) and dressings of treated wounds for approximately 24 hours following treatment. Wear protective gloves when assisting subjects with changing wound dressings and handling the disposal. In the event of accidental exposure (eg, through a splash to the eyes or mucous membranes), flush with clean water for at least 15 minutes.

Clean all surfaces that may have come in contact with VYJUVEK biological suspension or gel and treat all spills with a virucidal agent.

Dispose all materials (eg, vial, syringe, needle, cleaning materials) that may have come in contact with VYJUVEK biological suspension or gel into a biohazard bag or container.

ADVERSE REACTIONS

The most common adverse reactions (>5%) were itching, chills, redness, rash, cough, and runny nose.

Please click here for Full Prescribing Information.



Please click here to access the **Important Safety Information** and **Full Prescribing Information**.

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