Neulasta® (pegfilgrastim) injection Onpro® kit

NURSE CHECKLIST

This nurse checklist does not replace the Neulasta® Onpro® Instructions For Use (IFU) which should be reviewed prior to application.

The Patient IFU should be referred to when discussing Neulasta® Onpro® application with the patient. **Please provide the patient with the Neulasta® Onpro® Patient IFU (included in the kit).** You may also provide the patient with a Wallet Card. The patient may carry the Wallet Card as it also is a reminder for time of Neulasta® dose delivery the following day as well as other helpful information.

Please be aware that Wallet Card is not intended to replace the Neulasta® Onpro® Patient IFU.

1 CONVERSATION with the Patient (on-body injector is for adult patients only)

- ☐ Ask the patient what their activities look like the following day.
- ☐ Inform the patient that a caregiver should be present if the **on-body injector** is placed on the arm.
- ☐ Patients should NOT expose the **on-body injector** to direct sunlight. If the **on-body injector** is exposed to direct sunlight for more than 1 hour, it may affect your medicine. Patients should wear the **on-body injector** under clothing.
- ☐ Ask the patient to avoid tight clothing around the **on-body injector** and advise not to dislodge when removing clothing. The patient should call their healthcare provider immediately if the **on-body injector** becomes dislodged, a red light flashes, or if the **on-body injector** is noticeably wet (saturated), as they may need a replacement dose. Do not use other materials to hold it in place that could cover audio/visual indicators or compress the **on-body injector** against the patient's skin, as this could dislodge the cannula and lead to a missed or incomplete dose of Neulasta®.
- ☐ Patient should avoid getting body lotions, creams, oils or cleaning agents near the **on-body injector** as these products may loosen the adhesive. Before your next scheduled Neulasta® dose, avoid use of lotions, creams, or oils on your arms and stomach area (abdomen).
- ☐ Confirm with the patient that they are not planning on receiving radiation, CT scan, MRI, ultrasound or any other imaging tests or procedures while the **on-body injector** is on their body.
- ☐ Be sure the patient knows the day/time Neulasta® dose delivery is expected to start.
- ☐ Remind patients that the **on-body-injector** is on their body for the next 27 hours and they SHOULD NOT peel off, disturb, or bump the **on-body injector** before their full dose is complete. This may result in a missed or incomplete dose of Neulasta®. They should call your healthcare provider immediately if this happens.
- ☐ Tell the patient not to sleep on the **on-body injector** or apply pressure during wearing, especially during dose delivery. This may affect **on-body injector** performance.
- ☐ Patient should keep the **on-body injector** dry the last 3 hours prior to dose delivery start. They can immerse the **on-body injector** up to hour 24 as it is waterproof up to 8 feet for 1 hour.
- ☐ Patient should keep the **on-body injector** 4 inches away from electrical equipment such as cell phones, cordless phones, and common household appliances including microwave.
- ☐ If the patient plans to fly, they should ask the TSA agent for a manual pat-down.
- ☐ Patient should avoid activities and places that may interfere with monitoring during the dosing of Neulasta® administered by the **on-body injector** (hours 26-29) such as traveling, driving or operating heavy machinery.
- ☐ Advise the patient not to use a bath tub, hot tub, whirlpool or sauna while wearing the **on-body injector**. The **on-body injector** should only be exposed to temperatures between 41-104° Fahrenheit.
- ☐ Educate the patient that if they have an allergic reaction, they should remove the **on-body injector** by grabbing the edge of the adhesive pad and peeling off the **on-body injector**. Instruct the patient to get emergency medical help right away.
- ☐ Inform the patient that the **on-body injector** will produce a series of beeps after about 27 hours to signal patients that dose delivery is about to begin. Dose delivery will take about 45 minutes to complete. **Patients should not remove the on-body injector at this time.**
- ☐ Advise patients that a long beep will sound and the status light will be SOLID GREEN when dose delivery is complete.

  Patients can remove the **on-body injector** when the status light is SOLID GREEN or has switched off, and the fill indicator is EMPTY.

Please note: The prefilled syringe co-packaged in the **Neulasta® Onpro® kit** contains additional solution to compensate for liquid loss during delivery through the **on-body injector**. If this syringe is used for manual subcutaneous injection, the patient will receive an overdose. If the prefilled syringe for manual use is used with the **on-body injector**, the patient may receive less than the recommended dose.

Neulasta® Onpro®
Nurse Checklist

The information provided here is an abbreviated summary and does not replace the Instructions for Use, please consult the Healthcare Provider Instructions for Use for information in the kit prior to using the Neulasta® Onpro® kit.

2 PREPARE the Application Site

- ☐ When selecting application site, confirm with the patient there are no tight fits between the **on-body injector** and clothing to minimize the chance of dislodging the **on-body injector**.
- ☐ Thoroughly clean the site with alcohol to enhance **on-body injector** adherence to the skin. **Only** use alcohol to clean the skin.

3 FILL the On-Body Injector

- ☐ Remove air bubbles from Neulasta® prefilled syringe included in the Neulasta® Onpro® kit without expelling medicine. Injecting air bubbles into the on-body injector could interfere with the full-dose delivery.
- ☐ Center the needle directly over the medicine port at a 90° angle. Insert all the way into the port, avoiding sides.
- ☐ Remove blue needle cover from back of **on-body injector** only AFTER the **on-body injector** is filled.

4 CONFIRM On-Body Injector Activation

- ☐ Ensure amber light flashes and the fill indicator is at FULL, you'll have 3 minutes to apply to patient. Cannula will deploy in 3 minutes whether **on-body injector** is on the patient or not.

5 APPLY the On-Body Injector

- ☐ Grasp the **on-body injector's** plastic case with your fingertips and only by sides, keeping fingers off of the adhesive.
- ☐ Securely apply the **on-body injector** without bending, folding, wrinkling or curling the adhesive. **Important:** Once on the skin, **press firmly on the on-body injector** to ensure proper adhesion to the patient's skin.
- ☐ If additional adhesion is deemed appropriate, an adhesive extender that fits around the **on-body injector** can be obtained by calling 1-844-MYNEULASTA (1-844-696-3852).
  - **Do not** use other materials to secure the **on-body injector** to the patient that could cover audio/visual indicators or compress the **on-body injector** against the patient's skin.
- ☐ If the adhesive is wrinkled in front of the cannula window or has folds anywhere that prevent the **on-body injector** from securely adhering, remove the **on-body injector**. Start again with a new kit and call Amgen at 1-800-772-6436 or your Amgen representative.
- ☐ Be sure a free Sharps container is ordered. Visit NeulastaOnpro.com/DisposalContainer or call 1-844-MYNEULASTA.

Advise patient to **NOT** pull off device **EARLY**:

☐ Listen for the long beep ☐ SOLID GREEN light ☐ Fill indicator is EMPTY

If patient is unsure their dose of Neulasta® delivered properly via the **on-body injector**, patient should immediately call their healthcare provider.

For more information, go to NeulastaHCP.com or call 1-844-696-3852 (1-844-MYNEULASTA).

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