Neulasta® is a prescription medicine used to help reduce the chance of infection due to a low white blood cell count, in people with certain types of cancer (non-myeloid), who receive anti-cancer medicines (chemotherapy) that can cause fever and low blood cell count.

If the On-body Injector was placed on the back of your arm, a caregiver must be available to monitor the On-body Injector.

Please see Important Safety Information on pages 10-11.
About your cancer care

Being told you have cancer can feel overwhelming—you’ll have lots of new information to understand, appointments to keep, and decisions to make. Your healthcare team is there to help support and guide you throughout your journey, and to help you get the treatment option that is right for you.

Chemotherapy, or “chemo,” is one of those treatment options—it is given to help stop or slow down the growth of cancer, which is caused when cells in your body grow out of control.¹,²

The strong chemotherapy your doctor prescribed may reduce the number of white blood cells in your body.³ White blood cells are a key part of your immune system—your body’s natural defenses against infection. As your white blood cell count goes down, your risk of infection goes up.⁴

Important Safety Information

Do not take Neulasta® if you have had a serious allergic reaction to Neulasta® (pegfilgrastim) or NEUPOGEN® (filgrastim).

Please see additional Important Safety Information on pages 10-11.
Neulasta® is a prescription medication used to help protect against the risk of infection (initially marked by fever) in patients with some tumors receiving strong chemotherapy that decreases the number of infection-fighting white blood cells.\(^5\)

Neulasta® is similar to a protein made by your body and is sometimes called a white blood cell booster because it causes your body to make more white blood cells. This helps increase your natural defenses against infection.\(^5,6\)

Neulasta® is given once for each cycle of chemotherapy.

Neulasta® should only be administered on the day your healthcare provider has determined and should not be delivered until approximately 24 hours after receiving chemotherapy.

### Important Safety Information

Tell your healthcare provider if you:

- Have sickle cell trait or sickle cell disease
- Have had severe skin reactions to acrylic adhesives
- Are allergic to latex
- Have problems with your kidneys
- Have any other medical problems
- Are pregnant or plan to become pregnant. It is not known if Neulasta® may harm your unborn baby
- Are breastfeeding or plan to breastfeed. It is not known if Neulasta® passes into your breast milk

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements.

Please see additional Important Safety Information on pages 10-11.
The Neulasta® Onpro® is a kit that includes a single dose of Neulasta and a single-use On-body Injector, which your healthcare provider activates by filling with your Neulasta® dose. Your healthcare provider then applies the On-body Injector to your skin, either on the back of the arm or abdomen. It’s applied the same day you have chemotherapy, and you leave the doctor’s office wearing the On-body Injector.

The On-body Injector has been designed to deliver your Neulasta® approximately 27 hours after your healthcare provider has applied it on your skin.

Important Information

While the On-body Injector is in place you should avoid:

- Traveling, driving or operating heavy machinery during hour 26 through hour 29 after the On-body Injector is applied.
- Sleeping on the On-body Injector or applying pressure on the On-body Injector. The On-body Injector may not work properly.
- Bumping the On-body Injector or knocking it off your body.
- Getting body lotion, creams, oils, and skin cleansing products near the On-body Injector. These products may loosen the adhesive that holds the On-body Injector onto your body.
- Using hot tubs, whirlpools, or saunas, and direct sunlight. These may affect Neulasta®.
- Peeling off or disturbing the On-body Injector adhesive before you receive your full dose of Neulasta®.

Please see pages 8-9 for additional important information about the On-body Injector.
Understanding the On-body Injector

This summary does not replace the Patient Instructions for Use. It is important that you review the Patient Instructions for Use that came with the On-body Injector and contact your healthcare provider if you have any questions.

Flashing green
On-body Injector is working properly

Solid green (or turned off)
Medication delivery should be complete. Check to see if fill indicator reads “empty” and there is no noticeably wet adhesive

Flashing red
On-body Injector error—call your healthcare provider immediately

Status light
Adhesive pad
Fill indicator

Audio
On-body Injector will beep before and after dose delivery.

Cannula window
Allows you to view the cannula (a short, soft tube) that your Neulasta® passes through during the 45-minute dose delivery.

Actual size.

Please see Important Safety Information for Neulasta® on pages 10-11.
At the doctor’s office:

The day that the On-body Injector (pegfilgrastim) is applied

On the day of your chemotherapy, your healthcare provider will prepare an area of skin and apply the On-body Injector.

Abdomen  Back of upper arm

What happens immediately after the On-body Injector is applied?

Once the On-body Injector is placed on your skin, it will beep and the yellow light will flash. A short needle will insert the cannula, which is designed to allow Neulasta® to be delivered under your skin 27 hours later. The needle is designed to go back into the On-body Injector, but the cannula will remain in place until the On-body Injector is removed.

For the next 27 hours, the green light will flash every 5 seconds. This means the On-body Injector is working properly.

Important Information

Keep the On-body Injector at least 4 inches away from electrical equipment such as cell phones, cordless telephones, microwaves and other common appliances.

If the On-body Injector is too close to electrical equipment, it may not work correctly and can lead to a missed or incomplete dose of Neulasta®.

Please see pages 8-9 for additional important information about the On-body Injector.

Please see Important Safety Information for Neulasta® on pages 10-11.
At home or other setting:
The day of Neulasta® delivery

On the day of Neulasta® delivery, you will be responsible for the following (as described in detail in the Patient Instructions for Use):

It is important that you thoroughly review the Patient Instructions For Use. These instructions cover everything you need to know about the On-body Injector. The information in this brochure does not replace the Patient Instructions For Use. Please contact your healthcare provider if you have any questions.

- **Check the status light occasionally.** It should be flashing green for about 27 hours.
  
  If you hear beeping, check the status light. If the light is flashing RED, call your healthcare provider immediately.

- **Monitor the dose delivery.** Remember:
  
  - A caregiver should be with you the first time that you receive Neulasta® with the On-body Injector
  
  - If the On-body Injector was placed on the back of your arm, always have a caregiver available to monitor the On-body Injector

- **Remove the On-body Injector.**

- **Confirm the dose delivery.**

- **Dispose of the On-body Injector** according to local laws and regulations to avoid accidental needle stick. To participate in Amgen’s voluntary disposal program, please fill out and send the mailing card attached to the right of this page. You may also call 1-844-MYNEULASTA (1-844-696-3852) or visit www.NeulastaOnpro.com to enroll.
What can I expect when receiving Neulasta® with the On-body Injector?5

- See the Instructions for Use for the On-body Injector for information about the On-body Injector.
  - Know the time that delivery of your dose of Neulasta® is expected to start.
  - Avoid traveling, driving, or operating heavy machinery during hour 26 through hour 29 after the On-body Injector is applied. Avoid activities and places that may interfere with monitoring during the 45-minute period that Neulasta® is expected to be delivered by the On-body Injector, and for 1 hour after delivery.
- A caregiver should be with you the first time that you receive Neulasta® with the On-body Injector.
- If you have an allergic reaction during the delivery of Neulasta®, remove the On-body Injector by grabbing the edge of the adhesive pad and peeling off the On-body Injector. Get emergency medical help right away.

- You should only receive a dose of Neulasta® on the day your healthcare provider tells you.
- You should not receive your dose of Neulasta® any sooner than 24 hours after you finish receiving your chemotherapy. The On-body Injector is programmed to deliver your dose about 27 hours after your healthcare provider places the On-body Injector on your skin.
- Do not expose the On-body Injector to the following because the On-body Injector may be damaged and you could be injured:
  - MRI
  - X-ray
  - CT scan
  - Ultrasound
  - Oxygen-rich environments, such as hyperbaric chambers
• Avoid airport X-ray scans. Request a manual pat down instead. Use care during a manual pat down to help prevent the On-body Injector from being accidentally removed.

• **Keep the On-body Injector at least 4 inches away from electrical equipment such as cell phones, cordless telephones, microwaves and other common appliances.** If the On-body Injector is too close to electrical equipment, it may not work correctly and can lead to a missed or incomplete dose of Neulasta®.

• The On-body Injector is for adult patients only.

• **Call your healthcare provider right away if the:**
  – On-body Injector comes off before or during a dose delivery. **Do not re-apply it.**
  – On-body Injector is leaking.
  – Adhesive on your On-body Injector becomes noticeably wet (saturated) with fluid, or there is dripping. This may mean that Neulasta® is leaking out of your On-body Injector. If this happens you may only receive some of your dose of Neulasta®, or you may not receive a dose at all.
  – On-body Injector status light is flashing red.

Please review the Patient Instructions for Use for instructions and information about the On-body Injector. Discuss any questions you have with your healthcare provider. The information in this guide is intended as a summary. It is not intended to replace any instructions from your healthcare provider or the Instructions For Use which came packaged with the On-body Injector.

Please see Important Safety Information on pages 10-11.
Who should not take Neulasta®?
Do not take Neulasta® if you have had a serious allergic reaction to Neulasta® (pegfilgrastim) or NEUPOGEN® (filgrastim).

What should I tell my health care provider before taking Neulasta®? Tell your healthcare provider if you:
• Have sickle cell trait or sickle cell disease
• Have had severe skin reactions to acrylic adhesives
• Are allergic to latex
• Have problems with your kidneys
• Have any other medical problems
• Are pregnant or plan to become pregnant. It is not known if Neulasta® may harm your unborn baby
• Are breastfeeding or plan to breastfeed. It is not known if Neulasta® passes into your breast milk

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements.

What are possible serious side effects of Neulasta®?
• Spleen Rupture. Your spleen may become enlarged and can rupture while taking Neulasta®. A ruptured spleen can cause death. Call your doctor right away if you have pain in the left upper stomach area or left shoulder tip area. This pain could mean your spleen is enlarged or ruptured.

• A serious lung problem called Acute Respiratory Distress Syndrome (ARDS). Call your healthcare provider or get emergency medical help right away if you get any of these symptoms of ARDS: fever, shortness of breath, trouble breathing, or a fast rate of breathing.

• Serious Allergic Reactions. Get emergency medical help right away if you get any of these symptoms of a serious allergic reaction with Neulasta®: shortness of breath, wheezing, dizziness, swelling around the mouth or eyes, fast pulse, sweating, and hives.

If you have an allergic reaction during the delivery of Neulasta®, remove the On-body Injector by grabbing the edge of the adhesive pad and peeling off the On-body Injector. Get emergency medical help right away.
• **Sickle Cell Crises.** Severe sickle cell crises, and sometimes death, can happen in people with sickle cell trait or disease who receive filgrastim, a medicine similar to Neulasta®.

• **Kidney injury (glomerulonephritis).** Kidney injury has been seen in patients who received Neulasta®. You should notify your healthcare provider right away if you experience puffiness in your face or ankles, blood in your urine or brown colored urine or you notice you urinate less than usual.

• **Increased white blood cell count (leukocytosis).** Your doctor will check your blood during treatment with Neulasta®.

• **Capillary Leak Syndrome.** Neulasta® can cause fluid to leak from blood vessels into your body’s tissues. This condition is called “Capillary Leak Syndrome” (CLS). CLS can quickly cause you to have symptoms that may become life-threatening. Get emergency medical help right away if you develop any of the following symptoms:
  - swelling or puffiness and are urinating less often
  - trouble breathing
  - swelling of your stomach-area (abdomen) and feeling of fullness
  - dizziness or feeling faint
  - a general feeling of tiredness

The most common side effect of Neulasta® is pain in the bones and in your arms and legs. Tell your healthcare provider if you have any side effect that bothers you or that does not go away. These are not all the possible side effects of Neulasta®. Call your doctor for medical advice about side effects. You may report negative side effects to the FDA at 1-800-FDA-1088 (1-800-332-1088).

For more information about Neulasta®, talk with your healthcare provider or pharmacist; go to www.NeulastaOnpro.com, or call 1-844-696-3852 (1-844-MYNEULASTA).

Please see accompanying Neulasta® Patient Information.

**References:**
Because you’d rather be home

The Neulasta® Onpro®
includes a single dose of Neulasta® and a single-use On-body Injector

• Applied the same day as chemotherapy is administered6
• Designed to automatically administer Neulasta®
  the next day (approximately 27 hours after application)6

Visit www.NeulastaOnpro.com to learn more about
Neulasta® Onpro® or call 1-844-MYNEULASTA (1-844-696-3852).

• The most common side effect of Neulasta® is pain in the
  bones and in your arms and legs.

If you are traveling by plane while wearing the On-body Injector for Neulasta®
download a copy of the TSA form at NeulastaOnpro.com/TSA.

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