

KEYTRUDA®

(pembrolizumab)

May cause some serious side effects which can sometimes become life-threatening and lead to death. These may happen any time during treatment or even after your treatment has ended. You may experience more than one side effect at the same time.

Contact your doctor right away if you develop any signs or symptoms including those not listed on this card. Your doctor may give you other medicines in order to prevent more severe complications and reduce your symptoms. Your doctor may withhold the next dose of KEYTRUDA or stop treatment with KEYTRUDA.

Lungs

- Shortness of breath
- Chest pain
- Coughing

Intestines

- Diarrhea or more bowel movements than usual
- Stools that are black, tarry, sticky, or contain blood or mucus
- Severe stomach pain or tenderness
- Nausea or vomiting

Liver

- Nausea or vomiting
- Feeling less hungry
- Pain on the right side of stomach
- Yellowing of skin or whites of eyes
- Dark urine
- Bleeding or bruising more easily than normal

Kidneys

- Changes in the amount or color of your urine

Type 1 diabetes, including diabetic ketoacidosis

- Feeling more hungry or thirsty
- Needing to urinate more often
- Weight loss
- Feeling tired or feeling sick
- Stomach pain
- Fast and deep breathing
- Confusion
- Unusual sleepiness
- A sweet smell to your breath
- A sweet or metallic taste in your mouth
- A different odour to your urine or sweat

Hormone glands

- Rapid heartbeat
- Weight loss or weight gain
- Increased sweating
- Hair loss
- Feeling cold
- Constipation
- Deeper voice
- Muscle aches
- Dizziness or fainting
- Headaches that will not go away or unusual headache

IMPORTANT

- Do not attempt to diagnose or treat side effects yourself.
- Do not stop your treatment with KEYTRUDA unless you have discussed this with your doctor
- **Take this card with you at all times**, especially when you travel, whenever you go to the emergency room, or when you must see another doctor.
- Be sure to notify any health care professional you see that
- you are being treated with KEYTRUDA and show them this card.

- **If you get any side effects, talk to your doctor, pharmacist or nurse. This includes any possible side effects not listed in the package leaflet. You can also report side effects directly to www.lakemedelsverket.se.**
- **By reporting side effects, you can help provide more information on the safety of this medicine.**

For more information, consult the Package Leaflet for KEYTRUDA containing information for the patient. Available at www.fass.se and www.ema.europa.eu.

Important Contact Information

My Name

My Phone

Emergency Contact (Name and Phone)

Name of Doctor

Office Phone

After-hours Phone

Important Information for Health Care Providers

This patient is being treated with KEYTRUDA (*pembrolizumab*), which can cause immune-mediated adverse reactions that may appear any time during treatment or even after treatment. Assess patients for signs and symptoms of immune-mediated adverse reactions. Early diagnosis and appropriate management are essential to minimise any consequences of immune-mediated adverse reactions.

For suspected immune-mediated adverse reactions, ensure adequate evaluation to confirm aetiology or exclude other causes. Based on the severity of the adverse reaction, withhold KEYTRUDA and administer corticosteroids. Specific guidelines for managing immune-mediated adverse reactions are available in the Summary of Product Characteristics for KEYTRUDA.

Consultation with an oncologist or other medical specialist may be helpful for management of organ-specific immune-mediated adverse reactions.

Please consult the Summary of Product Characteristics for KEY TRUDA at www.fass.se and www.ema.europa.eu or call Medical Information at +46 77 570 04 88 for more information.

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