

YERVOY®
(ipilimumab)

Patient Alert Card

Date of MMA approval: AUG 2020
Local approval number: 731MT2005891-01



Important Information for Patients

Carry this card with you at all times to inform healthcare professionals that you are receiving treatment with YERVOY® alone.



If you have any signs or symptoms, tell your doctor right away.

POSSIBLE SIDE EFFECTS



BOWEL AND STOMACH

- diarrhoea (watery, loose or soft stools), bloody or dark-coloured stools
- more frequent bowel movements than usual
- pain or tenderness in your stomach or abdomen area, nausea, vomiting



EYE

- redness in the eye
- pain in the eye
- vision problems or blurry vision



LIVER

- eye or skin yellowing (jaundice)
- pain on the right side of your stomach area
- dark urine



NERVES

- muscle weakness
- numbness or tingling in legs, arms or face
- dizziness, loss of consciousness or difficulty waking up



SKIN

- skin rash with or without itching, dry skin
- blisters and/or peeling of the skin, mouth sores
- swelling of the face or lymph glands



GENERAL

- fever, headache, tiredness
- bleeding
- behavioural changes (e.g. less sex drive, being irritable or forgetful)
- dehydration, low blood pressure, shock



IMPORTANT

- Tell your doctor of any previous medical conditions.
- Early treatment of side effects reduces the likelihood that YERVOY® treatment will need to be temporarily or permanently stopped.
- Signs and symptoms that may appear mild can quickly worsen if left untreated.
- DO NOT try to treat these symptoms yourself.
- Signs and symptoms may be delayed and may occur weeks to months after your last YERVOY® injection.

For more information, read the YERVOY® Package Leaflet at www.ema.europa.eu

My Doctor's Contact Information (*who prescribed YERVOY®*)

Name of Doctor:

Office Phone:

After-Hours Phone:

My Contact Information

My Name and Phone:

Caregiver Name and Phone (in case of emergency):

IMPORTANT Information for Healthcare Professionals

- This patient is treated with **YERVOY**[®] monotherapy.
- Immune-related adverse reactions (irARs) may appear at any time during treatment or months after its discontinuation.
- Early diagnosis and appropriate management are essential to minimise life-threatening complications.
- Consultation with an oncologist or other medical specialist may be helpful for management of organ-specific irARs.
- Healthcare professionals should refer to the YERVOY[®] Summary of Product Characteristics (SmPC) at www.ema.europa.eu or call Medical Information on **00356 2397 6505** for more information.



The healthcare professional treating this patient with YERVOY[®] should complete the 'My Doctor's Contact Information' section of this Patient Alert Card.