My Doctor's Contact Information (who prescribed OPDIVO® or OPDIVO® in combination with YERVOY®) Name of Doctor: Office Phone: Out-of-Hours Phone:

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Date of Health Authority review:

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My Contact **IMPORTANT** Information Information for Healthcare Mv Name: OPDIVO® or OPDIVO® in

My Phone Number:

Emergency Contact

(name and phone number):

- **Professionals** This patient is treated with
- combination with YERVOY®. 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. complete the 'My Doctor's Contact Consultation with an oncologist Information' section of this Patient or other medical specialist may

Alert Card.

be helpful for management of organ-specific irARs. Healthcare professionals

should refer to the OPDIVO® Summary of Product Characteristics (SmPC) at www.fass.se or www.ema.europa.eu or call Bristol-Myers Squibb Medical Information on 08-585 07 304. The Patient Alert Card is also available for download or order on www.immunonkologi.se/ materialbeställning.

The healthcare professional treating this patient with OPDIVO® or OPDIVO® in combination with YERVOY® should

OPDIVO® (nivolumab)

Patient Alert Card

Bristol Myers Squibb

Important Information for Patients

Please carry this card with you at all times to inform healthcare professionals that you are receiving treatment with OPDIVO® or OPDIVO® in combination with YERVOY® (ipilimumab).



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



IMPORTANT

OPDIVO® treatment may increase the risk of serious or even lifethreatening immune-related sideeffects, which may affect different parts of the body, for example:

SUEST (beaut and burne

CHEST (heart and lungs): breathing difficulties, cough, wheeze, chest pain, irregular heartbeat, palpitations (increased awareness of your heartbeat)

GUT (stomach and bowels):diarrhoea (watery, loose or soft stools), blood or mucus in stools, dark-coloured stools, pain or

tenderness in your stomach or abdominal area

LIVER: eye or skin yellowing (jaundice), pain on the right side of your stomach area

KIDNEYS: change in amount and/or frequency of urine

HORMONE-PRODUCING

GLANDS (including Diabetes):
headaches, blurry or double vision,
fatigue (tiredness), weight
changes, behavioural changes (e.g.
lower sex drive, irritability or
forgetfulness), excessive thirst,
increased appetite with weight
loss, weakness, drowsiness,
depression, irritability, feeling
unwell, change in amount and/or
frequency of urine

SKIN: rash, itching, blisters and/or peeling of the skin (possibly fatal), ulcers, dry skin, skin nodules

OTHER: weakness, fatigue (tiredness), decreased appetite, nausea, vomiting, tingling or numbness in arms and legs, difficulty walking, fever, swollen lymph nodes, headache, seizures, stiff neck, confusion, drowsiness, muscle pain, stiffness, dark urine, eye pain or redness, blurry vision, or other vision problems



IMPORTANT

- Tell your doctor of previous medical conditions, including if you have had a stem cell transplant that used donor stem cells (allogeneic).
- Early assessment and management of side-effects by your doctor reduces the likelihood that treatment with OPDIVO® or OPDIVO® in combination with YERVOY® 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 injection.

For more information, read the OPDIVO® Package Leaflet included with the package, as well as on www.fass.se and www.ema.europa.eu or call Medical Information on 08-585 07 304