

# Tecentriq®



## Atezolizumab

Concentrate for solution for infusion

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### NAME OF THE MEDICINAL PRODUCT

Tecentriq

### QUALITATIVE AND QUANTITATIVE COMPOSITION

One vial of 20 mL of concentrate contains 1,200 mg atezolizumab.

After dilution, the final concentration of the diluted solution should be between 3.2 and 16.8 mg/mL.

One 14 mL vial of concentrate contains 840 mg of atezolizumab. After dilution, the final concentration of the diluted solution should be between 3.2 and 16.8 mg/mL.

For dilution and other handling recommendations, see section 2.3.

For the full list of excipients, see section 11.

### PHARMACEUTICAL FORM

Concentrate for solution for infusion (sterile concentrate).

Clear, colourless to slightly yellowish liquid.

### CLINICAL PARTICULARS

#### Patient safety information card and brochure

The marketing of TECENTRIQ is subject to a risk management plan (RMP) including patient safety information materials (patient information card and patient brochure). These materials emphasize important safety information that the patient should be aware of before and during treatment. Please explain to the patient the need to review these materials before starting treatment.

## 1 INDICATIONS AND USAGE

### 1.1 Urothelial Carcinoma

- TECENTRIQ (atezolizumab) is indicated for the treatment of patients with locally advanced or metastatic urothelial carcinoma who are not eligible for cisplatin-containing chemotherapy and whose tumours have a PD-L1 expression  $\geq 5\%$ .
- TECENTRIQ is indicated for the treatment of patients with locally advanced or metastatic urothelial carcinoma who have disease progression during or following any platinum-containing chemotherapy, or within 12 months of neoadjuvant or adjuvant chemotherapy

### 1.2 Non-Small Cell Lung Cancer

- TECENTRIQ, as a single agent, is indicated for the first-line treatment of adult patients with metastatic non-small cell lung cancer (NSCLC) whose tumors have high PD-L1 expression

(PD-L1 stained  $\geq 50\%$  of tumor cells [TC  $\geq 50\%$ ] or PD-L1 stained tumor-infiltrating immune cells [IC] covering  $\geq 10\%$  of the tumor area [IC  $\geq 10\%$ ]), as determined by an approved test, with no EGFR or ALK genomic tumor aberrations.

- TECENTRIQ, in combination with bevacizumab, paclitaxel, and carboplatin, is indicated for the first-line treatment of adult patients with metastatic non-squamous non-small cell lung cancer (NSCLC). In patients with EGFR mutant or ALK-positive NSCLC, TECENTRIQ, in combination with bevacizumab, paclitaxel, and carboplatin, is indicated only after failure of appropriate targeted therapies.
- TECENTRIQ, in combination with paclitaxel protein-bound and carboplatin, is indicated for the first-line treatment of adult patients with metastatic non-squamous NSCLC with no EGFR or ALK genomic tumor aberrations.
- TECENTRIQ is indicated for the treatment of patients with metastatic NSCLC who are naïve to anti-PD-L1 or anti-PD-1 therapies and have disease progression during or following platinum-containing chemotherapy. Patients with EGFR or ALK genomic tumor aberrations should have disease progression on approved therapy for NSCLC harboring these aberrations prior to receiving TECENTRIQ.

### **1.3 Locally Advanced or Metastatic Triple-Negative Breast Cancer**

TECENTRIQ, in combination with nab-paclitaxel, is indicated for the treatment of patients with unresectable locally advanced or metastatic triple-negative breast cancer (TNBC) whose tumors have PD-L1 expression  $\geq 1\%$  and who have not received prior chemotherapy for metastatic disease.

### **1.4 Small Cell Lung Cancer**

TECENTRIQ, in combination with carboplatin and etoposide, is indicated for the first-line treatment of adult patients with extensive-stage small cell lung cancer (ES-SCLC).

### **1.5 Hepatocellular Carcinoma**

TECENTRIQ, in combination with bevacizumab, is indicated for the treatment of patients with unresectable or metastatic hepatocellular carcinoma (HCC) who have not received prior systemic therapy.

### **1.6 Melanoma**

TECENTRIQ, in combination with cobimetinib and vemurafenib, is indicated for the treatment of patients with BRAF V600 mutation-positive unresectable or metastatic melanoma [*see Dosage and Administration (2.1)*].

## **2 DOSAGE AND ADMINISTRATION**

### **2.1 Patient Selection for Treatment of Urothelial Carcinoma, Triple-Negative Breast Cancer, Non-Small Cell Lung Cancer and Melanoma**

Select cisplatin-ineligible patients with previously untreated locally advanced or metastatic urothelial carcinoma for treatment with TECENTRIQ based on the PD-L1 expression on tumor-infiltrating immune cells [*see Clinical Studies (14.1)*].

Select patients with first-line metastatic non-small cell lung cancer for treatment with TECENTRIQ as a single agent based on the PD-L1 expression on tumor cells or on tumor infiltrating immune cells [*see Clinical Studies (14.2)*].

Select patients with locally advanced or metastatic triple-negative breast cancer for treatment with TECENTRIQ in combination with paclitaxel protein-bound based on the PD-L1 expression on tumor infiltrating immune cells [*see Clinical Studies (14.3)*].

Select patients with unresectable or metastatic melanoma for treatment with TECENTRIQ in combination with cobimetinib and vemurafenib after confirming the presence of a BRAF V600 mutation [*see Clinical Studies (14.5)*].

## **2.2 Recommended Dosage for Urothelial Carcinoma**

The recommended dosage of TECENTRIQ is:

- 840 mg every 2 weeks or
- 1200 mg every 3 weeks or
- 1680 mg every 4 weeks

administered intravenously over 60 minutes until disease progression or unacceptable toxicity. If the first infusion is tolerated, all subsequent infusions may be delivered over 30 minutes.

## **2.3 Recommended Dosage for NSCLC**

### Single Agent

The recommended dosage of TECENTRIQ is:

- 840 mg every 2 weeks or
- 1200 mg every 3 weeks or
- 1680 mg every 4 weeks

administered intravenously over 60 minutes until disease progression or unacceptable toxicity. If the first infusion is tolerated, all subsequent infusions may be delivered over 30 minutes.

### TECENTRIQ with Platinum-based Chemotherapy

The recommended dosage of TECENTRIQ is 1200 mg intravenously every 3 weeks until disease progression or unacceptable toxicity.

Administer TECENTRIQ prior to chemotherapy and bevacizumab when given on the same day. Refer to the Prescribing Information for the chemotherapy agents or bevacizumab administered in combination with TECENTRIQ for recommended dosing information.

Following completion of 4-6 cycles of chemotherapy, and if bevacizumab is discontinued, the recommended dosage of TECENTRIQ is:

- 840 mg every 2 weeks or
- 1200 mg every 3 weeks or
- 1680 mg every 4 weeks

administered intravenously until disease progression or unacceptable toxicity.

Administer the initial infusion of TECENTRIQ over 60 minutes. If the first infusion is tolerated, all subsequent infusions may be delivered over 30 minutes.

## **2.4 Recommended Dosage for Locally Advanced or Metastatic TNBC**

The recommended dosage of TECENTRIQ is 840 mg administered intravenously over 60 minutes, followed by 100 mg/m<sup>2</sup> paclitaxel protein-bound.

For each 28 day cycle, TECENTRIQ is administered on days 1 and 15, and paclitaxel protein-bound is administered on days 1, 8, and 15 until disease progression or unacceptable toxicity.

TECENTRIQ and paclitaxel protein-bound may be discontinued for toxicity independently of each other.

If the first infusion is tolerated, all subsequent infusions may be delivered over 30 minutes. Refer to the Prescribing Information for paclitaxel protein-bound for recommended dosing information.

## **2.5 Recommended Dosage for SCLC**

The recommended dosage of TECENTRIQ is 1200 mg intravenously every 3 weeks, when administered in combination with carboplatin and etoposide, until disease progression or unacceptable toxicity.

Administer TECENTRIQ prior to chemotherapy when given on the same day. Refer to the Prescribing Information for the chemotherapy agents administered in combination with TECENTRIQ for recommended dosing information.

Following completion of 4 cycles of carboplatin and etoposide, the recommended dosage of TECENTRIQ is 1200 mg every 3 weeks administered intravenously until disease progression or unacceptable toxicity.

Administer the initial infusion of TECENTRIQ over 60 minutes. If the first infusion is tolerated, all subsequent infusions may be delivered over 30 minutes.

## **2.6 Recommended Dosage for HCC**

The recommended dosage of TECENTRIQ is 1,200 mg administered as an intravenous infusion over 60 minutes, followed by 15 mg/kg of bevacizumab on the same day, every 3 weeks until disease progression or unacceptable toxicity.

Refer to the Prescribing Information for bevacizumab prior to initiation.

If bevacizumab is discontinued for toxicity, the recommended dosage of TECENTRIQ is 1,200 mg every 3 weeks administered intravenously until disease progression or unacceptable toxicity.

If the first infusion of TECENTRIQ is tolerated, all subsequent infusions may be delivered over 30 minutes.

Tecentriq in combination with bevacizumab has not been studied in HCC patients with severe hepatic impairment (Child Pugh C) and is therefore not recommended for use in these patients.

## **2.7 Recommended Dosage for Melanoma**

Prior to initiating TECENTRIQ, patients should receive a 28 day treatment cycle of cobimetinib 60 mg orally once daily (21 days on and 7 days off) and vemurafenib 960 mg orally twice daily from Days 1-21 and vemurafenib 720 mg orally twice daily from Days 22-28.

The recommended dose of TECENTRIQ is 840 mg administered as an intravenous infusion over 60 minutes every 2 weeks until disease progression or unacceptable toxicity, when administered with cobimetinib 60 mg orally once daily (21 days on and 7 days off) and vemurafenib 720 mg orally twice daily.

If the first infusion of TECENTRIQ is tolerated, all subsequent infusions may be delivered over 30 minutes.

Refer to the Prescribing Information for cobimetinib and vemurafenib prior to initiation.

## **2.8 Dosage Modifications for Adverse Reactions**

No dose reduction of TECENTRIQ is recommended.

In general, withhold TECENTRIQ for severe (Grade 3) immune-mediated adverse reactions. Permanently discontinue TECENTRIQ for life-threatening (Grade 4) immune-mediated adverse reactions, recurrent severe (Grade 3) immune-mediated reactions that require systemic

immunosuppressive treatment, or an inability to reduce corticosteroid dose to 10 mg or less of prednisone or equivalent per day within 12 weeks of initiating steroids.

Dosage modifications for TECENTRIQ for adverse reactions that require management different from these general guidelines are summarized in Table 1.

**Table 1: Recommended Dosage Modifications for Adverse Reactions**

| <b>Adverse Reaction</b>   | <b>Severity<sup>a</sup></b>  | <b>Dosage Modification</b>   |
|---|--|--|
| <b>Immune-Mediated Adverse Reactions [see Warnings and Precautions (5.1)]</b> |  |  |
| Pneumonitis   | Grade 2  | Withhold <sup>b</sup>  |
|   | Grades 3 or 4  | Permanently discontinue  |
| Colitis   | Grades 2 or 3  | Withhold <sup>b</sup>  |
|   | Grade 4  | Permanently discontinue  |
| Hepatitis with no tumor involvement of the liver                              | AST or ALT increases to more than 3 and up to 8 times ULN or Total bilirubin increases to more than 1.5 and up to 3 times ULN  | Withhold <sup>b</sup>  |
|   | AST or ALT increases to more than 8 times ULN or Total bilirubin increases to more than 3 times ULN  | Permanently discontinue  |
| Hepatitis with tumor involvement of the liver <sup>c</sup>                    | Baseline AST or ALT is more than 1 and up to 3 times ULN and increases to more than 5 and up to 10 times ULN<br>or<br>Baseline AST or ALT is more than 3 and up to 5 times ULN and increases to more than 8 and up to 10 times ULN | Withhold <sup>b</sup>  |
|   | AST or ALT increases to more than 10 times ULN or<br>Total bilirubin increases to more than 3 times ULN  | Permanently discontinue  |
| Endocrinopathies  | Grades 3, or 4   | Withhold until clinically stable or permanently discontinue depending on severity. |
| Nephritis with Renal Dysfunction  | Grades 2 or 3 increased blood creatinine   | Withhold <sup>b</sup>  |
|   | Grade 4 increased blood creatinine   | Permanently discontinue  |
|   | Suspected SJS, TEN, or DRESS   | Withhold   |

| <b>Adverse Reaction</b>  | <b>Severity<sup>a</sup></b>  | <b>Dosage Modification</b>             |
|--|------------------------------|--|
| Exfoliative Dermatologic Conditions                                | Confirmed SJS, TEN, or DRESS | Permanently discontinue                |
| Myocarditis  | Grades 2, 3, or 4            | Permanently discontinue                |
| Neurological Toxicities  | Grade 2                      | Withhold <sup>b</sup>                  |
|  | Grades 3 or 4                | Permanently discontinue                |
| <b>Other Adverse Reactions</b>                                     |                              |  |
| Infusion-Related Reactions<br>[see Warnings and Precautions (5.2)] | Grades 1 or 2                | Interrupt or slow the rate of infusion |
|  | Grades 3 or 4                | Permanently discontinue                |

<sup>a</sup> Based on Common Terminology Criteria for Adverse Events (CTCAE) version 4

<sup>b</sup> Resume in patients with complete or partial resolution (Grade 0 to 1) after corticosteroid taper. Permanently discontinue if no complete or partial resolution within 12 weeks of initiating steroids or inability to reduce prednisone to 10 mg per day or less (or equivalent) within 12 weeks of initiating steroids.

<sup>c</sup> If AST and ALT are less than or equal to ULN at baseline, withhold or permanently discontinue TECENTRIQ based on recommendations for hepatitis with no liver involvement.

ALT = alanine aminotransferase, AST = aspartate aminotransferase, ULN = upper limit normal, DRESS = Drug Rash with Eosinophilia and Systemic Symptoms, SJS = Stevens Johnson syndrome, TEN = toxic epidermal necrolysis

## 2.9 Preparation and Administration

### Preparation

Visually inspect drug product for particulate matter and discoloration prior to administration, whenever solution and container permit. Discard the vial if the solution is cloudy, discolored, or visible particles are observed. Do not shake the vial.

Prepare the solution for infusion as follows:

- Select the appropriate vial(s) based on the prescribed dose.
- Withdraw the required volume of TECENTRIQ from the vial(s) using sterile needle and syringe.
- Dilute to a final concentration between 3.2 mg/mL and 16.8 mg/mL in a polyvinyl chloride (PVC), polyethylene (PE), or polyolefin (PO) infusion bag containing 0.9% Sodium Chloride Injection, USP.
- Dilute with only 0.9% Sodium Chloride Injection
- Mix diluted solution by gentle inversion. Do not shake.
- Discard used or empty vials of TECENTRIQ.

### Storage of Infusion Solution

This product does not contain a preservative.

Administer immediately once prepared. If diluted TECENTRIQ infusion solution is not used immediately, store solution either:

- At  $\leq 30^{\circ}\text{C}$  for up to 24 hours from the time of preparation. This includes room temperature storage of the infusion in the infusion bag and time for administration of the infusion, or

- Under refrigeration at 2°C-8°C for up to 30 days from time of preparation.
- From a microbiological point of view, the prepared solution for infusion should be used immediately. If not used immediately, in-use storage times and conditions prior to use are the responsibility of the user and would normally not be longer than 24 hours at 2 °C to 8 °C or 8 hours at ambient temperature ( $\leq 25$  °C) unless dilution has taken place in controlled and validated aseptic conditions.

Do not freeze.

Do not shake.

### Administration

Administer the initial infusion over 60 minutes through an intravenous line with or without a sterile, non-pyrogenic, low-protein binding in-line filter (pore size of 0.2–0.22 micron). If the first infusion is tolerated, all subsequent infusions may be delivered over 30 minutes.

Do not coadminister other drugs through the same intravenous line.

Do not administer as an intravenous push or bolus.

## **3. DOSAGE FORMS AND STRENGTHS**

Injection: 840 mg/14 mL (60 mg/mL) and 1200 mg/20 mL (60 mg/mL) colorless to slightly yellow solution in a single-dose vial.

## **4. CONTRAINDICATIONS**

Hypersensitivity to the active substance or to any of the excipients.

## **5. WARNINGS AND PRECAUTIONS**

### **5.1 Severe and Fatal Immune-Mediated Adverse Reactions**

TECENTRIQ is a monoclonal antibody that belongs to a class of drugs that bind to either the programmed death-receptor 1 (PD-1) or the PD-ligand 1 (PD-L1), blocking the PD-1/PD-L1 pathway, thereby removing inhibition of the immune response, potentially breaking peripheral tolerance and inducing immune-mediated adverse reactions. Important immune-mediated adverse reactions listed under Warnings and Precautions may not include all possible severe and fatal immune-mediated reactions.

Immune-mediated adverse reactions, which may be severe or fatal, can occur in any organ system or tissue. Immune-mediated adverse reactions can occur at any time after starting a PD-1/PD-L1 blocking antibody. While immune-mediated adverse reactions usually manifest during treatment with PD-1/PD-L1 blocking antibodies, immune-mediated adverse reactions can also manifest after discontinuation of PD-1/PD-L1 blocking antibodies.

Early identification and management of immune-mediated adverse reactions are essential to ensure safe use of PD-1/PD-L1 blocking antibodies. Monitor patients closely for symptoms and signs that may be clinical manifestations of underlying immune-mediated adverse reactions. Evaluate liver enzymes, creatinine, and thyroid function at baseline and periodically during treatment. In cases of suspected immune-mediated adverse reactions, initiate appropriate workup

to exclude alternative etiologies, including infection. Institute medical management promptly, including specialty consultation as appropriate.

Withhold or permanently discontinue TECENTRIQ depending on severity [*see Dosage and Administration (2.8)*]. In general, if TECENTRIQ requires interruption or discontinuation, administer systemic corticosteroid therapy (1 to 2 mg/kg/day prednisone or equivalent) until improvement to Grade 1 or less. Upon improvement to Grade 1 or less, initiate corticosteroid taper and continue to taper over at least 1 month. Consider administration of other systemic immunosuppressants in patients whose immune-mediated adverse reactions are not controlled with corticosteroid therapy.

Toxicity management guidelines for adverse reactions that do not necessarily require systemic steroids (e.g., endocrinopathies and dermatologic reactions) are discussed below.

### **Immune-Mediated Pneumonitis**

TECENTRIQ can cause immune-mediated pneumonitis. The incidence of pneumonitis is higher in patients who have received prior thoracic radiation.

#### *TECENTRIQ as a Single Agent:*

Immune-mediated pneumonitis occurred in 3% (83/2616) of patients receiving TECENTRIQ as a single agent, including fatal (<0.1%), Grade 4 (0.2%), Grade 3 (0.8%), and Grade 2 (1.1%) adverse reactions. Pneumonitis led to permanent discontinuation of TECENTRIQ in 0.5% and withholding of TECENTRIQ in 1.5% of patients.

Systemic corticosteroids were required in 55% (46/83) of patients with pneumonitis. Pneumonitis resolved in 69% of the 83 patients. Of the 39 patients in whom TECENTRIQ was withheld for pneumonitis, 25 reinitiated TECENTRIQ after symptom improvement; of these, 4% had recurrence of pneumonitis.

#### *TECENTRIQ in Combination with Cobimetinib and Vemurafenib:*

Immune-mediated pneumonitis occurred in 13% (29/230) of patients receiving TECENTRIQ in combination with cobimetinib and vemurafenib, including Grade 3 (1.3%) and Grade 2 (7%) adverse reactions. Pneumonitis led to permanent discontinuation of TECENTRIQ in 2.6% and withholding of TECENTRIQ in 7.4% of patients.

Systemic corticosteroids were required in 55% (16/29) of patients with pneumonitis. Pneumonitis resolved in 97% of the 29 patients. Of the 17 patients in whom TECENTRIQ was withheld for pneumonitis, 10 reinitiated TECENTRIQ after symptom improvement; of these, 50% had recurrence of pneumonitis.

### **Immune-Mediated Colitis**

TECENTRIQ can cause immune-mediated colitis. Colitis can present with diarrhea, abdominal pain, and lower gastrointestinal (GI) bleeding. Cytomegalovirus (CMV) infection/reactivation has been reported in patients with corticosteroid-refractory immune-mediated colitis. In cases of corticosteroid-refractory colitis, consider repeating infectious workup to exclude alternative etiologies.

#### *TECENTRIQ as a Single Agent:*

Immune-mediated colitis occurred in 1% (26/2616) of patients receiving TECENTRIQ as a single agent, including Grade 3 (0.5%) and Grade 2 (0.3%) adverse reactions. Colitis led to

permanent discontinuation of TECENTRIQ in 0.2% and withholding of TECENTRIQ in 0.5% of patients.

Systemic corticosteroids were required in 50% (13/26) of patients with colitis. Colitis resolved in 73% of the 26 patients. Of the 12 patients in whom TECENTRIQ was withheld for colitis, 8 reinitiated treatment with TECENTRIQ after symptom improvement; of these, 25% had recurrence of colitis.

### **Immune-Mediated Hepatitis**

TECENTRIQ can cause immune-mediated hepatitis.

Immune-mediated hepatitis occurred in 1.8% (48/2616) of patients receiving TECENTRIQ as a single agent, including fatal (<0.1%), Grade 4 (0.2%), Grade 3 (0.5%), and Grade 2 (0.5%) adverse reactions. Hepatitis led to permanent discontinuation of TECENTRIQ in 0.2% and withholding of TECENTRIQ in 0.2% of patients.

Systemic corticosteroids were required in 25% (12/48) of patients with hepatitis. Hepatitis resolved in 50% of the 48 patients. Of the 6 patients in whom TECENTRIQ was withheld for hepatitis, 4 reinitiated treatment with TECENTRIQ after symptom improvement; of these, none had recurrence of hepatitis.

#### *TECENTRIQ in Combination with Cobimetinib and Vemurafenib:*

Immune-mediated hepatitis occurred in 6.1% (14/230) of patients receiving TECENTRIQ in combination with cobimetinib and vemurafenib, including Grade 4 (1.3%), Grade 3 (1.7%) and Grade 2 (1.3%) adverse reactions. Hepatitis led to permanent discontinuation of TECENTRIQ in 2.2% and withholding of TECENTRIQ in 1.7% of patients.

Systemic corticosteroids were required in 50% (7/14) of patients with hepatitis. Hepatitis resolved in 93% of the 14 patients. Of the 4 patients in whom TECENTRIQ was withheld for hepatitis, 3 reinitiated TECENTRIQ after symptom improvement; of these, 33% had recurrence of hepatitis.

### **Immune-Mediated Endocrinopathies**

#### *Adrenal Insufficiency*

TECENTRIQ can cause primary or secondary adrenal insufficiency. For Grade 2 or higher adrenal insufficiency, initiate symptomatic treatment, including hormone replacement as clinically indicated. Withhold or permanently discontinue TECENTRIQ depending on severity [see *Dosage and Administration* (2.8)].

Adrenal insufficiency occurred in 0.4% (11/2616) of patients receiving TECENTRIQ as a single agent, including Grade 3 (<0.1%) and Grade 2 (0.2%) adverse reactions. Adrenal insufficiency led to permanent discontinuation of TECENTRIQ in one patient and withholding of TECENTRIQ in one patient.

Systemic corticosteroids were required in 82% (9/11) of patients with adrenal insufficiency, of these, 3 patients remained on systemic corticosteroids. The single patient in whom TECENTRIQ was withheld for adrenal insufficiency did not reinitiate TECENTRIQ.

#### *Hypophysitis*

TECENTRIQ can cause immune-mediated hypophysitis. Hypophysitis can present with acute symptoms associated with mass effect such as headache, photophobia, or visual field cuts. Hypophysitis can cause hypopituitarism. Initiate hormone replacement as clinically indicated. Withhold or permanently discontinue TECENTRIQ depending on severity [see *Dosage and Administration* (2.8)].

Hypophysitis occurred in <0.1% (2/2616) of patients receiving TECENTRIQ as a single agent, including Grade 2 (1 patient, <0.1%) adverse reactions. Hypophysitis led to permanent discontinuation of TECENTRIQ in one patient and no patients required withholding of TECENTRIQ.

Systemic corticosteroids were required in 50% (1/2) of patients with hypophysitis. Hypophysitis did not resolve in these 2 patients.

*Thyroid disorders:*

TECENTRIQ can cause immune-mediated thyroid disorders. Thyroiditis can present with or without endocrinopathy. Hypothyroidism can follow hyperthyroidism. Initiate hormone replacement for hypothyroidism or medical management for hyperthyroidism as clinically indicated. Withhold or permanently discontinue TECENTRIQ depending on severity [see *Dosage and Administration* (2.8)].

*Thyroiditis:*

Thyroiditis occurred in 0.2% (4/2616) of patients receiving TECENTRIQ as a single agent, including Grade 2 (<0.1%) adverse reactions. Thyroiditis did not lead to permanent discontinuation of TECENTRIQ in any of these patients, but led to withholding of TECENTRIQ in one patient.

Hormone replacement therapy was required in 75% (3/4) of patients with thyroiditis. Systemic corticosteroids were required in 25% (1/4) of patients with thyroiditis. Thyroiditis resolved in 50% of patients. The single patient in whom TECENTRIQ was withheld for thyroiditis reinitiated TECENTRIQ; this patient did not have recurrence of thyroiditis.

*Hyperthyroidism:*

*TECENTRIQ as a Single Agent:*

Hyperthyroidism occurred in 0.8% (21/2616) of patients receiving TECENTRIQ as a single agent, including Grade 2 (0.4%) adverse reactions. Hyperthyroidism did not lead to permanent discontinuation of TECENTRIQ in any of these patients, but led to withholding of TECENTRIQ in 0.1% of patients.

Antithyroid therapy was required in 29% (6/21) of patients with hyperthyroidism. Of these 6 patients, the majority remained on antithyroid treatment. Of the 3 patients in whom TECENTRIQ was withheld for hyperthyroidism, one patient reinitiated TECENTRIQ; this patient did not have recurrence of hyperthyroidism.

*TECENTRIQ in Combination with Cobimetinib and Vemurafenib:*

Hyperthyroidism occurred in 19% (43/230) of patients receiving TECENTRIQ in combination with cobimetinib and vemurafenib, including Grade 3 (0.9%) and Grade 2 (7.8%) adverse reactions. Hyperthyroidism led to permanent discontinuation of TECENTRIQ in 0.4% and withholding of TECENTRIQ in 10% of patients.

Antithyroid therapy was required in 53% (23/43) of patients with hyperthyroidism. Of these 23 patients, the majority remained on antithyroid treatment. Of the 24 patients in whom TECENTRIQ was withheld for hyperthyroidism, 18 patients reinitiated TECENTRIQ; of these, 28% had recurrence of hyperthyroidism.

*Hypothyroidism:*

*TECENTRIQ as a Single Agent:*

Hypothyroidism occurred in 4.9% (128/2616) of patients receiving TECENTRIQ as a single agent, including Grade 3 (0.2%) and Grade 2 (3.4%) adverse reactions. Hypothyroidism did not

lead to permanent discontinuation of TECENTRIQ in any of these patients, but led to withholding of TECENTRIQ in 0.6% of patients.

Hormone replacement therapy was required in 81% (104/128) of patients with hypothyroidism. The majority of patients with hypothyroidism remained on thyroid hormone replacement. Of the 17 patients in whom TECENTRIQ was withheld for hypothyroidism, 8 reinitiated TECENTRIQ after symptom improvement.

#### *TECENTRIQ in Combination with Platinum-based Chemotherapy:*

Hypothyroidism occurred in 11% (277/2421) of patients with NSCLC and SCLC receiving TECENTRIQ in combination with platinum-based chemotherapy, including Grade 4 (<0.1%), Grade 3 (0.3%), and Grade 2 (5.7%) adverse reactions. Hypothyroidism led to permanent discontinuation of TECENTRIQ in 0.1% and withholding of TECENTRIQ in 1.6% of patients.

Hormone replacement therapy was required in 71% (198/277) of patients with hypothyroidism. The majority of patients with hypothyroidism remained on thyroid hormone replacement. Of the 39 patients in whom TECENTRIQ was withheld for hypothyroidism, 9 reinitiated TECENTRIQ after symptom improvement.

#### *TECENTRIQ in Combination with Cobimetinib and Vemurafenib:*

Hypothyroidism occurred in 26% (60/230) of patients receiving TECENTRIQ in combination with cobimetinib and vemurafenib, including Grade 2 (9.1%) adverse reactions. Hypothyroidism did not lead to permanent discontinuation of TECENTRIQ in any of these patients, but led to withholding of TECENTRIQ in 2.6% of patients.

Hormone replacement therapy was required in 52% (31/60) of patients with hypothyroidism. The majority of patients with hypothyroidism remained on thyroid hormone replacement. Of the 6 patients in whom TECENTRIQ was withheld for hypothyroidism, 4 reinitiated TECENTRIQ after symptom improvement. The majority of patients with hypothyroidism required long term thyroid replacement.

#### *Type 1 Diabetes Mellitus , which can present with Diabetic Ketoacidosis*

Monitor patients for hyperglycemia or other signs and symptoms of diabetes. Initiate treatment with insulin as clinically indicated. Withhold or permanently discontinue TECENTRIQ depending on severity [see *Dosage and Administration (2.8)*].

Type 1 diabetes mellitus occurred in 0.3% (7/2616) of patients receiving TECENTRIQ, including Grade 3 (0.2%) and Grade 2 (<0.1%) adverse reactions. Type 1 diabetes mellitus led to permanent discontinuation of TECENTRIQ in one patient and withholding of TECENTRIQ in two patients.

Treatment with insulin was required for all patients with confirmed Type 1 diabetes mellitus and insulin therapy was continued long-term. Of the 2 patients in whom TECENTRIQ was withheld for Type 1 diabetes mellitus, both re-initiated TECENTRIQ treatment.

### **Immune-Mediated Nephritis with Renal Dysfunction**

TECENTRIQ can cause immune-mediated nephritis.

#### *TECENTRIQ as a Single Agent:*

Immune-mediated nephritis with renal dysfunction occurred in <0.1% (1/2616) of patients receiving TECENTRIQ as a single agent, and this adverse reaction was a Grade 3 (<0.1%) adverse reaction. Nephritis led to permanent discontinuation of TECENTRIQ in this patient.

This patient required systemic corticosteroids. In this patient, nephritis did not resolve.

#### *TECENTRIQ in Combination with Cobimetinib and Vemurafenib:*

Immune-mediated nephritis with renal dysfunction occurred in 1.3% (3/230) of patients receiving TECENTRIQ in combination with cobimetinib and vemurafenib, including Grade 2 (1.3%) adverse reactions. Nephritis led to permanent discontinuation of TECENTRIQ in 0.4% and withholding of TECENTRIQ in 0.9% of patients.

Systemic corticosteroids were required in 67% (2/3) of patients with nephritis. Nephritis resolved in all 3 of these patients. Of the 2 patients in whom TECENTRIQ was withheld for nephritis, both reinitiated TECENTRIQ after symptom improvement and neither had recurrence of nephritis.

### **Immune-Mediated Dermatologic Adverse Reactions**

TECENTRIQ can cause immune-mediated rash or dermatitis. Exfoliative dermatitis, including Stevens-Johnson syndrome (SJS), DRESS, and toxic epidermal necrolysis (TEN), has occurred with PD-1/PD-L1 blocking antibodies. Topical emollients and/or topical corticosteroids may be adequate to treat mild to moderate non-exfoliative rashes. Withhold or permanently discontinue TECENTRIQ depending on severity [*see Dosage and Administration (2.8)*].

Immune-mediated dermatologic adverse reactions occurred in 0.6% (15/2616) of patients receiving TECENTRIQ as a single agent, including Grade 3 (<0.1%) and Grade 2 (0.2%) adverse reactions. Dermatologic adverse reactions led to permanent discontinuation of TECENTRIQ in 0.1% and withholding of TECENTRIQ in 0.2% of patients.

Systemic corticosteroids were required in 20% (3/15) of patients with dermatologic adverse reactions. Dermatologic adverse reactions resolved in 87% of the 15 patients. Of the 4 patients in whom TECENTRIQ was withheld for immune-mediated dermatologic adverse reactions, none re-initiated TECENTRIQ.

### **Other Immune-Mediated Adverse Reactions**

The following clinically significant immune-mediated adverse reactions occurred at an incidence of < 1% (unless otherwise noted) in patients who received TECENTRIQ or were reported with the use of other PD-1/PD-L1 blocking antibodies. Severe or fatal cases have been reported for some of these adverse reactions.

*Cardiac/Vascular:* Myocarditis, pericarditis, vasculitis

*Nervous System:* Meningitis, encephalitis, myelitis and demyelination, myasthenic syndrome/myasthenia gravis (including exacerbation), Guillain-Barré syndrome, nerve paresis, autoimmune neuropathy

*Ocular:* Uveitis, iritis, and other ocular inflammatory toxicities can occur. Some cases can be associated with retinal detachment. Various grades of visual impairment, including blindness, can occur. If uveitis occurs in combination with other immune-mediated adverse reactions, consider a Vogt-Koyanagi-Harada-like syndrome, as this may require treatment with systemic steroids to reduce the risk of permanent vision loss.

*Gastrointestinal:* Pancreatitis to include increases in serum amylase and lipase levels, gastritis, duodenitis.

*Musculoskeletal and Connective Tissue:* Myositis/polymyositis, rhabdomyolysis and associated sequelae including renal failure, arthritis, polymyalgia rheumatic.

*Endocrine:* Hypoparathyroidism

*Other (Hematologic/Immune):* Hemolytic anemia, aplastic anemia, hemophagocytic lymphohistiocytosis, systemic inflammatory response syndrome, histiocytic necrotizing lymphadenitis (Kikuchi lymphadenitis), sarcoidosis, immune thrombocytopenic purpura, solid organ transplant rejection.

## 5.2 Infusion-Related Reactions

TECENTRIQ can cause severe or life-threatening infusion-related reactions. Monitor for signs and symptoms of infusion-related reactions. Interrupt, slow the rate of, or permanently discontinue TECENTRIQ based on the severity [see *Dosage and Administration (2.8)*]. For Grade 1 or 2 infusion-related reactions, consider using pre-medications with subsequent doses.

In clinical studies enrolling 2616 patients with various cancers who received TECENTRIQ as a single-agent [see *Adverse Reactions (6.1)*], infusion-related reactions occurred in 1.3% of patients, including Grade 3 (0.2%). The frequency and severity of infusion-related reactions were similar whether TECENTRIQ was given as a single-agent in patients with various cancers, in combination with other antineoplastic drugs in NSCLC and SCLC, and across the recommended dose range (840 mg Q2W to 1680 mg Q4W).

## 5.3 Complications of Allogeneic HSCT after PD-1/PD-L1 Inhibitors

Fatal and other serious complications can occur in patients who receive allogeneic hematopoietic stem cell transplantation (HSCT) before or after being treated with a PD-1/PD-L1 blocking antibody. Transplant-related complications include hyperacute graft-versus-host disease (GVHD), acute GVHD, chronic GVHD, hepatic veno-occlusive disease (VOD) after reduced intensity conditioning, and steroid-requiring febrile syndrome (without an identified infectious cause). These complications may occur despite intervening therapy between PD-1/PD-L1 blockage and allogeneic HSCT.

Follow patients closely for evidence of transplant-related complications and intervene promptly. Consider the benefits versus risks of treatment with a PD-1/PD-L1 blocking antibody prior to or after an allogeneic HSCT.

## 5.4 Embryo-Fetal Toxicity

Based on its mechanism of action, TECENTRIQ can cause fetal harm when administered to a pregnant woman. There are no available data on the use of TECENTRIQ in pregnant women. Animal studies have demonstrated that inhibition of the PD-L1/PD-1 pathway can lead to increased risk of immune-related rejection of the developing fetus resulting in fetal death.

Verify pregnancy status of females of reproductive potential prior to initiating TECENTRIQ. Advise females of reproductive potential of the potential risk to a fetus. Advise females of reproductive potential to use effective contraception during treatment with TECENTRIQ and for at least 5 months after the last dose [see *Use in Specific Populations (8.1, 8.3)*].

## 6 ADVERSE REACTIONS

The following clinically significant adverse reactions are described elsewhere in the labeling:

- Severe and Fatal Immune-Mediated Adverse Reactions [see *Warnings and Precautions (5.1)*]
- Infusion-Related Reactions [see *Warnings and Precautions (5.2)*]
- Complications of Allogeneic HSCT after PD-1/PD-L1 Inhibitors [see *Warnings and Precautions (5.3)*]

## 6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

The data described in WARNINGS AND PRECAUTIONS reflect exposure to TECENTRIQ as a single-agent in 2616 patients in two randomized, active-controlled studies (POPLAR, OAK) and four open-label, single arm studies (PCD4989g, IMvigor210, BIRCH, FIR) which enrolled 524 patients with metastatic urothelial carcinoma, 1636 patients with metastatic NSCLC, and 456 patients with other tumor types. TECENTRIQ was administered at a dose of 1200 mg intravenously every 3 weeks in all studies except PCD4989g. Among the 2616 patients who received a single-agent TECENTRIQ, 36% were exposed for longer than 6 months and 20% were exposed for longer than 12 months. Using the dataset described for patients who received TECENTRIQ as a single-agent, the most common adverse reactions in  $\geq 20\%$  of patients were fatigue/asthenia (48%), decreased appetite (25%), nausea (24%), cough (22%), and dyspnea (22%).

In addition, the data reflect exposure to TECENTRIQ in combination with other antineoplastic drugs in 2421 patients with NSCLC (N = 2223) or SCLC (N = 198) enrolled in five randomized, active-controlled trials, including IMpower150, IMpower130 and IMpower133. Among the 2421 patients, 53% were exposed to TECENTRIQ for longer than 6 months and 29% were exposed to TECENTRIQ for longer than 12 months. Among the 2421 patients with NSCLC and SCLC who received TECENTRIQ in combination with other antineoplastic drugs, the most common adverse reactions in  $\geq 20\%$  of patients were fatigue/asthenia (49%), nausea (38%), alopecia (35%), constipation (29%), diarrhea (28%) and decreased appetite (27%).

The data also reflect exposure to TECENTRIQ administered in combination with cobimetinib and vemurafenib in 230 patients enrolled in IMspire150. Among the 230 patients, 62% were exposed to TECENTRIQ for longer than 6 months and 42% were exposed to TECENTRIQ for longer than 12 months.

### Urothelial Carcinoma

#### *Cisplatin-Ineligible Patients with Locally Advanced or Metastatic Urothelial Carcinoma*

The safety of TECENTRIQ was evaluated in IMvigor 210 (Cohort 1), a multicenter, open-label, single-arm trial that included 119 patients with locally advanced or metastatic urothelial carcinoma who were ineligible for cisplatin-containing chemotherapy and were either previously untreated or had disease progression at least 12 months after neoadjuvant or adjuvant chemotherapy [see *Clinical Studies (14.1)*]. Patients received TECENTRIQ 1200 mg intravenously every 3 weeks until either unacceptable toxicity or disease progression. The median duration of exposure was 15 weeks (0 to 87 weeks).

Five patients (4.2%) who were treated with TECENTRIQ experienced one of the following events which led to death: sepsis, cardiac arrest, myocardial infarction, respiratory failure, or respiratory distress. One additional patient (0.8%) was experiencing herpetic meningoencephalitis and disease progression at the time of death.

Serious adverse reactions occurred in 37% of patients. The most frequent serious adverse reactions ( $\geq 2\%$ ) were diarrhea, intestinal obstruction, sepsis, acute kidney injury, and renal failure.

TECENTRIQ was discontinued for adverse reactions in 4.2% of patients. The adverse reactions leading to discontinuation were diarrhea/colitis (1.7%), fatigue (0.8%), hypersensitivity (0.8%), and dyspnea (0.8%).

Adverse reactions leading to interruption occurred in 35% of patients; the most common ( $\geq 1\%$ ) were intestinal obstruction, fatigue, diarrhea, urinary tract infection, infusion-related reaction, cough, abdominal pain, peripheral edema, pyrexia, respiratory tract infection, upper respiratory tract infection, creatinine increase, decreased appetite, hyponatremia, back pain, pruritus, and venous thromboembolism.

Tables 2 and 3 summarize the adverse reactions and Grades 3–4 selected laboratory abnormalities, respectively, in patients who received TECENTRIQ in IMvigor210 (Cohort 1).

**Table 2: Adverse Reactions in  $\geq 10\%$  of Patients with Urothelial Carcinoma in IMvigor210 (Cohort 1)**

| Adverse Reaction                             | TECENTRIQ<br>N = 119 |                |
|--|----------------------|----------------|
|  | All Grades (%)       | Grades 3–4 (%) |
| <b>General</b>                               |                      |                |
| Fatigue <sup>1</sup>                         | 52                   | 8              |
| Peripheral edema <sup>2</sup>                | 17                   | 2              |
| Pyrexia                                      | 14                   | 0.8            |
| <b>Gastrointestinal</b>                      |                      |                |
| Diarrhea <sup>3</sup>                        | 24                   | 5              |
| Nausea                                       | 22                   | 2              |
| Vomiting                                     | 16                   | 0.8            |
| Constipation                                 | 15                   | 2              |
| Abdominal pain <sup>4</sup>                  | 15                   | 0.8            |
| <b>Metabolism and Nutrition</b>              |                      |                |
| Decreased appetite <sup>5</sup>              | 24                   | 3              |
| <b>Musculoskeletal and Connective Tissue</b> |                      |                |
| Back/Neck pain                               | 18                   | 3              |
| Arthralgia                                   | 13                   | 0              |
| <b>Skin and Subcutaneous Tissue</b>          |                      |                |
| Pruritus                                     | 18                   | 0.8            |
| Rash <sup>6</sup>                            | 17                   | 0.8            |
| <b>Infections</b>                            |                      |                |
| Urinary tract infection <sup>7</sup>         | 17                   | 5              |

| <b>Respiratory, Thoracic, and Mediastinal</b> |    |   |
|---|----|---|
| Cough <sup>8</sup>                            | 14 | 0 |
| Dyspnea <sup>9</sup>                          | 12 | 0 |

<sup>1</sup> Includes fatigue, asthenia, lethargy, and malaise

<sup>2</sup> Includes edema peripheral, scrotal edema, lymphedema, and edema

<sup>3</sup> Includes diarrhea, colitis, frequent bowel movements, autoimmune colitis

<sup>4</sup> Includes abdominal pain, upper abdominal pain, lower abdominal pain, and flank pain

<sup>5</sup> Includes decreased appetite and early satiety

<sup>6</sup> Includes rash, dermatitis, dermatitis acneiform, rash maculo-papular, rash erythematous, rash pruritic, rash macular, and rash papular

<sup>7</sup> Includes urinary tract infection, urinary tract infection bacterial, cystitis, and urosepsis

<sup>8</sup> Includes cough and productive cough

<sup>9</sup> Includes dyspnea and exertional dyspnea

**Table 3: Grades 3–4 Laboratory Abnormalities in  $\geq$  1% of Patients with Urothelial Carcinoma in IMvigor210 (Cohort 1)**

| <b>Laboratory Abnormality</b>  | <b>Grades 3–4 (%)</b> |
|--------------------------------|-----------------------|
| <b>Chemistry</b>               |                       |
| Hyponatremia                   | 15                    |
| Hyperglycemia                  | 10                    |
| Increased Alkaline Phosphatase | 7                     |
| Increased Creatinine           | 5                     |
| Hypophosphatemia               | 4                     |
| Increased ALT                  | 4                     |
| Increased AST                  | 4                     |
| Hyperkalemia                   | 3                     |
| Hypermagnesemia                | 3                     |
| Hyperbilirubinemia             | 3                     |
| <b>Hematology</b>              |                       |
| Lymphopenia                    | 9                     |
| Anemia                         | 7                     |

Graded per NCI CTCAE v4.0.

#### *Previously Treated Locally Advanced or Metastatic Urothelial Carcinoma*

The safety of TECENTRIQ was evaluated in IMvigor210 (Cohort 2), a multicenter, open-label, single-arm trial that included 310 patients with locally advanced or metastatic urothelial carcinoma who had disease progression during or following at least one platinum-containing chemotherapy regimen or who had disease progression within 12 months of treatment with a platinum-containing neoadjuvant or adjuvant chemotherapy regimen [see *Clinical Studies (14.1)*]. Patients received TECENTRIQ 1200 mg intravenously every 3 weeks until unacceptable toxicity or either radiographic or clinical progression. The median duration of exposure was 12.3 weeks (0.1 to 46 weeks).

Three patients (1%) who were treated with TECENTRIQ experienced one of the following events which led to death: sepsis, pneumonitis, or intestinal obstruction.

TECENTRIQ was discontinued for adverse reactions in 3.2% of patients. Sepsis led to discontinuation in 0.6% of patients.

Serious adverse reactions occurred in 45% of patients. The most frequent serious adverse reactions (> 2%) were urinary tract infection, hematuria, acute kidney injury, intestinal obstruction, pyrexia, venous thromboembolism, urinary obstruction, pneumonia, dyspnea, abdominal pain, sepsis, and confusional state.

Adverse reactions leading to interruption occurred in 27% of patients; the most common (> 1%) were liver enzyme increase, urinary tract infection, diarrhea, fatigue, confusional state, urinary obstruction, pyrexia, dyspnea, venous thromboembolism, and pneumonitis.

Tables 4 and 5 summarize the adverse reactions and Grades 3–4 selected laboratory abnormalities, respectively, in patients who received TECENTRIQ in IMvigor210 (Cohort 2).

**Table 4: Adverse Reactions in  $\geq 10\%$  of Patients with Urothelial Carcinoma in IMvigor210 (Cohort 2)**

| Adverse Reaction                              | TECENTRIQ<br>N = 310 |                |
|---|----------------------|----------------|
|   | All Grades (%)       | Grades 3–4 (%) |
| <b>General</b>                                |                      |                |
| Fatigue                                       | 52                   | 6              |
| Pyrexia                                       | 21                   | 1              |
| Peripheral edema                              | 18                   | 1              |
| <b>Metabolism and Nutrition</b>               |                      |                |
| Decreased appetite                            | 26                   | 1              |
| <b>Gastrointestinal</b>                       |                      |                |
| Nausea  | 25                   | 2              |
| Constipation                                  | 21                   | 0.3            |
| Diarrhea                                      | 18                   | 1              |
| Abdominal pain                                | 17                   | 4              |
| Vomiting                                      | 17                   | 1              |
| <b>Infections</b>                             |                      |                |
| Urinary tract infection                       | 22                   | 9              |
| <b>Respiratory, Thoracic, and Mediastinal</b> |                      |                |
| Dyspnea                                       | 16                   | 4              |
| Cough   | 14                   | 0.3            |
| <b>Musculoskeletal and Connective Tissue</b>  |                      |                |
| Back/Neck pain                                | 15                   | 2              |
| Arthralgia                                    | 14                   | 1              |
| <b>Skin and Subcutaneous Tissue</b>           |                      |                |
| Rash  | 15                   | 0.3            |

|                          |    |     |
|--------------------------|----|-----|
| Pruritus                 | 13 | 0.3 |
| <b>Renal and Urinary</b> |    |     |
| Hematuria                | 14 | 3   |

**Table 5: Grades 3–4 Laboratory Abnormalities in  $\geq 1\%$  of Patients with Urothelial Carcinoma in IMvigor210 (Cohort 2)**

| Laboratory Abnormality         | Grades 3–4 (%) |
|--------------------------------|----------------|
| <b>Chemistry</b>               |                |
| Hyponatremia                   | 10             |
| Hyperglycemia                  | 5              |
| Increased Alkaline Phosphatase | 4              |
| Increased Creatinine           | 3              |
| Increased ALT                  | 2              |
| Increased AST                  | 2              |
| Hypoalbuminemia                | 1              |
| <b>Hematology</b>              |                |
| Lymphopenia                    | 10             |
| Anemia                         | 8              |

Graded per NCI CTCAE v4.0.

### Non-small Cell Lung Cancer (NSCLC)

#### *IMpower110*

The safety of TECENTRIQ was evaluated in IMpower110, a multicenter, international, randomized, open-label study in 549 chemotherapy-naïve patients with stage IV NSCLC, including those with EGFR or ALK genomic tumor aberrations. Patients received TECENTRIQ 1200 mg every 3 weeks (n=286) or platinum-based chemotherapy consisting of carboplatin or cisplatin with either pemetrexed or gemcitabine (n=263) until disease progression or unacceptable toxicity [see *Clinical Studies (14.2)*]. IMpower110 enrolled patients whose tumors express PD-L1 (PD-L1 stained  $\geq 1\%$  of tumor cells [TC] or PD-L1 stained tumor-infiltrating immune cells [IC] covering  $\geq 1\%$  of the tumor area). The median duration of exposure to TECENTRIQ was 5.3 months (0 to 33 months).

Fatal adverse reactions occurred in 3.8% of patients receiving TECENTRIQ; these included death (reported as unexplained death and death of unknown cause), aspiration, chronic obstructive pulmonary disease, pulmonary embolism, acute myocardial infarction, cardiac arrest, mechanical ileus, sepsis, cerebral infarction, and device occlusion (1 patient each).

Serious adverse reactions occurred in 28% of patients receiving TECENTRIQ. The most frequent serious adverse reactions ( $>2\%$ ) were pneumonia (2.8%), chronic obstructive pulmonary disease (2.1%) and pneumonitis (2.1%)

TECENTRIQ was discontinued due to adverse reactions in 6% of patients; the most common adverse reactions ( $\geq 2$  patients) leading to TECENTRIQ discontinuation were peripheral neuropathy and pneumonitis.

Adverse reactions leading to interruption of TECENTRIQ occurred in 26% of patients; the most common (>1%) were ALT increased (2.1%), AST increased (2.1%), pneumonitis (2.1%), pyrexia (1.4%), pneumonia (1.4%) and upper respiratory tract infection (1.4%).

Tables 6 and 7 summarize adverse reactions and selected laboratory abnormalities in patients receiving TECENTRIQ in IMpower110.

**Table 6: Adverse Reactions Occurring in  $\geq 10\%$  of Patients with NSCLC Receiving TECENTRIQ in IMpower110**

| Adverse Reaction                             | TECENTRIQ<br>N = 286 |             | Platinum-Based Chemotherapy<br>N = 263 |                |
|--|----------------------|-------------|--|----------------|
|  | All Grades (%)       | Grades 3–4* | All Grades*                            | Grades 3–4 (%) |
| <b>Gastrointestinal</b>                      |                      |             |  |                |
| Nausea                                       | 14                   | 0.3         | 34                                     | 1.9            |
| Constipation                                 | 12                   | 1.0         | 22                                     | 0.8            |
| Diarrhea                                     | 11                   | 0           | 12                                     | 0.8            |
| <b>General</b>                               |                      |             |  |                |
| Fatigue/asthenia                             | 25                   | 1.4         | 34                                     | 4.2            |
| Pyrexia                                      | 14                   | 0           | 9                                      | 0.4            |
| <b>Metabolism and Nutrition</b>              |                      |             |  |                |
| Decreased appetite                           | 15                   | 0.7         | 19                                     | 0              |
| <b>Respiratory, Thoracic and Mediastinal</b> |                      |             |  |                |
| Dyspnea                                      | 14                   | 0.7         | 10                                     | 0              |
| Cough  | 12                   | 0.3         | 10                                     | 0              |

Graded per NCI CTCAE v4.0

**Table 7: Laboratory Abnormalities Worsening from Baseline Occurring in  $\geq 20\%$  of Patients Receiving TECENTRIQ in IMpower110**

| Laboratory Abnormality         | TECENTRIQ      |                | Platinum Based Chemotherapy |                |
|--------------------------------|----------------|----------------|-----------------------------|----------------|
|                                | All Grades (%) | Grades 3–4 (%) | All Grades (%)              | Grades 3–4 (%) |
| <b>Hematology</b>              |                |                |                             |                |
| Anemia                         | 69             | 1.8            | 94                          | 20             |
| Lymphopenia                    | 47             | 9              | 59                          | 17             |
| <b>Chemistry</b>               |                |                |                             |                |
| Hypoalbuminemia                | 48             | 0.4            | 39                          | 2              |
| Increased alkaline phosphatase | 46             | 2.5            | 42                          | 1.2            |
| Hyponatremia                   | 44             | 9              | 36                          | 7              |
| Increased ALT                  | 38             | 3.2            | 32                          | 0.8            |
| Increased AST                  | 36             | 3.2            | 32                          | 0.8            |
| Hyperkalemia                   | 29             | 3.9            | 36                          | 2.7            |
| Hypocalcemia                   | 24             | 1.4            | 24                          | 2.7            |
| Increased blood creatinine     | 24             | 0.7            | 33                          | 1.5            |
| Hypophosphatemia               | 23             | 3.6            | 21                          | 2              |

Each test incidence is based on the number of patients who had at least one on-study laboratory measurement available: TECENTRIQ (range: 278-281); platinum-based chemotherapy (range:256-260). Graded per NCI CTCAE v4.0. Increased blood creatinine only includes patients with test results above the normal range.

### *IMpower150*

The safety of TECENTRIQ with bevacizumab, paclitaxel and carboplatin was evaluated in IMpower150, a multicenter, international, randomized, open-label trial in which 393 chemotherapy-naïve patients with metastatic non-squamous NSCLC received TECENTRIQ 1200 mg with bevacizumab 15 mg/kg, paclitaxel 175 mg/m<sup>2</sup> or 200 mg/m<sup>2</sup>, and carboplatin AUC 6 mg/mL/min intravenously every 3 weeks for a maximum of 4 or 6 cycles, followed by TECENTRIQ 1200 mg with bevacizumab 15 mg/kg intravenously every 3 weeks until disease progression or unacceptable toxicity [see *Clinical Studies (14.2)*]. The median duration of exposure to TECENTRIQ was 8.3 months in patients receiving TECENTRIQ with bevacizumab, paclitaxel, and carboplatin.

Fatal adverse reactions occurred in 6% of patients receiving TECENTRIQ; these included hemoptysis, febrile neutropenia, pulmonary embolism, pulmonary hemorrhage, death, cardiac arrest, cerebrovascular accident, pneumonia, aspiration pneumonia, chronic obstructive pulmonary disease, intracranial hemorrhage, intestinal angina, intestinal ischemia, intestinal obstruction and aortic dissection.

Serious adverse reactions occurred in 44%. The most frequent serious adverse reactions (>2%) were febrile neutropenia, pneumonia, diarrhea, and hemoptysis.

TECENTRIQ was discontinued due to adverse reactions in 15% of patients; the most common adverse reaction leading to discontinuation was pneumonitis (1.8%).

Adverse reactions leading to interruption of TECENTRIQ occurred in 48%; the most common (>1%) were neutropenia, thrombocytopenia, fatigue/asthenia, diarrhea, hypothyroidism, anemia, pneumonia, pyrexia, hyperthyroidism, febrile neutropenia, increased ALT, dyspnea, dehydration and proteinuria.

Tables 8 and 9 summarize adverse reactions and laboratory abnormalities in patients receiving TECENTRIQ with bevacizumab, paclitaxel, and carboplatin in IMpower150.

**Table 8: Adverse Reactions Occurring in ≥15% of Patients with NSCLC Receiving TECENTRIQ in IMpower150**

| Adverse Reaction                             | TECENTRIQ with Bevacizumab, Paclitaxel, and Carboplatin<br>N = 393 |                | Bevacizumab, Paclitaxel and Carboplatin<br>N = 394 |                |
|--|--|----------------|--|----------------|
|  | All Grades (%)   | Grades 3–4 (%) | All Grades (%)                                     | Grades 3–4 (%) |
| <b>Nervous System</b>                        |  |                |  |                |
| Neuropathy <sup>1</sup>                      | 56   | 3              | 47   | 3              |
| Headache                                     | 16   | 0.8            | 13   | 0              |
| <b>General</b>                               |  |                |  |                |
| Fatigue/Asthenia                             | 50   | 6              | 46   | 6              |
| Pyrexia                                      | 19   | 0.3            | 9  | 0.5            |
| <b>Skin and Subcutaneous Tissue</b>          |  |                |  |                |
| Alopecia                                     | 48   | 0              | 46   | 0              |
| Rash <sup>2</sup>                            | 23   | 2              | 10   | 0.3            |
| <b>Musculoskeletal and Connective Tissue</b> |  |                |  |                |
| Myalgia/Pain <sup>3</sup>                    | 42   | 3              | 34   | 2              |
| Arthralgia                                   | 26   | 1              | 22   | 1              |
| <b>Gastrointestinal</b>                      |  |                |  |                |
| Nausea                                       | 39   | 4              | 32   | 2              |
| Diarrhea <sup>4</sup>                        | 33   | 6              | 25   | 0.5            |
| Constipation                                 | 30   | 0.3            | 23   | 0.3            |
| Vomiting                                     | 19   | 2              | 18   | 1              |
| <b>Metabolism and Nutrition</b>              |  |                |  |                |
| Decreased appetite                           | 29   | 4              | 21   | 0.8            |
| <b>Vascular</b>                              |  |                |  |                |
| Hypertension                                 | 25   | 9              | 22   | 8              |
| <b>Respiratory</b>                           |  |                |  |                |
| Cough  | 20   | 0.8            | 19   | 0.3            |
| Epistaxis                                    | 17   | 1              | 22   | 0.3            |
| <b>Renal</b>                                 |  |                |  |                |
| Proteinuria <sup>5</sup>                     | 16   | 3              | 15   | 3              |

Graded per NCI CTCAE v4.0

<sup>1</sup> Includes neuropathy peripheral, peripheral sensory neuropathy, hypoesthesia, paraesthesia, dysesthesia, polyneuropathy.

<sup>2</sup> Includes rash, rash maculo-papular, drug eruption, eczema, eczema asteatotic, dermatitis, contact dermatitis, rash erythematous, rash macular, pruritic rash, seborrheic dermatitis, dermatitis psoriasiform.

<sup>3</sup> Includes pain in extremity, musculoskeletal chest pain, musculoskeletal discomfort, neck pain, backpain, myalgia, and bone pain.

<sup>4</sup> Includes diarrhea, gastroenteritis, colitis, enterocolitis.

<sup>5</sup> Data based on Preferred Terms since laboratory data for proteinuria were not systematically collected.

**Table 9: Laboratory Abnormalities Worsening from Baseline Occurring in  $\geq 20\%$  of Patients with NSCLC Receiving TECENTRIQ in IMpower150**

| Laboratory Abnormality         | TECENTRIQ with Bevacizumab, Paclitaxel, and Carboplatin |                 | Bevacizumab, Paclitaxel and Carboplatin |                 |
|--------------------------------|---|-----------------|---|-----------------|
|                                | All Grades (%)  | Grades 3–4 (%)  | All Grades (%)                          | Grades 3–4 (%)  |
| <b>Hematology</b>              |   |                 |   |                 |
| Anemia                         | 83  | 10              | 83                                      | 9               |
| Neutropenia                    | 52  | 31              | 45                                      | 26              |
| Lymphopenia                    | 48  | 17              | 38                                      | 13              |
| <b>Chemistry</b>               |   |                 |   |                 |
| Hyperglycemia                  | 61  | 0               | 60                                      | 0               |
| Increased BUN                  | 52  | NA <sup>1</sup> | 44                                      | NA <sup>1</sup> |
| Hypomagnesemia                 | 42  | 2               | 36                                      | 1               |
| Hypoalbuminemia                | 40  | 3               | 31                                      | 2               |
| Increased AST                  | 40  | 4               | 28                                      | 0.8             |
| Hyponatremia                   | 38  | 10              | 36                                      | 9               |
| Increased Alkaline Phosphatase | 37  | 2               | 32                                      | 1               |
| Increased ALT                  | 37  | 6               | 28                                      | 0.5             |
| Increased TSH                  | 30  | NA <sup>1</sup> | 20                                      | NA <sup>1</sup> |
| Hyperkalemia                   | 28  | 3               | 25                                      | 2               |
| Increased Creatinine           | 28  | 1               | 19                                      | 2               |
| Hypocalcemia                   | 26  | 3               | 21                                      | 3               |
| Hypophosphatemia               | 25  | 4               | 18                                      | 4               |
| Hypokalemia                    | 23  | 7               | 14                                      | 4               |
| Hyperphosphatemia              | 25  | NA <sup>1</sup> | 19                                      | NA <sup>1</sup> |

Each test incidence is based on the number of patients who had both baseline and at least one on-study laboratory measurement available: TECENTRIQ with bevacizumab, paclitaxel, and carboplatin range: 337-380); bevacizumab, paclitaxel, and carboplatin (range: 337-382). Graded per NCI CTCAE v4.0

<sup>1</sup> NA = Not applicable. NCI CTCAE does not provide a Grades 3-4 definition for these laboratory abnormalities

### *IMpower130*

The safety of TECENTRIQ with paclitaxel protein-bound and carboplatin was evaluated in IMpower130, a multicenter, international, randomized, open-label trial in which 473 chemotherapy-naïve patients with metastatic non-squamous NSCLC received TECENTRIQ 1200 mg and carboplatin AUC 6 mg/mL/min intravenously on Day 1 and paclitaxel protein-bound 100 mg/m<sup>2</sup> intravenously on Day 1, 8, and 15 of each 21-day cycle for a maximum of 4 or 6 cycles, followed by TECENTRIQ 1200 mg intravenously every 3 weeks until disease progression or unacceptable toxicity [see *Clinical Studies (14.2)*]. Among patients receiving TECENTRIQ, 55% were exposed for 6 months or longer and 3.5% were exposed for greater than one year.

Fatal adverse reactions occurred in 5.3% of patients receiving TECENTRIQ; these included pneumonia (1.1%), pulmonary embolism (0.8%), myocardial infarction (0.6%), cardiac arrest

(0.4%), pneumonitis (0.4%) and sepsis, septic shock, staphylococcal sepsis, aspiration, respiratory distress, cardiorespiratory arrest, ventricular tachycardia, death (not otherwise specified), and hepatic cirrhosis (0.2% each).

Serious adverse reactions occurred in 51% of patients receiving TECENTRIQ. The most frequent serious adverse reactions ( $\geq 2\%$ ) were pneumonia (6%), diarrhea (3%), lung infection (3%), pulmonary embolism (3%), chronic obstructive pulmonary disease exacerbation (2.5%), dyspnea (2.3%), and febrile neutropenia (1.9%).

TECENTRIQ was discontinued due to adverse reactions in 13% of patients; the most common adverse reactions leading to discontinuation were pneumonia (0.8%), pulmonary embolism (0.8%), fatigue (0.6%), dyspnea (0.6%), pneumonitis (0.6%), neutropenia (0.4%), nausea (0.4%), renal failure (0.4%), cardiac arrest (0.4%), and septic shock (0.4%).

Adverse reactions leading to interruption of TECENTRIQ occurred in 62% of patients; the most common ( $>1\%$ ) were neutropenia, thrombocytopenia, anemia, diarrhea, fatigue/asthenia, pneumonia, dyspnea, pneumonitis, pyrexia, nausea, acute kidney injury, vomiting, pulmonary embolism, arthralgia, infusion-related reaction, abdominal pain, chronic obstructive pulmonary disease exacerbation, dehydration, and hypokalemia.

Tables 10 and 11 summarize adverse reactions and laboratory abnormalities in patients receiving TECENTRIQ with paclitaxel protein-bound and carboplatin in IMpower130.

**Table 10: Adverse Reactions Occurring in  $\geq 20\%$  of Patients with NSCLC Receiving TECENTRIQ in IMpower130**

| Adverse Reaction                             | TECENTRIQ with Paclitaxel Protein-Bound and Carboplatin<br>N = 473 |                | Paclitaxel Protein-Bound and Carboplatin<br>N = 232 |                |
|--|--|----------------|---|----------------|
|  | All Grades (%)   | Grades 3–4 (%) | All Grades (%)                                      | Grades 3–4 (%) |
| <b>General</b>                               |  |                |   |                |
| Fatigue/Asthenia                             | 61   | 11             | 60  | 8              |
| <b>Gastrointestinal</b>                      |  |                |   |                |
| Nausea                                       | 50   | 3.4            | 46  | 2.2            |
| Diarrhea <sup>1</sup>                        | 43   | 6              | 32  | 6              |
| Constipation                                 | 36   | 1.1            | 31  | 0              |
| Vomiting                                     | 27   | 2.7            | 19  | 2.2            |
| <b>Musculoskeletal and Connective Tissue</b> |  |                |   |                |
| Myalgia/Pain <sup>2</sup>                    | 38   | 3              | 22  | 0.4            |
| <b>Nervous System</b>                        |  |                |   |                |
| Neuropathy <sup>3</sup>                      | 33   | 2.5            | 28  | 2.2            |
| <b>Respiratory, Thoracic and Mediastinal</b> |  |                |   |                |
| Dyspnea <sup>4</sup>                         | 32   | 4.9            | 25  | 1.3            |
| Cough  | 27   | 0.6            | 17  | 0              |
| <b>Skin and Subcutaneous Tissue</b>          |  |                |   |                |
| Alopecia                                     | 32   | 0              | 27  | 0              |
| Rash <sup>5</sup>                            | 20   | 0.6            | 11  | 0.9            |

| <b>Metabolism and Nutrition</b> |    |     |    |     |
|---------------------------------|----|-----|----|-----|
| Decreased appetite              | 30 | 2.1 | 26 | 2.2 |

Graded per NCI CTCAE v4.0

<sup>1</sup> Includes diarrhea, colitis, and gastroenteritis

<sup>2</sup> Includes back pain, pain in extremity, myalgia, musculoskeletal chest pain, bone pain, neck pain and musculoskeletal discomfort

<sup>3</sup> Includes neuropathy peripheral, peripheral sensory neuropathy, hypoesthesia, paresthesia, dysesthesia, polyneuropathy

<sup>4</sup> Includes dyspnea, dyspnea exertional and wheezing

<sup>5</sup> Includes rash, rash maculo-papular, eczema, rash pruritic, rash erythematous, dermatitis, dermatitis contact, drug eruption, seborrheic dermatitis and rash macular.

**Table 11: Laboratory Abnormalities Worsening from Baseline Occurring in ≥20% of Patients Receiving TECENTRIQ in IMpower130**

| Laboratory Abnormality         | TECENTRIQ with Paclitaxel Protein-Bound and Carboplatin<br>N = 473 |                 | Paclitaxel Protein-Bound and Carboplatin<br>N = 232 |                 |
|--------------------------------|--|-----------------|---|-----------------|
|                                | All Grades (%)   | Grades 3–4 (%)  | All Grades (%)                                      | Grades 3–4 (%)  |
| <b>Hematology</b>              |  |                 |   |                 |
| Anemia                         | 92   | 33              | 87  | 25              |
| Neutropenia                    | 75   | 50              | 67  | 39              |
| Thrombocytopenia               | 73   | 19              | 59  | 13              |
| Lymphopenia                    | 71   | 23              | 61  | 16              |
| <b>Chemistry</b>               |  |                 |   |                 |
| Hyperglycemia                  | 75   | 8               | 66  | 8               |
| Hypomagnesemia                 | 50   | 3.4             | 42  | 3.2             |
| Hyponatremia                   | 37   | 9               | 28  | 7               |
| Hypoalbuminemia                | 35   | 1.3             | 31  | 0               |
| Increased ALT                  | 31   | 2.8             | 24  | 3.9             |
| Hypocalcemia                   | 31   | 2.6             | 27  | 1.8             |
| Hypophosphatemia               | 29   | 6               | 20  | 3.2             |
| Increased AST                  | 28   | 2.2             | 24  | 1.8             |
| Increased TSH                  | 26   | NA <sup>1</sup> | 5   | NA <sup>1</sup> |
| Hypokalemia                    | 26   | 6               | 24  | 4.4             |
| Increased Alkaline Phosphatase | 25   | 2.6             | 22  | 1.3             |
| Increased Blood Creatinine     | 23   | 2.8             | 16  | 0.4             |
| Hyperphosphatemia              | 21   | NA <sup>1</sup> | 13  | NA <sup>1</sup> |

Each test incidence is based on the number of patients who had both baseline and at least one on-study laboratory measurement available: TECENTRIQ with paclitaxel protein bound and carboplatin (range: 423 - 467); paclitaxel protein bound and carboplatin (range: 218- 229). Graded per NCI CTCAE v4.0.

<sup>1</sup> NA = Not applicable. NCI CTCAE does not provide a Grades 3-4 definition for these laboratory abnormalities

### Previously Treated Metastatic NSCLC

The safety of TECENTRIQ was evaluated in OAK, a multicenter, international, randomized, open-label trial in patients with metastatic NSCLC who progressed during or following a platinum-containing regimen, regardless of PD-L1 expression [see *Clinical Studies (14.2)*]. A total of 609 patients received TECENTRIQ 1200 mg intravenously every 3 weeks until unacceptable toxicity, radiographic progression, or clinical progression or docetaxel (n=578) 75 mg/m<sup>2</sup> intravenously every 3 weeks until unacceptable toxicity or disease progression. The study excluded patients with active or prior autoimmune disease or with medical conditions that required systemic corticosteroids. The median duration of exposure was 3.4 months (0 to 26 months) in TECENTRIQ-treated patients and 2.1 months (0 to 23 months) in docetaxel-treated patients.

The study population characteristics were: median age of 63 years (25 to 85 years), 46% age 65 years or older, 62% male, 71% White, 20% Asian, 68% former smoker, 16% current smoker, and 63% had ECOG performance status of 1.

Fatal adverse reactions occurred in 1.6% of patients; these included pneumonia, sepsis, septic shock, dyspnea, pulmonary hemorrhage, sudden death, myocardial ischemia or renal failure.

Serious adverse reactions occurred in 33.5% of patients. The most frequent serious adverse reactions (>1%) were pneumonia, sepsis, dyspnea, pleural effusion, pulmonary embolism, pyrexia and respiratory tract infection.

TECENTRIQ was discontinued due to adverse reactions in 8% of patients. The most common adverse reactions leading to TECENTRIQ discontinuation were fatigue, infections and dyspnea. Adverse reactions leading to interruption of TECENTRIQ occurred in 25% of patients; the most common (>1%) were pneumonia, liver function test abnormality, dyspnea, fatigue, pyrexia, and back pain.

Tables 12 and 13 summarize adverse reactions and laboratory abnormalities, respectively, in OAK.

**Table 12: Adverse Reactions Occurring in ≥10% of Patients with NSCLC Receiving TECENTRIQ in OAK**

| Adverse Reaction                | TECENTRIQ<br>N = 609 |                   | Docetaxel<br>N = 578 |                   |
|---------------------------------|----------------------|-------------------|----------------------|-------------------|
|                                 | All Grades<br>(%)    | Grades 3-4<br>(%) | All Grades<br>(%)    | Grades 3-4<br>(%) |
| <b>General</b>                  |                      |                   |                      |                   |
| Fatigue/Asthenia <sup>1</sup>   | 44                   | 4                 | 53                   | 6                 |
| Pyrexia                         | 18                   | <1                | 13                   | <1                |
| <b>1. RESPIRATORY</b>           |                      |                   |                      |                   |
| Cough <sup>2</sup>              | 26                   | <1                | 21                   | <1                |
| Dyspnea                         | 22                   | 2.8               | 21                   | 2.6               |
| <b>Metabolism and Nutrition</b> |                      |                   |                      |                   |
| Decreased appetite              | 23                   | <1                | 24                   | 1.6               |
| <b>2. MUSCULOSKELETAL</b>       |                      |                   |                      |                   |
| Myalgia/Pain <sup>3</sup>       | 20                   | 1.3               | 20                   | <1                |
| Arthralgia                      | 12                   | 0.5               | 10                   | 0.2               |

| <b>3. GASTROINTESTINAL</b> |    |    |    |    |
|----------------------------|----|----|----|----|
| Nausea                     | 18 | <1 | 23 | <1 |
| Constipation               | 18 | <1 | 14 | <1 |
| Diarrhea                   | 16 | <1 | 24 | 2  |
| <b>4. SKIN</b>             |    |    |    |    |
| Rash <sup>4</sup>          | 12 | <1 | 10 | 0  |

Graded per NCI CTCAE v4.0

<sup>1</sup> Includes fatigue and asthenia

<sup>2</sup> Includes cough and exertional cough

<sup>3</sup> Includes musculoskeletal pain, musculoskeletal stiffness, musculoskeletal chest pain, myalgia

<sup>4</sup> Includes rash, erythematous rash, generalized rash, maculopapular rash, papular rash, pruritic rash, pustular rash, pemphigoid

**Table 13: Laboratory Abnormalities Worsening from Baseline Occurring in  $\geq 20\%$  of Patients with NSCLC Receiving TECENTRIQ in OAK**

| Laboratory Abnormality         | TECENTRIQ      |                | Docetaxel      |                |
|--------------------------------|----------------|----------------|----------------|----------------|
|                                | All Grades (%) | Grades 3-4 (%) | All Grades (%) | Grades 3-4 (%) |
| <b>Hematology</b>              |                |                |                |                |
| Anemia                         | 67             | 3              | 82             | 7              |
| Lymphocytopenia                | 49             | 14             | 60             | 21             |
| <b>Chemistry</b>               |                |                |                |                |
| Hypoalbuminemia                | 48             | 4              | 50             | 3              |
| Hyponatremia                   | 42             | 7              | 31             | 6              |
| Increased Alkaline Phosphatase | 39             | 2              | 25             | 1              |
| Increased AST                  | 31             | 3              | 16             | 0.5            |
| Increased ALT                  | 27             | 3              | 14             | 0.5            |
| Hypophosphatemia               | 27             | 5              | 23             | 4              |
| Hypomagnesemia                 | 26             | 1              | 21             | 1              |
| Increased Creatinine           | 23             | 2              | 16             | 1              |

Each test incidence is based on the number of patients who had both baseline and at least one on-study laboratory measurement available: TECENTRIQ (range: 546–585) and docetaxel (range: 532–560). Graded according to NCI CTCAE version 4.0

### Metastatic Triple Negative Breast Cancer (TNBC)

The safety of TECENTRIQ in combination with paclitaxel protein-bound was evaluated in IMpassion130, a multicenter, international, randomized, double-blinded placebo-controlled trial in patients with locally advanced or metastatic TNBC who have not received prior chemotherapy for metastatic disease [see *Clinical Studies (14.3)*]. Patients received TECENTRIQ 840 mg (n=452) or placebo (n=438) intravenously followed by paclitaxel protein-bound (100 mg/m<sup>2</sup>) intravenously. For each 28 day cycle, TECENTRIQ was administered on days 1 and 15 and paclitaxel protein-bound was administered on days 1, 8, and 15 until disease progression or unacceptable toxicity. In the safety-evaluable population, the median duration of exposure to TECENTRIQ was 5.5 months (range: 0-32 months) and paclitaxel protein-bound was 5.1 months (range: 0-31.5 months) in the TECENTRIQ and paclitaxel protein-bound arm. The median duration of exposure to placebo was 5.1 months (range: 0-25.1 months) and paclitaxel protein-bound was 5.0 months (range: 0-23.7 months) in the placebo and paclitaxel protein-bound arm.

Fatal adverse reactions occurred in 1.3%) of patients in the TECENTRIQ and paclitaxel protein-bound arm; these included septic shock, mucosal inflammation, auto-immune hepatitis, aspiration, pneumonia, pulmonary embolism.

Serious adverse reactions occurred in 23% of patients. The most frequent serious adverse reactions were pneumonia (2%), urinary tract infection (1%), dyspnea (1%), and pyrexia (1%).

Adverse reactions leading to discontinuation of TECENTRIQ occurred in 6% (29/452) of patients in the TECENTRIQ and paclitaxel protein-bound arm. The most common adverse reaction leading to TECENTRIQ discontinuation was peripheral neuropathy (<1%).

Adverse reactions leading to interruption of TECENTRIQ occurred in 31% of patients; the most common ( $\geq 2\%$ ) were neutropenia, neutrophil count decreased, hyperthyroidism, and pyrexia.

Immune-related adverse reactions requiring systemic corticosteroid therapy occurred in 13% (59/452) of patients in the TECENTRIQ and paclitaxel protein-bound arm.

Tables 14 and 15 summarize adverse reactions and selected laboratory abnormalities worsening from baseline in the TECENTRIQ treated patients.

**Table 14: Adverse Reactions Occurring in  $\geq 10\%$  of Patients with TNBC in IMpassion130**

| Adverse Reaction                     | TECENTRIQ with Paclitaxel Protein-Bound<br>N = 452 |                | Placebo with Paclitaxel Protein-Bound<br>N = 438 |                |
|--------------------------------------|--|----------------|--|----------------|
|                                      | All Grades (%)                                     | Grades 3-4 (%) | All Grades (%)                                   | Grades 3-4 (%) |
| <b>Skin and Subcutaneous Tissue</b>  |  |                |  |                |
| Alopecia                             | 56   | <1             | 58   | <1             |
| Rash                                 | 17   | <1             | 16   | <1             |
| Pruritus                             | 14   | 0              | 10   | 0              |
| <b>Nervous System</b>                |  |                |  |                |
| Peripheral neuropathies <sup>1</sup> | 47   | 9              | 44   | 5              |
| Headache                             | 23   | <1             | 22   | <1             |
| Dysgeusia                            | 14   | 0              | 14   | 0              |
| Dizziness                            | 14   | 0              | 11   | 0              |
| <b>General</b>                       |  |                |  |                |
| Fatigue                              | 47   | 4              | 45   | 3.4            |
| Pyrexia                              | 19   | <1             | 11   | 0              |
| Peripheral Edema                     | 15   | <1             | 16   | 1.4            |
| Asthenia                             | 12   | <1             | 11   | <1             |
| <b>Gastrointestinal</b>              |  |                |  |                |
| Nausea                               | 46   | 1.1            | 38   | 1.8            |
| Diarrhea                             | 33   | 1.3            | 34   | 2.1            |
| Constipation                         | 25   | <1             | 25   | <1             |
| Vomiting                             | 20   | <1             | 17   | 1.1            |
| Abdominal pain                       | 10   | <1             | 12   | <1             |

| <b>Respiratory, Thoracic, and Mediastinal</b> |    |     |     |    |
|---|----|-----|-----|----|
| Cough   | 25 | 0   | 19  | 0  |
| Dyspnea                                       | 16 | <1  | 15  | <1 |
| <b>Metabolism and Nutrition</b>               |    |     |     |    |
| Decreased Appetite                            | 20 | <1  | 18  | <1 |
| <b>Musculoskeletal and Connective Tissue</b>  |    |     |     |    |
| Arthralgia                                    | 18 | <1  | 16  | <1 |
| Back pain                                     | 15 | 1.3 | 13  | <1 |
| Myalgia                                       | 14 | <1  | 15  | <1 |
| Pain in extremity                             | 11 | <1  | 10  | <1 |
| <b>Endocrine</b>                              |    |     |     |    |
| Hypothyroidism                                | 14 | 0   | 3.4 | 0  |
| <b>Infections</b>                             |    |     |     |    |
| Urinary tract infection                       | 12 | <1  | 11  | <1 |
| Upper respiratory tract infection             | 11 | 1.1 | 9   | 0  |
| Nasopharyngitis                               | 11 | 0   | 8   | 0  |

Graded per NCI CTCAE v4.0

<sup>1</sup> Includes peripheral neuropathy, peripheral sensory neuropathy, paresthesia, and polyneuropathy

**Table 15: Laboratory Abnormalities Worsening from Baseline Occurring in  $\geq 20\%$  of Patients with TNBC in IMpassion130**

| Laboratory Abnormality         | TECENTRIQ with Paclitaxel Protein-Bound |                | Placebo in combination with Paclitaxel Protein-Bound |                |
|--------------------------------|---|----------------|--|----------------|
|                                | All Grades (%)                          | Grades 3–4 (%) | All Grades (%)                                       | Grades 3–4 (%) |
| <b>Hematology</b>              |   |                |  |                |
| Decreased Hemoglobin           | 79                                      | 3.8            | 73   | 3              |
| Decreased Leukocytes           | 76                                      | 14             | 71   | 9              |
| Decreased Neutrophils          | 58                                      | 13             | 54   | 13             |
| Decreased Lymphocytes          | 54                                      | 13             | 47   | 8              |
| Increased Prothrombin INR      | 25                                      | <1             | 25   | <1             |
| <b>Chemistry</b>               |   |                |  |                |
| Increased ALT                  | 43                                      | 6              | 34   | 2.7            |
| Increased AST                  | 42                                      | 4.9            | 34   | 3.4            |
| Decreased Calcium              | 28                                      | 1.1            | 26   | <1             |
| Decreased Sodium               | 27                                      | 4.2            | 25   | 2.7            |
| Decreased Albumin              | 27                                      | <1             | 25   | <1             |
| Increased Alkaline Phosphatase | 25                                      | 3.3            | 22   | 2.7            |
| Decreased Phosphate            | 22                                      | 3.6            | 19   | 3.7            |
| Increased Creatinine           | 21                                      | <1             | 16   | <1             |

Each test incidence is based on the number of patients who had at least one on-study laboratory measurement available: TECENTRIQ with paclitaxel protein-bound (range: 316-452); placebo with paclitaxel protein-bound (range: 299-438). Graded per NCI CTCAE v4.0, except for increased creatinine which only includes patients with creatinine increase based on upper limit of normal definition for Grade 1 events (NCI CTCAE v5.0).

## Small Cell Lung Cancer (SCLC)

The safety of TECENTRIQ with carboplatin and etoposide was evaluated in IMpower133, a randomized, multicenter, double-blind, placebo-controlled trial in which 198 patients with ES-SCLC received TECENTRIQ 1200 mg and carboplatin AUC 5 mg/mL/min on Day 1 and etoposide 100 mg/m<sup>2</sup> intravenously on Days 1, 2 and 3 of each 21-day cycle for a maximum of 4 cycles, followed by TECENTRIQ 1200 mg every 3 weeks until disease progression or unacceptable toxicity [see *Clinical Studies (14.4)*]. Among 198 patients receiving TECENTRIQ, 32% were exposed for 6 months or longer and 12% were exposed for 12 months or longer.

Fatal adverse reactions occurred in 2% of patients receiving TECENTRIQ. These included pneumonia, respiratory failure, neutropenia, and death (1 patient each).

Serious adverse reactions occurred in 37% of patients receiving TECENTRIQ. Serious adverse reactions in >2% were pneumonia (4.5%), neutropenia (3.5%), febrile neutropenia (2.5%), and thrombocytopenia (2.5%).

TECENTRIQ was discontinued due to adverse reactions in 11% of patients. The most frequent adverse reaction requiring permanent discontinuation in >2% of patients was infusion-related reactions (2.5%).

Adverse reactions leading to interruption of TECENTRIQ occurred in 59% of patients; the most common (>1%) were neutropenia (22%), anemia (9%), leukopenia (7%), thrombocytopenia (5%), fatigue (4.0%), infusion-related reaction (3.5%), pneumonia (2.0%), febrile neutropenia (1.5%), increased ALT (1.5%), and nausea (1.5%).

Tables 16 and 17 summarize adverse reactions and laboratory abnormalities, respectively, in patients who received TECENTRIQ with carboplatin and etoposide in IMpower133.

**Table 16: Adverse Reactions Occurring in ≥20% of Patients with SCLC Receiving TECENTRIQ in IMpower133**

| Adverse Reaction                    | TECENTRIQ with Carboplatin and Etoposide<br>N = 198 |                | Placebo with Carboplatin and Etoposide<br>N = 196 |                |
|-------------------------------------|---|----------------|---|----------------|
|                                     | All Grades (%)                                      | Grades 3–4 (%) | All Grades (%)                                    | Grades 3–4 (%) |
| <b>General</b>                      |   |                |   |                |
| Fatigue/asthenia                    | 39  | 5              | 33  | 3              |
| <b>Gastrointestinal</b>             |   |                |   |                |
| Nausea                              | 38  | 1              | 33  | 1              |
| Constipation                        | 26  | 1              | 30  | 1              |
| Vomiting                            | 20  | 2              | 17  | 3              |
| <b>Skin and Subcutaneous Tissue</b> |   |                |   |                |
| Alopecia                            | 37  | 0              | 35  | 0              |
| <b>Metabolism and Nutrition</b>     |   |                |   |                |
| Decreased appetite                  | 27  | 1              | 18  | 0              |

Graded per NCI CTCAE v4.0

**Table 17: Laboratory Abnormalities Worsening from Baseline Occurring in  $\geq 20\%$  of Patients with SCLC Receiving TECENTRIQ in IMpower133**

| Laboratory Abnormality         | TECENTRIQ with Carboplatin and Etoposide |                 | Placebo with Carboplatin and Etoposide |                 |
|--------------------------------|--|-----------------|--|-----------------|
|                                | All Grades (%)                           | Grades 3–4 (%)  | All Grades (%)                         | Grades 3–4 (%)  |
| <b>Hematology</b>              |  |                 |  |                 |
| Anemia                         | 94                                       | 17              | 93                                     | 19              |
| Neutropenia                    | 73                                       | 45              | 76                                     | 48              |
| Thrombocytopenia               | 58                                       | 20              | 53                                     | 17              |
| Lymphopenia                    | 46                                       | 14              | 38                                     | 11              |
| <b>Chemistry</b>               |  |                 |  |                 |
| Hyperglycemia                  | 67                                       | 10              | 65                                     | 8               |
| Increased Alkaline Phosphatase | 38                                       | 1               | 35                                     | 2               |
| Hyponatremia                   | 34                                       | 15              | 33                                     | 11              |
| Hypoalbuminemia                | 32                                       | 1               | 30                                     | 0               |
| Decreased TSH <sup>2</sup>     | 28                                       | NA <sup>1</sup> | 15                                     | NA <sup>1</sup> |
| Hypomagnesemia                 | 31                                       | 5               | 35                                     | 6               |
| Hypocalcemia                   | 26                                       | 3               | 28                                     | 5               |
| Increased ALT                  | 26                                       | 3               | 31                                     | 1               |
| Increased AST                  | 22                                       | 1               | 21                                     | 2               |
| Increased Blood Creatinine     | 22                                       | 4               | 15                                     | 1               |
| Hyperphosphatemia              | 21                                       | NA <sup>1</sup> | 23                                     | NA <sup>1</sup> |
| Increased TSH <sup>2</sup>     | 21                                       | NA <sup>1</sup> | 7                                      | NA <sup>1</sup> |

Each test incidence is based on the number of patients who had both baseline and at least one on-study laboratory measurement available: TECENTRIQ (range: 181-193); Placebo (range: 181-196). Graded per NCI CTCAE v4.0  
<sup>1</sup> NA= Not applicable. <sup>2</sup> TSH = thyroid-stimulating hormone. NCI CTCAE v4.0 does not include these laboratories.

### Hepatocellular Carcinoma (HCC)

The safety of TECENTRIQ in combination with bevacizumab was evaluated in IMbrave150, a multicenter, international, randomized, open-label trial in patients with locally advanced or metastatic or unresectable hepatocellular carcinoma who have not received prior systemic treatment [see *Clinical Studies (14.5)*]. Patients received 1,200 mg of TECENTRIQ intravenously followed by 15 mg/kg bevacizumab (n=329) every 3 weeks, or 400 mg of sorafenib (n=156) given orally twice daily, until disease progression or unacceptable toxicity. The median duration of exposure to TECENTRIQ was 7.4 months (range: 0-16 months) and to bevacizumab was 6.9 months (range: 0-16 months).

Fatal adverse reactions occurred in 4.6% of patients in the TECENTRIQ and bevacizumab arm. The most common adverse reactions leading to death were gastrointestinal and esophageal varices hemorrhage (1.2%) and infections (1.2%).

Serious adverse reactions occurred in 38% of patients in the TECENTRIQ and bevacizumab arm. The most frequent serious adverse reactions ( $\geq 2\%$ ) were gastrointestinal hemorrhage (7%), infections (6%), and pyrexia (2.1%).

Adverse reactions leading to discontinuation of TECENTRIQ occurred in 9% of patients in the TECENTRIQ and bevacizumab arm. The most common adverse reactions leading to TECENTRIQ discontinuation were hemorrhages (1.2%), including gastrointestinal, subarachnoid, and pulmonary hemorrhages; increased transaminases or bilirubin (1.2%); infusion-related reaction/cytokine release syndrome (0.9%); and autoimmune hepatitis (0.6%).

Adverse reactions leading to interruption of TECENTRIQ occurred in 41% of patients in the TECENTRIQ and bevacizumab arm; the most common ( $\geq 2\%$ ) were liver function laboratory abnormalities including increased transaminases, bilirubin, or alkaline phosphatase (8%); infections (6%); gastrointestinal hemorrhages (3.6%); thrombocytopenia/decreased platelet count (3.6%); hyperthyroidism (2.7%); and pyrexia (2.1%).

Immune-related adverse reactions requiring systemic corticosteroid therapy occurred in 12% of patients in the TECENTRIQ and bevacizumab arm.

Tables 18 and 19 summarize adverse reactions and laboratory abnormalities, respectively, in patients who received TECENTRIQ and bevacizumab in IMbrave150.

**Table 18: Adverse Reactions Occurring in  $\geq 10\%$  of Patients with HCC Receiving TECENTRIQ in IMbrave150**

| Adverse Reaction  | TECENTRIQ in combination with bevacizumab (n = 329) |                             | Sorafenib (n=156)           |                             |
|---|---|-----------------------------|-----------------------------|-----------------------------|
|   | All Grades <sup>2</sup> (%)                         | Grades 3–4 <sup>2</sup> (%) | All Grades <sup>2</sup> (%) | Grades 3–4 <sup>2</sup> (%) |
| <b>Vascular Disorders</b>                                   |   |                             |                             |                             |
| Hypertension  | 30  | 15                          | 24                          | 12                          |
| <b>General Disorders and Administration Site Conditions</b> |   |                             |                             |                             |
| Fatigue/asthenia <sup>1</sup>                               | 26  | 2                           | 32                          | 6                           |
| Pyrexia   | 18  | 0                           | 10                          | 0                           |
| <b>Renal and Urinary Disorders</b>                          |   |                             |                             |                             |
| Proteinuria   | 20  | 3                           | 7                           | 0.6                         |
| <b>Investigations</b>                                       |   |                             |                             |                             |
| Weight Decreased  | 11  | 0                           | 10                          | 0                           |
| <b>Skin and Subcutaneous Tissue Disorders</b>               |   |                             |                             |                             |
| Pruritus  | 19  | 0                           | 10                          | 0                           |
| Rash  | 12  | 0                           | 17                          | 2.6                         |
| <b>Gastrointestinal Disorders</b>                           |   |                             |                             |                             |
| Diarrhea  | 19  | 1.8                         | 49                          | 5                           |
| Constipation  | 13  | 0                           | 14                          | 0                           |
| Abdominal Pain  | 12  | 0                           | 17                          | 0                           |
| Nausea  | 12  | 0                           | 16                          | 0                           |
| Vomiting  | 10  | 0                           | 8                           | 0                           |
| <b>Metabolism and Nutrition Disorders</b>                   |   |                             |                             |                             |
| Decreased Appetite  | 18  | 1.2                         | 24                          | 3.8                         |

| <b>Respiratory, Thoracic and Mediastinal Disorders</b> |    |     |     |   |
|--|----|-----|-----|---|
| Cough  | 12 | 0   | 10  | 0 |
| Epistaxis  | 10 | 0   | 4.5 | 0 |
| <b>Injury, Poisoning and Procedural Complications</b>  |    |     |     |   |
| Infusion Related Reaction                              | 11 | 2.4 | 0   | 0 |

<sup>1</sup> Includes fatigue and asthenia

<sup>2</sup> Graded per NCI CTCAE v4.0

**Table 19: Laboratory Abnormalities Worsening from Baseline Occurring in  $\geq 20\%$  of Patients with HCC Receiving TECENTRIQ in IMbrave150**

| Laboratory Abnormality         | TECENTRIQ in combination with bevacizumab (n=329) |                             | Sorafenib (n=156)           |                             |
|--------------------------------|---|-----------------------------|-----------------------------|-----------------------------|
|                                | All Grades <sup>1</sup> (%)                       | Grades 3–4 <sup>1</sup> (%) | All Grades <sup>1</sup> (%) | Grades 3–4 <sup>1</sup> (%) |
| <b>Chemistry</b>               |   |                             |                             |                             |
| Increased AST                  | 86  | 16                          | 90                          | 16                          |
| Increased Alkaline Phosphatase | 70  | 4                           | 76                          | 4.6                         |
| Increased ALT                  | 62  | 8                           | 70                          | 4.6                         |
| Decreased Albumin              | 60  | 1.5                         | 54                          | 0.7                         |
| Decreased Sodium               | 54  | 13                          | 49                          | 9                           |
| Increased Glucose              | 48  | 9                           | 43                          | 4.6                         |
| Decreased Calcium              | 30  | 0.3                         | 35                          | 1.3                         |
| Decreased Phosphorus           | 26  | 4.7                         | 58                          | 16                          |
| Increased Potassium            | 23  | 1.9                         | 16                          | 2                           |
| Hypomagnesemia                 | 22  | 0                           | 22                          | 0                           |
| <b>Hematology</b>              |   |                             |                             |                             |
| Decreased Platelet             | 68  | 7                           | 63                          | 4.6                         |
| Decreased Lymphocytes          | 62  | 13                          | 58                          | 11                          |
| Decreased Hemoglobin           | 58  | 3.1                         | 62                          | 3.9                         |
| Increased Bilirubin            | 57  | 8                           | 59                          | 14                          |
| Decreased Leukocyte            | 32  | 3.4                         | 29                          | 1.3                         |
| Decreased Neutrophil           | 23  | 2.3                         | 16                          | 1.1                         |

Each test incidence is based on the number of patients who had both baseline and at least one on-study laboratory measurement available: TECENTRIQ plus bevacizumab (222-323) and sorafenib (90-153)

<sup>1</sup> Graded per NCI CTCAE v4.0

### Melanoma

The safety of TECENTRIQ, administered with cobimetinib and vemurafenib was evaluated in IMspire150, a double-blind, randomized (1:1), placebo-controlled study conducted in patients with previously untreated BRAF V600 mutation-positive metastatic or unresectable melanoma [see *Clinical Studies (14.5)*]. Patients received TECENTRIQ with cobimetinib and vemurafenib (N=230) or placebo with cobimetinib and vemurafenib (n=281).

Among the 230 patients who received TECENTRIQ administered with cobimetinib and vemurafenib, the median duration of exposure to TECENTRIQ was 9.2 months (range: 0-30

months) to cobimetinib was 10.0 months (range: 1-31 months) and to vemurafenib was 9.8 months (range: 1-31 months).

Fatal adverse reactions occurred in 3% of patients in the TECENTRIQ plus cobimetinib and vemurafenib arm. Adverse reactions leading to death were hepatic failure, fulminant hepatitis, sepsis, septic shock, pneumonia, and cardiac arrest.

Serious adverse reactions occurred in 45% of patients in the TECENTRIQ plus cobimetinib and vemurafenib arm. The most frequent ( $\geq 2\%$ ) serious adverse reactions were hepatotoxicity (7%), pyrexia (6%), pneumonia (4.3%), malignant neoplasms (2.2%), and acute kidney injury (2.2%).

Adverse reactions leading to discontinuation of TECENTRIQ occurred in 21% of patients in the TECENTRIQ plus cobimetinib and vemurafenib arm. The most frequent ( $\geq 2\%$ ) adverse reactions leading to TECENTRIQ discontinuation were increased ALT (2.2%) and pneumonitis (2.6%).

Adverse reactions leading to interruption of TECENTRIQ occurred in 68% of patients in the TECENTRIQ plus cobimetinib and vemurafenib arm. The most frequent ( $\geq 2\%$ ) adverse reactions leading to TECENTRIQ interruption were pyrexia (14%), increased ALT (13%), hyperthyroidism (10%), increased AST (10%), increased lipase (9%), increased amylase (7%), pneumonitis (5%), increased CPK (4.3%), diarrhea (3.5%), pneumonia (3.5%), asthenia (3%), rash (3%), influenza (3%), arthralgia (2.6%), fatigue (2.2%), dyspnea (2.2%), cough (2.2%), peripheral edema (2.2%), uveitis (2.2%), bronchitis (2.2%), hypothyroidism (2.2%), and respiratory tract infection (2.2%).

Tables 20 and 21 summarizes the incidence of adverse reactions and laboratory abnormalities in Study IMspire150.

**Table 20: Adverse Reactions Occurring in  $\geq 10\%$  of Patients on the TECENTRIQ plus Cobimetinib and Vemurafenib Arm or the Placebo plus Cobimetinib and Vemurafenib Arm and at a Higher Incidence (Between Arm Difference of  $\geq 5\%$  All Grades or  $\geq 2\%$  Grades 3-4 TECENTRIQ in IMspire150)**

| Adverse Reaction  | TECENTRIQ in combination with Cobimetinib and Vemurafenib (n=230) |               | Placebo with Cobimetinib and Vemurafenib (n=281) |               |
|---|---|---------------|--|---------------|
|   | All Grades (%)  | Grade 3-4 (%) | All Grades (%)                                   | Grade 3-4 (%) |
| <b>Skin and Subcutaneous Tissue Disorders</b>               |   |               |  |               |
| Rash <sup>1</sup>   | 75  | 27            | 72   | 23            |
| Pruritus  | 26  | <1            | 17   | <1            |
| Photosensitivity reaction                                   | 21  | <1            | 25   | 3.2           |
| <b>General Disorders and Administration Site Conditions</b> |   |               |  |               |
| Fatigue <sup>2</sup>  | 51  | 3             | 45   | 1.8           |
| Pyrexia <sup>3</sup>  | 49  | 1.7           | 35   | 2.1           |
| Edema <sup>4</sup>  | 26  | <1            | 21   | 0             |
| <b>Gastrointestinal Disorders</b>                           |   |               |  |               |
| Hepatotoxicity <sup>5</sup>                                 | 50  | 21            | 36   | 13            |
| Nausea  | 30  | <1            | 32   | 2.5           |
| Stomatitis <sup>6</sup>                                     | 23  | 1.3           | 15   | <1            |
| <b>Musculoskeletal and Connective Tissue Disorders</b>      |   |               |  |               |
| Musculoskeletal pain <sup>7</sup>                           | 62  | 4.3           | 48   | 3.2           |
| <b>Endocrine Disorders</b>                                  |   |               |  |               |
| Hypothyroidism <sup>8</sup>                                 | 22  | 0             | 10   | 0             |
| Hyperthyroidism   | 18  | <1            | 8  | 0             |
| <b>Injury, Poisoning and Procedural Complications</b>       |   |               |  |               |

|  |    |     |    |    |
|--|----|-----|----|----|
| Infusion related reaction <sup>9</sup>                 | 10 | 2.6 | 8  | <1 |
| <b>Respiratory, Thoracic and Mediastinal Disorders</b> |    |     |    |    |
| Pneumonitis <sup>10</sup>                              | 12 | 1.3 | 6  | <1 |
| <b>Vascular Disorders</b>                              |    |     |    |    |
| Hypertension <sup>11</sup>                             | 17 | 10  | 18 | 7  |

<sup>1</sup> Includes rash, rash maculo-papular, dermatitis acneiform, rash macular, rash erythematous, eczema, skin exfoliation, rash papular, rash pustular, palmar-plantar erythrodysesthesia syndrome, dermatitis, dermatitis contact, erythema multiforme, rash pruritic, drug eruption, nodular rash, dermatitis allergic, exfoliative rash, dermatitis exfoliative generalised and rash morbilliform

<sup>2</sup> Includes fatigue, asthenia and malaise

<sup>3</sup> Includes pyrexia and hyperpyrexia

<sup>4</sup> Includes edema peripheral, lymphoedema, oedema, face oedema, eyelid oedema, periorbital oedema, lip oedema and generalised oedema

<sup>5</sup> Includes alanine aminotransferase increased, aspartate aminotransferase increased, blood bilirubin increased, transaminases increased, hepatitis, hepatic enzyme increased, hepatotoxicity, hypertransaminasaemia, bilirubin conjugated increased, hepatocellular injury, hyperbilirubinaemia, liver function test increased, hepatic failure, hepatitis fulminant and liver function test abnormal

<sup>6</sup> Includes stomatitis, mucosal inflammation, aphthous ulcer, mouth ulceration, cheilitis and glossitis

<sup>7</sup> Includes arthralgia, myalgia, pain in extremity, back pain, musculoskeletal pain, arthritis, neck pain, musculoskeletal chest pain, musculoskeletal stiffness, bone pain, spinal pain, immune-mediated arthritis, joint stiffness and non-cardiac chest pain

<sup>8</sup> Includes hypothyroidism and blood thyroid stimulating hormone increased

<sup>9</sup> Includes infusion related reaction and hypersensitivity

<sup>10</sup> Includes pneumonitis and interstitial lung disease

<sup>11</sup> Includes hypertension, blood pressure increased, hypertensive crisis

Clinically important adverse reactions in < 10% of patients who received TECENTRIQ plus cobimetinib and vemurafenib were:

*Cardiac Disorders:* Arrhythmias, ejection fraction decreased, electrocardiogram QT prolonged

*Eye Disorders:* Uveitis

*Gastrointestinal disorders:* Pancreatitis

*Infections and infestations:* Pneumonia, urinary tract infection

*Metabolism and nutrition disorders:* Hyperglycemia

*Nervous system Disorders:* Dizziness, dysgeusia, syncope

*Respiratory, thoracic and mediastinal disorders:* Dyspnea, oropharyngeal pain

*Skin and Subcutaneous Tissue Disorders:* Vitiligo

**Table 21: Laboratory Abnormalities Worsening from Baseline Occurring in  $\geq 20\%$  of Patients Receiving TECENTRIQ plus Cobimetinib and Vemurafenib Arm or the Placebo plus Cobimetinib and Vemurafenib Arm and at a Higher Incidence (Between Arm Difference of  $\geq 5\%$  All Grades or  $\geq 2\%$  Grades 3-4) in IMspire150**

| Laboratory Abnormality | TECENTRIQ in combination with Cobimetinib and Vemurafenib (n=230) |               | Placebo with Cobimetinib and Vemurafenib (n=281) |               |
|------------------------|---|---------------|--|---------------|
|                        | All Grades (%)  | Grade 3-4 (%) | All Grades (%)                                   | Grade 3-4 (%) |
| <b>Hematology</b>      |   |               |  |               |
| Decreased Lymphocytes  | 80  | 24            | 72   | 17            |
| Decreased Hemoglobin   | 77  | 2.6           | 72   | 2.2           |
| Decreased Platelet     | 34  | 1.3           | 24   | 0.4           |
| Decreased Neutrophils  | 26  | 2.2           | 19   | 1.5           |

| <b>Chemistry</b>                                   |    |                 |    |                 |
|--|----|-----------------|----|-----------------|
| Increased Creatine Kinase                          | 88 | 22              | 81 | 18              |
| Increased AST                                      | 80 | 13              | 68 | 6               |
| Increased ALT                                      | 79 | 18              | 62 | 12              |
| Increased Triacylglycerol Lipase                   | 75 | 46              | 62 | 35              |
| Increased Alkaline Phosphatase                     | 73 | 6               | 63 | 2.9             |
| Decreased Phosphorus                               | 67 | 22              | 64 | 14              |
| Increased Amylase                                  | 51 | 13              | 45 | 13              |
| Increased Blood Urea Nitrogen                      | 47 | NA <sup>1</sup> | 37 | NA <sup>1</sup> |
| Decreased Albumin                                  | 43 | 0.9             | 34 | 1.5             |
| Increased Bilirubin                                | 42 | 3.1             | 33 | 0.7             |
| Decreased Calcium                                  | 41 | 1.3             | 28 | 0               |
| Decreased Sodium                                   | 40 | 5               | 34 | 7               |
| Decreased Thyroid-Stimulating Hormone              | 38 | NA <sup>1</sup> | 23 | NA <sup>1</sup> |
| Increased Thyroid-Stimulating Hormone <sup>2</sup> | 37 | NA <sup>1</sup> | 33 | NA <sup>1</sup> |
| Decreased Potassium                                | 36 | 5               | 22 | 4.3             |
| Increased Triiodothyronine                         | 33 | NA <sup>1</sup> | 18 | NA <sup>1</sup> |
| Increased Free Thyroxine                           | 32 | NA <sup>1</sup> | 21 | NA <sup>1</sup> |
| Decreased Total Triiodothyronine                   | 32 | NA <sup>1</sup> | 8  | NA <sup>1</sup> |
| Increased Potassium                                | 29 | 1.3             | 19 | 1.4             |
| Decreased Triiodothyronine                         | 27 | NA <sup>1</sup> | 21 | NA <sup>1</sup> |
| Increased Sodium                                   | 20 | 0               | 13 | 0.4             |

Graded per NCI CTCAE v4.0.

Each test incidence is based on the number of patients who had both baseline and at least one on-study laboratory measurement available: TECENTRIQ plus cobimetinib and vemurafenib (28-277), placebo plus cobimetinib and vemurafenib arm (25-230).

<sup>1</sup> NA= Not applicable. NCI CTCAE v4.0 does not include these laboratories.

<sup>2</sup> Increased Thyroid Stimulating Hormone has a difference <5% (All Grades) between arms and is included for clinical completeness.

## 6.2 Immunogenicity

As with all therapeutic proteins, there is a potential for immunogenicity. The detection of antibody formation is highly dependent on the sensitivity and specificity of the assay. Additionally, the observed incidence of antibody (including neutralizing antibody) positivity in an assay may be influenced by several factors including assay methodology, sample handling, timing of sample collection, concomitant medications, and underlying disease. For these reasons, comparison of the incidence of antibodies to atezolizumab in the studies described above with the incidence of antibodies in other studies or to other products may be misleading.

Among 565 patients with NSCLC in OAK, 30% tested positive for treatment-emergent anti-drug antibodies (ADA) at one or more post-dose time points. The median onset time to ADA formation was 3 weeks. The ability of these binding ADA to neutralize atezolizumab is unknown. Patients who tested positive for treatment-emergent ADA also had decreased systemic atezolizumab exposure [see *Clinical Pharmacology (12.3)*]. Exploratory analyses showed that the subset of patients who were ADA positive by week 4 (21%; 118/560) appeared to have less efficacy (effect on overall survival) as compared to patients who tested negative for treatment-emergent ADA by week 4 [see *Clinical Studies (14.2)*]. The presence of ADA did not have a clinically significant effect on the incidence or severity of adverse reactions.

Among 275 patients with urothelial carcinoma in IMvigor210 (Cohort 2), 42% tested positive for treatment-emergent ADA at one or more post-dose time points. Among 111 patients in IMvigor210 (Cohort 1), 48% tested positive for treatment-emergent ADA at one or more post-dose time points. Patients who tested positive for treatment-emergent ADA also had decreased systemic atezolizumab exposures. The presence of ADA did not have a clinically significant effect on the incidence or severity of adverse reactions.

Among 364 ADA-evaluable patients with NSCLC who received TECENTRIQ with bevacizumab, paclitaxel and carboplatin in IMpower150, 36% (n=132) tested positive for treatment-emergent ADA at one or more post-dose time points and 83% of these 132 patients tested ADA positive prior to receiving the second dose of atezolizumab. The ability of these binding ADA to neutralize atezolizumab is unknown. Patients who tested positive for treatment-emergent ADA had lower systemic atezolizumab exposure as compared to patients who were ADA negative [see *Clinical Pharmacology (12.3)*]. The presence of ADA did not increase the incidence or severity of adverse reactions [see *Clinical Studies (14.2)*].

Among 434 patients with TNBC in IMpassion130, 13% tested positive for treatment-emergent ADA at one or more post-dose time points. Among 178 patients in PD-L1 positive subgroup with TNBC in IMpassion130, 12% tested positive for treatment-emergent ADA at one or more post-dose time points. Patients who tested positive for treatment-emergent ADA had decreased systemic atezolizumab exposure [see *Clinical Pharmacology (12.3)*]. There are insufficient numbers of patients in the PD-L1 positive subgroup with ADA to determine whether ADA alters the efficacy of atezolizumab. The presence of ADA did not have a clinically significant effect on the incidence or severity of adverse reactions.

Among 315 ADA-evaluable patients with HCC who received TECENTRIQ and bevacizumab in IMbrave150, 28% (n=88) tested positive for treatment-emergent ADA at one or more post-dose time points and 66% (58/88) of these 88 patients tested ADA-positive prior to receiving the third dose of TECENTRIQ. The ability of these binding ADA to neutralize atezolizumab is unknown. Patients who tested positive for treatment-emergent ADA had lower systemic atezolizumab exposure as compared to patients who were ADA-negative [see *Clinical Pharmacology (12.3)*]. Exploratory analyses showed that the subset of patients who were ADA-positive by week 6 (20%; 58/288) appeared to have less efficacy (effect on overall survival) as compared to patients who tested negative for treatment-emergent ADA by week 6; [see *Clinical Studies (14.5)*]. The presence of ADA did not have a clinically significant effect on the incidence or severity of adverse reactions.

Among 218 ADA-evaluable patients with melanoma who received TECENTRIQ in combination with cobimetinib and vemurafenib in IMspire150, 13% (n=29) tested positive for treatment-emergent ADA at one or more post-dose time points. Patients who tested positive for treatment-emergent ADA had decreased systemic atezolizumab exposure [see *Clinical Pharmacology (12.3)*]. There are insufficient numbers of patients with positive ADA to determine whether ADA alters the efficacy or incidence or severity of adverse reactions.

#### Reporting of suspected adverse reactions

Reporting suspected adverse reactions after authorisation of the medicinal product is important. It allows continued monitoring of the benefit/risk balance of the medicinal product. Any suspected adverse events should be reported to the Ministry of Health according to the National Regulation by using an online form: <http://sideeffects.health.gov.il>

## 8 USE IN SPECIFIC POPULATIONS

### 8.1 Pregnancy

#### Risk Summary

Based on its mechanism of action [*see Clinical Pharmacology (12.1)*], TECENTRIQ can cause fetal harm when administered to a pregnant woman. There are no available data on the use of TECENTRIQ in pregnant women.

Animal studies have demonstrated that inhibition of the PD-L1/PD-1 pathway can lead to increased risk of immune-related rejection of the developing fetus resulting in fetal death (*see Data*). Advise females of reproductive potential of the potential risk to a fetus.

In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2% to 4% and 15% to 20%, respectively.

#### Data

##### *Animal Data*

Animal reproduction studies have not been conducted with TECENTRIQ to evaluate its effect on reproduction and fetal development. A literature-based assessment of the effects on reproduction demonstrated that a central function of the PD-L1/PD-1 pathway is to preserve pregnancy by maintaining maternal immune tolerance to a fetus. Blockage of PD-L1 signaling has been shown in murine models of pregnancy to disrupt tolerance to a fetus and to result in an increase in fetal loss; therefore, potential risks of administering TECENTRIQ during pregnancy include increased rates of abortion or stillbirth. As reported in the literature, there were no malformations related to the blockade of PD-L1/PD-1 signaling in the offspring of these animals; however, immune-mediated disorders occurred in PD-1 and PD-L1 knockout mice. Based on its mechanism of action, fetal exposure to atezolizumab may increase the risk of developing immune-mediated disorders or altering the normal immune response.

### 8.2 Lactation

#### Risk Summary

There is no information regarding the presence of atezolizumab in human milk, the effects on the breastfed infant, or the effects on milk production. As human IgG is excreted in human milk, the potential for absorption and harm to the infant is unknown. Because of the potential for serious adverse reactions in breastfed infants from TECENTRIQ, advise women not to breastfeed during treatment and for at least 5 months after the last dose.

### 8.3 Females and Males of Reproductive Potential

#### Pregnancy Testing

Verify pregnancy status in females of reproductive potential prior to initiating TECENTRIQ [*see Use in Specific Populations (8.1)*].

#### Contraception

##### *Females*

Based on its mechanism of action, TECENTRIQ can cause fetal harm when administered to a pregnant woman [*see Use in Specific Populations (8.1)*]. Advise females of reproductive potential to use effective contraception during treatment with TECENTRIQ and for at least 5 months following the last dose.

## Infertility

### *Females*

Based on animal studies, TECENTRIQ may impair fertility in females of reproductive potential while receiving treatment [see *Nonclinical Toxicology (13.1)*].

### **8.4 Pediatric Use**

The safety and effectiveness of TECENTRIQ have not been established in pediatric patients.

The safety and antitumor activity of TECENTRIQ were assessed but not established in a single-arm, multi-center, multi-cohort trial (NCT02541604) in 60 pediatric patients aged 7 months to <17 years with relapsed or progressive solid tumors and lymphomas. No new safety signals were observed in pediatric patients in this study.

In pediatric patients who received TECENTRIQ 15 mg/kg with a maximum dose of 1200 mg every 3 weeks, the steady-state exposure (AUC) of atezolizumab in pediatric patients aged 12 years or older was comparable to that in adult patients who received TECENTRIQ 1200 mg every 3 weeks, while the exposure trended lower in pediatric patients less than 12 years old.

### **8.5 Geriatric Use**

Of 3040 patients with urothelial carcinoma, lung cancer, triple-negative breast cancer, hepatocellular carcinoma, and melanoma who were treated with TECENTRIQ in clinical studies, 43% were 65 years and over and 12% were 75 years and over. No overall differences in safety or effectiveness were observed between patients aged 65 years or older and younger patients.

## **11 DESCRIPTION**

Atezolizumab is a programmed cell death ligand 1 (PD-L1) blocking antibody. Atezolizumab is an Fc-engineered, humanized, non-glycosylated IgG1 kappa immunoglobulin that has a calculated molecular mass of 145 kDa.

TECENTRIQ (atezolizumab) injection for intravenous use is a sterile, preservative-free, colorless to slightly yellow solution in single-dose vials. Each mL contains 60mg/ml of atezolizumab and is formulated in sucrose, L-histidine, glacial acetic acid and polysorbate 20, with a pH 5.8.

## **12 CLINICAL PHARMACOLOGY**

### **12.1 Mechanism Of Action**

Pd-11 may be expressed on tumor cells and/or tumor infiltrating immune cells and can contribute to the inhibition of the anti-tumor immune response in the tumor microenvironment. Binding of pd-11 to the pd-1 and b7.1 receptors found on t cells and antigen presenting cells suppresses cytotoxic t-cell activity, t-cell proliferation and cytokine production.

Atezolizumab is a monoclonal antibody that binds to pd-11 and blocks its interactions with both pd-1 and b7.1 receptors. This releases the pd-11/pd-1 mediated inhibition of the immune response, including activation of the anti-tumor immune response without inducing antibody-dependent cellular cytotoxicity. In syngeneic mouse tumor models, blocking pd-11 activity resulted in decreased tumor growth.

In mouse models of cancer, dual inhibition of the PD-1/PD-L1 and MAPK pathways suppresses tumor growth and improves tumor immunogenicity through increased antigen presentation and T cell infiltration and activation compared to targeted therapy alone.

## 12.3 Pharmacokinetics

Patients' exposure to atezolizumab increased dose proportionally over the dose range of 1 mg/kg to 20 mg/kg, including a dose of 1200 mg administered every 3 weeks. The clearance (CV%) was 0.20 L/day (29%), the volume of distribution at steady state was 6.9 L, and the terminal half-life was 27 days. Steady state was achieved after 6 to 9 weeks following multiple doses. The systemic accumulation ratio for every 2 weeks administration and every 3 weeks administration was 3.3- and 1.9- fold, respectively. Atezolizumab clearance was found to decrease over time, with a mean maximal reduction (CV%) from baseline value of approximately 17% (41%); however, the decrease in clearance was not considered clinically relevant.

### Specific Populations

Age (21 to 89 years), body weight, sex, albumin levels, tumor burden, region or race, mild or moderate renal impairment [estimated glomerular filtration rate (eGFR) 30 to 89 mL/min/1.73 m<sup>2</sup>], mild hepatic impairment (bilirubin  $\leq$  ULN and AST  $>$  ULN or bilirubin  $>$  1 to 1.5  $\times$  ULN and any AST), moderate hepatic impairment (bilirubin  $>$ 1.5 to 3x ULN and any AST), level of PD-L1 expression, or performance status had no clinically significant effect on the systemic exposure of atezolizumab. Across clinical trials with TECENTRIQ, median atezolizumab clearance in patients who tested positive for treatment-emergent anti-drug antibodies (ADA) was 22% (range: 18% to 49%) higher as compared to atezolizumab clearance in patients who tested negative for treatment-emergent ADA.

### Drug Interaction Studies

The drug interaction potential of atezolizumab is unknown.

## 13 NONCLINICAL TOXICOLOGY

### 13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

No studies have been performed to test the potential of atezolizumab for carcinogenicity or genotoxicity.

Animal fertility studies have not been conducted with atezolizumab; however, an assessment of the male and female reproductive organs was included in a 26-week, repeat-dose toxicity study in cynomolgus monkeys. Weekly administration of atezolizumab to female monkeys at the highest dose tested caused an irregular menstrual cycle pattern and a lack of newly formed corpora lutea in the ovaries. This effect occurred at an estimated AUC approximately 6 times the AUC in patients receiving the recommended dose and was reversible. There was no effect on the male monkey reproductive organs.

### 13.2 Animal Toxicology and/or Pharmacology

In animal models, inhibition of PD-L1/PD-1 signaling increased the severity of some infections and enhanced inflammatory responses. M. tuberculosis-infected PD-1 knockout mice exhibit markedly decreased survival compared with wild-type controls, which correlated with increased bacterial proliferation and inflammatory responses in these animals. PD-L1 and PD-1 knockout mice and mice receiving PD-L1 blocking antibody have also shown decreased survival following infection with lymphocytic choriomeningitis virus.

## 14 CLINICAL STUDIES

### 14.1 Urothelial Carcinoma

#### Cisplatin-Ineligible Patients with Locally Advanced or Metastatic Urothelial Carcinoma

The efficacy of TECENTRIQ was investigated in IMvigor210 (Cohort 1) (NCT02951767), a multicenter, open-label, single-arm trial that included 119 patients with locally advanced or

metastatic urothelial carcinoma who were ineligible for cisplatin-containing chemotherapy and were either previously untreated or had disease progression at least 12 months after neoadjuvant or adjuvant chemotherapy. Patients were considered cisplatin-ineligible if they met any one of the following criteria at study entry: impaired renal function [creatinine clearance (CLcr) of 30 to 59 mL/min], Eastern Cooperative Oncology Group (ECOG) performance status (PS) of 2, hearing loss of  $\geq 25$  decibels (dB) at two contiguous frequencies, or Grades 2-4 peripheral neuropathy. This study excluded patients who had: a history of autoimmune disease; active or corticosteroid-dependent brain metastases; administration of a live, attenuated vaccine within 28 days prior to enrollment; or administration of systemic immunostimulatory agents within 6 weeks or systemic immunosuppressive medications within 2 weeks prior to enrollment. Patients received TECENTRIQ 1200 mg as an intravenous infusion every 3 weeks until unacceptable toxicity or disease progression. Tumor response assessments were conducted every 9 weeks for the first 54 weeks and every 12 weeks thereafter. Major efficacy outcome measures included confirmed overall response rate (ORR) as assessed by independent review facility (IRF) using Response Evaluation Criteria in Solid Tumors (RECIST v1.1), duration of response (DoR) and overall survival (OS).

In this study, the median age was 73 years, 81% were male, and 91% were White. Thirty-five percent of patients had non-bladder urothelial carcinoma and 66% had visceral metastases. Eighty percent of patients had an ECOG PS of 0 or 1. Reasons for ineligibility for cisplatin-containing chemotherapy were: 70% had impaired renal function, 20% had an ECOG PS of 2, 14% had a hearing loss of  $\geq 25$  dB, and 6% had Grades 2-4 peripheral neuropathy at baseline. Twenty percent of patients had disease progression following prior platinum-containing neoadjuvant or adjuvant chemotherapy.

Tumor specimens were evaluated prospectively using the VENTANA PD-L1 (SP142) Assay at a central laboratory, and the results were used to define subgroups for pre-specified analyses. Of the 119 patients, 27% were classified as having PD-L1 expression of  $\geq 5\%$  (defined as PD-L1 stained tumor-infiltrating immune cells [IC] covering  $\geq 5\%$  of the tumor area). The remaining 73% of patients were classified as having PD-L1 expression of  $< 5\%$  (PD-L1 stained tumor-infiltrating IC covering  $< 5\%$  of the tumor area).

Among the 32 patients with PD-L1 expression of  $\geq 5\%$ , median age was 67 years, 81% were male, 19% female, and 88% were White. Twenty-eight percent of patients had non-bladder urothelial carcinoma and 56% had visceral metastases. Seventy-two percent of patients had an ECOG PS of 0 or 1. Reasons for ineligibility for cisplatin-containing chemotherapy were: 66% had impaired renal function, 28% had an ECOG PS of 2, 16% had a hearing loss  $\geq 25$  dB, and 9% had Grades 2-4 peripheral neuropathy at baseline. Thirty-one percent of patients had disease progression following prior platinum-containing neoadjuvant or adjuvant chemotherapy.

Confirmed ORR in all patients and the two PD-L1 subgroups are summarized in Table 22. The median follow-up time for this study was 14.4 months. In 24 patients with disease progression following neoadjuvant or adjuvant therapy, the ORR was 33% (95% CI: 16%, 55%).

**Table 22: Efficacy Results in IMvigor210 (Cohort 1)**

|  | All Patients | PD-L1 Expression Subgroups                                |  |
|--|--------------|---|--|
|  | N = 119      | PD-L1 Expression of $< 5\%$ in ICs <sup>1</sup><br>N = 87 | PD-L1 Expression of $\geq 5\%$ in ICs <sup>1</sup><br>N = 32 |
| <b>Number of IRF-assessed Confirmed Responders</b> | 28           | 19  | 9  |

|  |                    |                    |                    |
|--|--------------------|--------------------|--------------------|
| <b>ORR % (95% CI)</b>  | 23.5% (16.2, 32.2) | 21.8% (13.7, 32)   | 28.1% (13.8, 46.8) |
| Complete Response  | 6.7%               | 6.9%               | 6.3%               |
| Partial Response   | 16.8%              | 14.9%              | 21.9%              |
| <b>Median DoR, months<br/>(range)</b>  | NR<br>(3.7, 16.6+) | NR<br>(3.7, 16.6+) | NR<br>(8.1, 15.6+) |
| NR = Not reached<br>+ Denotes a censored value<br><sup>1</sup> PD-L1 expression in tumor-infiltrating immune cells (ICs) |                    |                    |                    |

IMvigor130 (NCT02807636) is an ongoing multicenter, randomized study in previously untreated patients with metastatic urothelial carcinoma who are eligible for platinum-containing chemotherapy. The study contains three arms: TECENTRIQ monotherapy, TECENTRIQ with platinum-based chemotherapy (i.e., cisplatin or carboplatin with gemcitabine), and platinum-based chemotherapy alone (comparator). Both cisplatin-eligible and cisplatin-ineligible patients are included in the study. Tumor specimens were evaluated prospectively using the VENTANA PD-L1 (SP142) Assay at a central laboratory. The independent Data Monitoring Committee (iDMC) for the study conducted a review of early data and found that patients classified as having PD-L1 expression of <5% when treated with TECENTRIQ monotherapy had decreased survival compared to those who received platinum-based chemotherapy. The iDMC recommended closure of the monotherapy arm to further accrual of patients with low PD-L1 expression, however, no other changes were recommended for the study, including any change of therapy for patients who had already been randomized to and were receiving treatment in the monotherapy arm.

#### Previously Treated Locally Advanced or Metastatic Urothelial Carcinoma

The efficacy of TECENTRIQ was investigated in IMvigor210 (Cohort 2) (NCT02108652), a multicenter, open-label, single-arm trial that included 310 patients with locally advanced or metastatic urothelial carcinoma who had disease progression during or following a platinum-containing chemotherapy regimen or who had disease progression within 12 months of treatment with a platinum-containing neoadjuvant or adjuvant chemotherapy regimen. This study excluded patients who had: a history of autoimmune disease, active or corticosteroid-dependent brain metastases, administration of a live, attenuated vaccine within 28 days prior to enrollment, or administration of systemic immunostimulatory agents within 6 weeks or systemic immunosuppressive medications within 2 weeks prior to enrollment. Patients received TECENTRIQ 1200 mg intravenously every 3 weeks until unacceptable toxicity or either radiographic or clinical progression. Tumor response assessments were conducted every 9 weeks for the first 54 weeks and every 12 weeks thereafter. Major efficacy outcome measures included confirmed ORR as assessed by IRF using RECIST v1.1 and DoR.

In this study, the median age was 66 years, 78% were male, 91% of patients were White. Twenty-six percent had non-bladder urothelial carcinoma and 78% of patients had visceral metastases. Sixty-two percent of patients had an ECOG PS of 1 and 35% of patients had a baseline CLcr < 60 mL/min. Nineteen percent of patients had disease progression following prior platinum-containing neoadjuvant or adjuvant chemotherapy. Forty-one percent of patients had received 2 or more prior systemic regimens in the metastatic setting. Seventy-three percent of patients received prior cisplatin, 26% had prior carboplatin, and 1% were treated with other platinum-based regimens.

Tumor specimens were evaluated prospectively using the VENTANA PD-L1 (SP142) Assay at a central laboratory and the results were used to define subgroups for pre-specified analyses. Of

the 310 patients, 32% were classified as having PD-L1 expression of  $\geq 5\%$ . The remaining 68% of patients were classified as having PD-L1 expression of  $< 5\%$ .

Confirmed ORR and median DOR in all patients and the two PD-L1 subgroups are summarized in Table 23. The median follow-up time for this study was 32.9 months. In 59 patients with disease progression following neoadjuvant or adjuvant therapy, the ORR was 22.0% (95% CI: 12.3%, 34.7%).

**Table 23: Efficacy Results in IMvigor210 (Cohort 2)**

|   | All Patients          | PD-L1 Expression Subgroups                                |  |
|---|-----------------------|---|--|
|   | N = 310               | PD-L1 Expression of $< 5\%$ in IC <sup>1</sup><br>N = 210 | PD-L1 Expression of $\geq 5\%$ in IC <sup>1</sup><br>N = 100 |
| <b>Number of IRF-assessed Confirmed Responders</b>                    | 46                    | 20  | 26   |
| <b>ORR % (95% CI)</b>   | 14.8% (11.2, 19.3)    | 9.5% (5.9, 14.3)  | 26% (17.7, 35.7)   |
| Complete Response   | 5.5%                  | 2.4%  | 12.0%  |
| Partial Response  | 9.4%                  | 7.1%  | 14.0%  |
| <b>Median DOR, months (range)</b>                                     | 27.7<br>(2.1+, 33.4+) | 20.9<br>(2.1+, 33.4+)                                     | 29.7<br>(4.2, 31.2+)   |
| + Denotes a censored value  |                       |   |  |
| <sup>1</sup> PD-L1 expression in tumor-infiltrating immune cells (IC) |                       |   |  |

## 14.2 Non-Small Cell Lung Cancer

### Metastatic Chemotherapy-Naïve NSCLC with High PD-L1 Expression

The efficacy of TECENTRIQ was evaluated in IMpower110 (NCT02409342), a multicenter, international, randomized, open-label trial in patients with stage IV NSCLC whose tumors express PD-L1 (PD-L1 stained  $\geq 1\%$  of tumor cells [TC  $\geq 1\%$ ] or PD-L1 stained tumor-infiltrating immune cells [IC] covering  $\geq 1\%$  of the tumor area [IC  $\geq 1\%$ ]), who had received no prior chemotherapy for metastatic disease. PD-L1 tumor status was determined based on immunohistochemistry (IHC) testing using the VENTANA PD-L1 (SP142) Assay. The evaluation of efficacy is based on the subgroup of patients with high PD-L1 expression (TC  $\geq 50\%$  or IC  $\geq 10\%$ ), excluding those with EGFR or ALK genomic tumor aberrations. The trial excluded patients with a history of autoimmune disease, administration of a live attenuated vaccine within 28 days prior to randomization, active or untreated CNS metastases, administration of systemic immunostimulatory agents within 4 weeks or systemic immunosuppressive medications within 2 weeks prior to randomization.

Randomization was stratified by sex, ECOG performance status, histology (non-squamous vs. squamous) and PD-L1 expression (TC  $\geq 1\%$  and any IC vs. TC  $< 1\%$  and IC  $\geq 1\%$ ). Patients were randomized (1:1) to receive one of the following treatment arms:

- Arm A: TECENTRIQ 1200 mg every 3 weeks until disease progression or unacceptable toxicity
- Arm B: Platinum-based chemotherapy

Arm B platinum-based chemotherapy regimens for non-squamous NSCLC consisted of cisplatin (75 mg/m<sup>2</sup>) and pemetrexed (500 mg/m<sup>2</sup>) OR carboplatin (AUC 6 mg/mL/min) and pemetrexed

(500 mg/m<sup>2</sup>) on Day 1 of each 21-day cycle for a maximum of 4 or 6 cycles followed by pemetrexed 500 mg/m<sup>2</sup> until disease progression or unacceptable toxicity.

Arm B platinum-based chemotherapy regimens for squamous NSCLC consisted of cisplatin (75 mg/m<sup>2</sup>) on Day 1 with gemcitabine (1250 mg/m<sup>2</sup>) on Days 1 and 8 of each 21-day cycle OR carboplatin (AUC 5 mg/mL/min) on Day 1 with gemcitabine (1000 mg/m<sup>2</sup>) on Days 1 and 8 of each 21-day cycle for a maximum of 4 or 6 cycles followed by best supportive care until disease progression or unacceptable toxicity.

Administration of TECENTRIQ was permitted beyond RECIST-defined disease progression. Tumor assessments were conducted every 6 weeks for the first 48 weeks following Cycle 1, Day 1 and then every 9 weeks thereafter. Tumor specimens were evaluated prospectively using the VENTANA PD-L1 (SP142) Assay at a central laboratory and the results were used to define subgroups for pre-specified analyses.

The major efficacy outcome measure was overall survival (OS) sequentially tested in the following subgroups of patients, excluding those with EGFR or ALK genomic tumor aberrations: TC ≥50% or IC ≥10%; TC ≥5% or IC ≥5%; and TC ≥1% or IC ≥1%.

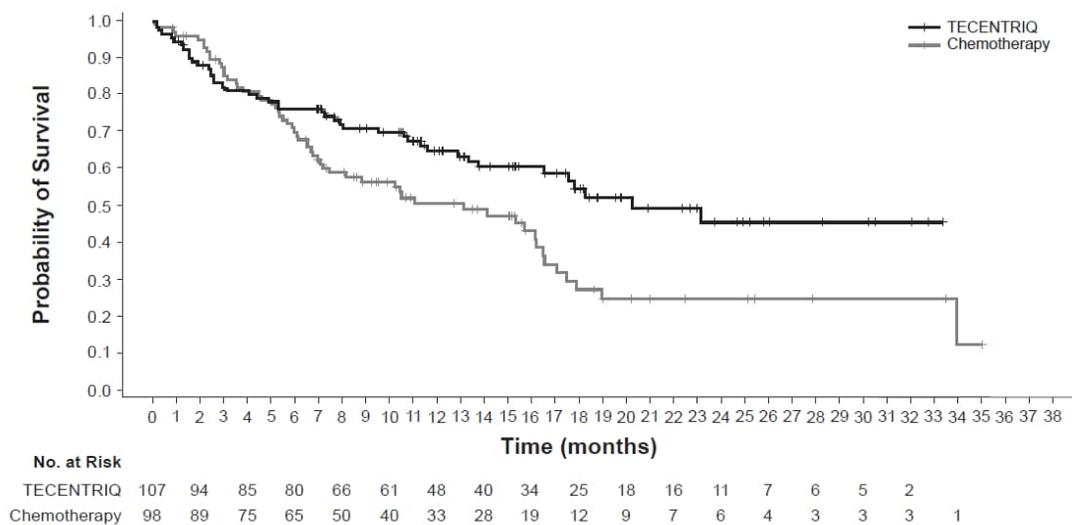
Among the 205 chemotherapy-naïve patients with stage IV NSCLC with high PD-L1 expression (TC ≥ 50% or IC ≥ 10%) excluding those with EGFR or ALK genomic tumor aberrations, the median age was 65.0 years (range: 33 to 87), and 70% of patients were male. The majority of patients were White (82%) and Asian (17%). Baseline ECOG performance status was 0 (36%) or 1 (64%); 88% were current or previous smokers; and 76% of patients had non-squamous disease while 24% of patients had squamous disease.

The trial demonstrated a statistically significant improvement in OS for patients with high PD-L1 expression (TC ≥50% or IC ≥10%) at the time of the OS interim analysis. There was no statistically significant difference in OS for the other two PD-L1 subgroups (TC ≥5% or IC ≥5%; and TC ≥1% or IC ≥1%) at the interim or final analyses. Efficacy results for patients with NSCLC with high PD-L1 expression are presented in Table 24 and Figure 1.

**Table 24: Efficacy Results from IMpower110 in Patients with NSCLC with High PD-L1 Expression (TC ≥ 50% or IC ≥ 10%) and without EGFR or ALK Genomic Tumor Aberrations**

|   | Arm A: TECENTRIQ<br>N = 107 | Arm B: Platinum-Based<br>Chemotherapy<br>N = 98 |
|---|-----------------------------|---|
| <b>Overall Survival<sup>1</sup></b>   |                             |   |
| Deaths (%)  | 44 (41%)                    | 57 (58%)  |
| Median, months  | 20.2                        | 13.1  |
| (95% CI)  | (16.5, NE)                  | (7.4, 16.5)                                     |
| Hazard ratio <sup>2</sup> (95% CI)  | 0.59 (0.40, 0.89)           |   |
| p-value <sup>3</sup>  | 0.0106 <sup>4</sup>         |   |
| <sup>1</sup> Based on OS interim analysis. The median survival follow-up time in patients was 15.7 months.<br><sup>2</sup> Stratified by sex and ECOG performance status<br><sup>3</sup> Based on the stratified log-rank test compared to Arm A<br><sup>4</sup> Compared to the allocated alpha of 0.0413 (two-sided) for this interim analysis.<br>CI=confidence interval; NE=not estimable |                             |   |

**Figure 1: Kaplan-Meier Plot of Overall Survival in IMpower110 in Patients with NSCLC with High PD-L1 Expression (TC ≥ 50% or IC ≥ 10%) and Without EGFR or ALK Genomic Tumor Aberrations**



Investigator-assessed PFS showed an HR of 0.63 (95% CI: 0.45, 0.88), with median PFS of 8.1 months (95% CI: 6.8, 11.0) in the TECENTRIQ arm and 5 months (95% CI: 4.2, 5.7) in the platinum-based chemotherapy arm. The investigator-assessed confirmed ORR was 38% (95% CI: 29%, 48%) in the TECENTRIQ arm and 29% (95% CI: 20%, 39%) in the platinum-based chemotherapy arm.

### Metastatic Chemotherapy-Naive Non-Squamous NSCLC

#### *IMpower150*

The efficacy of TECENTRIQ with bevacizumab, paclitaxel, and carboplatin was evaluated in IMpower150 (NCT02366143), a multicenter, international, randomized (1:1:1), open-label trial in patients with metastatic non-squamous NSCLC. Patients with stage IV non-squamous NSCLC who had received no prior chemotherapy for metastatic disease but could have received prior EGFR or ALK kinase inhibitor if appropriate, regardless of PD-L1 or T-effector gene (tGE) status and ECOG performance status 0 or 1 were eligible. The trial excluded patients with a history of autoimmune disease, administration of a live attenuated vaccine within 28 days prior to randomization, active or untreated CNS metastases, administration of systemic immunostimulatory agents within 4 weeks or systemic immunosuppressive medications within 2 weeks prior to randomization, or clear tumor infiltration into the thoracic great vessels or clear cavitation of pulmonary lesions as seen on imaging. Randomization was stratified by sex, presence of liver metastases, and PD-L1 expression status on tumor cells (TC) and tumor-infiltrating immune cells (IC) as follows: TC3 and any IC vs. TC0/1/2 and IC2/3 vs. TC0/1/2 and IC0/1. Patients were randomized to one of the following three treatment arms.

- Arm A: TECENTRIQ 1200 mg, paclitaxel 175 mg/m<sup>2</sup> or 200 mg/m<sup>2</sup> and carboplatin AUC 6 mg/mL/min on Day 1 of each 21-day cycle for a maximum of 4 or 6 cycles
- Arm B: TECENTRIQ 1200 mg, bevacizumab 15 mg/kg, paclitaxel 175 mg/m<sup>2</sup> or 200 mg/m<sup>2</sup>, and carboplatin AUC 6 mg/mL/min on Day 1 of each 21-day cycle for a maximum of 4 or 6 cycles
- Arm C: bevacizumab 15 mg/kg, paclitaxel 175 mg/m<sup>2</sup> or 200 mg/m<sup>2</sup>, and carboplatin AUC 6 mg/mL/min on Day 1 of each 21-day cycle for a maximum of 4 or 6 cycles

Patients who had not experienced disease progression following the completion or cessation of platinum-based chemotherapy, received:

- Arm A: TECENTRIQ 1200 mg intravenously on Day 1 of each 21-day cycle until disease progression or unacceptable toxicity
- Arm B: TECENTRIQ 1200 mg and bevacizumab 15 mg/kg intravenously on Day 1 of each 21-day cycle until disease progression or unacceptable toxicity
- Arm C: bevacizumab 15 mg/kg intravenously on Day 1 of each 21-day cycle until disease progression or unacceptable toxicity

Tumor assessments were conducted every 6 weeks for the first 48 weeks following Cycle 1, Day 1 and then every 9 weeks thereafter. Tumor specimens were evaluated prior to randomization for PD-L1 tumor expression using the VENTANA PD-L1 (SP142) assay at a central laboratory. Tumor tissue was collected at baseline for expression of tGE signature and evaluation was performed using a clinical trial assay in a central laboratory prior to the analysis of efficacy outcome measures.

Major efficacy outcome measures for comparison of Arms B and C were progression free survival (PFS) by RECIST v1.1 in the tGE-WT (patients with high expression of T-effector gene signature [tGE], excluding those with EGFR- and ALK-positive NSCLC [WT]) and in the ITT-WT subpopulations and overall survival (OS) in the ITT-WT subpopulation. Additional efficacy outcome measures for comparison of Arms B and C or Arms A and C were PFS and OS in the ITT population, OS in the tGE-WT subpopulation, and ORR/DoR in the tGE-WT and ITT-WT subpopulations.

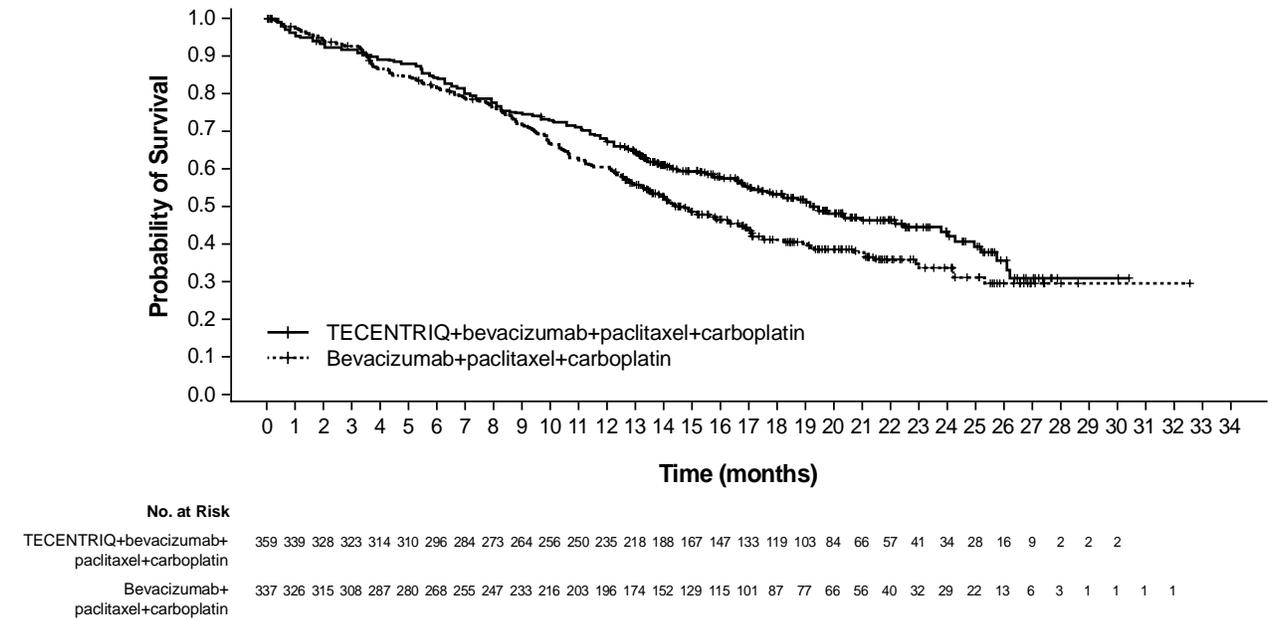
A total of 1202 patients were enrolled across the three arms of whom 1045 were in the ITT-WT subpopulation and 447 were in the tGE-WT subpopulation. The demographic information is limited to the 800 patients enrolled in Arms B and C where efficacy has been demonstrated. The median age was 63 years (range: 31 to 90), and 60% of patients were male. The majority of patients were White (82%), 13% of patients were Asian, 10% were Hispanic, and 2% of patients were Black. Clinical sites in Asia (enrolling 13% of the study population) received paclitaxel at a dose of 175 mg/m<sup>2</sup> while the remaining 87% received paclitaxel at a dose of 200 mg/m<sup>2</sup>. Approximately 14% of patients had liver metastases at baseline, and most patients were current or previous smokers (80%). Baseline ECOG performance status was 0 (43%) or 1 (57%). PD-L1 was TC3 and any IC in 12%, TC0/1/2 and IC2/3 in 13%, and TC0/1/2 and IC0/1 in 75%. The demographics for the 696 patients in the ITT-WT subpopulation were similar to the ITT population except for the absence of patients with EGFR- or ALK-positive NSCLC.

The trial demonstrated a statistically significant improvement in PFS between Arms B and C in both the tGE-WT and ITT-WT subpopulations, but did not demonstrate a significant difference for either subpopulation between Arms A and C based on the final PFS analyses. In the interim analysis of OS, a statistically significant improvement was observed for Arm B compared to Arm C, but not for Arm A compared to Arm C. Efficacy results for the ITT-WT subpopulation are presented in Table 25 and Figure 2.

**Table 25: Efficacy Results in ITT-WT Population in IMpower150**

|   | <b>Arm C: Bevacizumab,<br/>Paclitaxel and<br/>Carboplatin</b><br><br>N = 337 | <b>Arm B: TECENTRIQ<br/>with Bevacizumab,<br/>Paclitaxel, and<br/>Carboplatin</b><br><br>N = 359 | <b>Arm A: TECENTRIQ<br/>with Paclitaxel, and<br/>Carboplatin</b><br><br>N = 349 |
|---|--|--|---|
| <b>Overall Survival<sup>1</sup></b>   |  |  |   |
| Deaths (%)  | 197 (59%)  | 179 (50%)  | 179 (51%)   |
| Median, months  | 14.7   | 19.2   | 19.4  |
| (95% CI)  | (13.3, 16.9)   | (17.0, 23.8)   | (15.7, 21.3)  |
| Hazard ratio <sup>2</sup> (95% CI)  | ---  | 0.78 (0.64, 0.96)  | 0.84 (0.72, 1.08)   |
| p-value <sup>3</sup>  | ---  | 0.016 <sup>4</sup>   | 0.204 <sup>5</sup>  |
| <b>Progression-Free Survival<sup>6</sup></b>  |  |  |   |
| Number of events (%)  | 247 (73%)  | 247 (69%)  | 245 (70%)   |
| Median, months  | 7.0  | 8.5  | 6.7   |
| (95% CI)  | (6.3, 7.9)   | (7.3, 9.7)   | (5.6, 6.9)  |
| Hazard ratio <sup>2</sup> (95% CI)  | ---  | 0.71 (0.59, 0.85)  | 0.94 (0.79, 1.13)   |
| p-value <sup>3</sup>  | ---  | 0.0002 <sup>7</sup>  | 0.5219  |
| <b>Objective Response Rate<sup>6</sup></b>  |  |  |   |
| Number of responders (%)  | 142 (42%)  | 196 (55%)  | 150 (43%)   |
| (95% CI)  | (37, 48)   | (49, 60)   | (38, 48)  |
| Complete Response   | 3 (1%)   | 14 (4%)  | 9 (3%)  |
| Partial Response  | 139 (41%)  | 182 (51%)  | 141 (40%)   |
| <b>Duration of Response<sup>6</sup></b>   | n = 142  | n = 196  | n = 150   |
| Median, months  | 6.5  | 10.8   | 9.5   |
| (95% CI)  | (5.6, 7.6)   | (8.4, 13.9)  | (7.0, 13.0)   |
| <sup>1</sup> Based on OS interim analysis .<br><sup>2</sup> Stratified by sex, presence of liver metastases, and PD-L1 expression status on TC and IC<br><sup>3</sup> Based on the stratified log-rank test compared to Arm C<br><sup>4</sup> Compared to the allocated $\alpha=0.0174$ (two sided) for this interim analysis.<br><sup>5</sup> Compared to the allocated $\alpha=0.0128$ (two sided) for this interim analysis.<br><sup>6</sup> As determined by independent review facility (IRF) per RECIST v1.1 (Response Evaluation Criteria in Solid Tumors v1.1)<br><sup>7</sup> Compared to the allocated $\alpha=0.006$ (two sided) for the final PFS analysis.<br>CI=confidence interval |  |  |   |

**Figure 2: Kaplan-Meier Curves for Overall Survival in ITT-WT Population in IMpower150**



Exploratory analyses showed that the subset of patients in the four drug regimen arm who were ADA positive by week 4 (30%) appeared to have similar efficacy (effect on overall survival) as compared to patients who tested negative for treatment-emergent ADA by week 4 (70%) [see *Adverse Reactions (6.2), Clinical Pharmacology (12.3)*]. In an exploratory analysis, propensity score matching was conducted to compare ADA positive patients in the TECENTRIQ, bevacizumab, paclitaxel, and carboplatin arm with a matched population in the bevacizumab, paclitaxel, and carboplatin arm. Similarly ADA negative patients in the TECENTRIQ, bevacizumab, paclitaxel, and carboplatin arm were compared with a matched population in the bevacizumab, paclitaxel, and carboplatin arm. Propensity score matching factors were: baseline sum of longest tumor size (BSLD), baseline ECOG, baseline albumin, baseline LDH, sex, tobacco history, metastatic site, TC level, and IC level. The hazard ratio comparing the ADA-positive subgroup with its matched control was 0.69 (95% CI: 0.44, 1.07). The hazard ratio comparing the ADA-negative subgroup with its matched control was 0.64 (95% CI: 0.46, 0.90).

### IMpower130

The efficacy of TECENTRIQ with paclitaxel protein-bound and carboplatin was evaluated in IMpower130 (NCT02367781), a multicenter, randomized (2:1), open-label trial in patients with stage IV non-squamous NSCLC. Patients with Stage IV non-squamous NSCLC who had received no prior chemotherapy for metastatic disease, but could have received prior EGFR or ALK kinase inhibitor, if appropriate, were eligible. The trial excluded patients with history of autoimmune disease, administration of live attenuated vaccine within 28 days prior to randomization, administration of immunostimulatory agents within 4 weeks or systemic immunosuppressive medications within 2 weeks prior to randomization, and active or untreated CNS metastases. Randomization was stratified by sex, presence of liver metastases, and PD-L1 tumor expression according to the VENTANA PD-L1 (SP142) assay as follows: TC3 and any IC vs. TC0/1/2 and IC2/3 vs. TC0/1/2 and IC0/1. Patients were randomized to one of the following treatment regimens:

- TECENTRIQ 1200 mg on Day 1, paclitaxel protein-bound 100 mg/m<sup>2</sup> on Days 1, 8, and 15, and carboplatin AUC 6 mg/mL/min on Day 1 of each 21-day cycle for a maximum of 4 or 6

cycles followed by TECENTRIQ 1200 mg once every 3 weeks until disease progression or unacceptable toxicity, or

- Paclitaxel protein-bound 100 mg/m<sup>2</sup> on Days 1, 8 and 15 and carboplatin AUC 6 mg/mL/min on Day 1 of each 21-day cycle for a maximum of 4 or 6 cycles followed by best supportive care or pemetrexed.

Tumor assessments were conducted every 6 weeks for the first 48 weeks, then every 9 weeks thereafter. Major efficacy outcome measures were PFS by RECIST v1.1 and OS in the subpopulation of patients evaluated for and documented to have no EGFR or ALK genomic tumor aberrations (ITT-WT).

A total of 724 patients were enrolled; of these, 681 (94%) were in the ITT-WT population. The median age was 64 years (range: 18 to 86) and 59% were male. The majority of patients were white (90%), 2% of patients were Asian, 5% were Hispanic, and 4% were Black. Baseline ECOG performance status was 0 (41%) or 1 (58%). Most patients were current or previous smokers (90%). PD-L1 tumor expression was TC0/1/2 and IC0/1 in 73%; TC3 and any IC in 14%; and TC0/1/2 and IC2/3 in 13%.

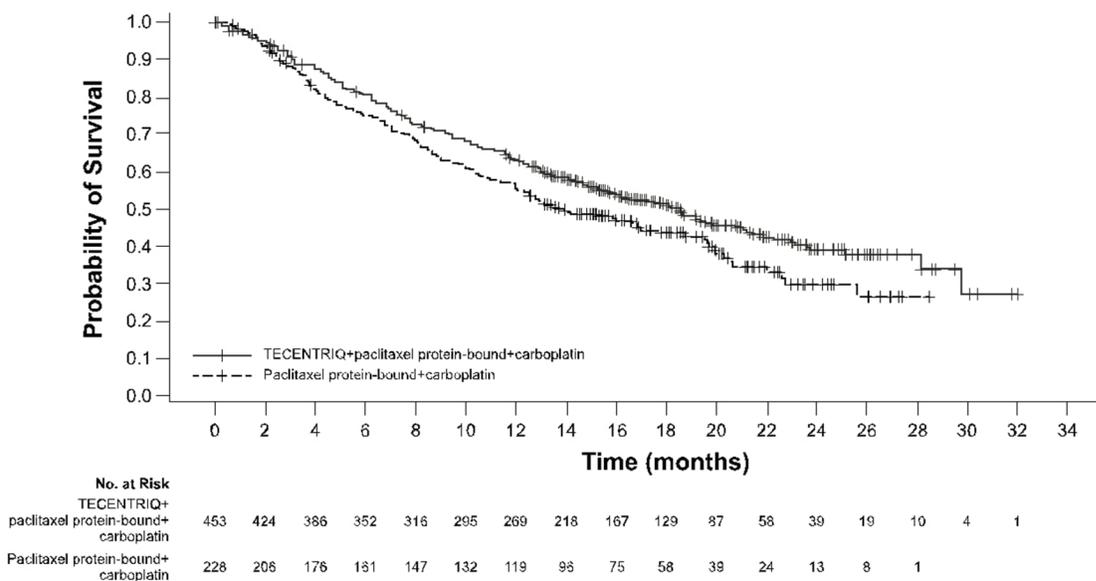
Efficacy results for the ITT-WT population are presented in Table 26 and Figure 3.

**Table 26: Efficacy Results from IMpower130**

|   | <b>TECENTRIQ with<br/>Paclitaxel Protein-Bound<br/>and Carboplatin</b> | <b>Paclitaxel Protein-Bound<br/>and Carboplatin</b> |
|---|--|---|
| <b>Overall Survival<sup>1</sup></b>   | n=453  | n=228   |
| Deaths (%)  | 228 (50%)  | 131 (57%)   |
| Median, months  | 18.6   | 13.9  |
| (95% CI)  | (15.7, 21.1)   | (12.0, 18.7)  |
| Hazard ratio <sup>2</sup> (95% CI)  | 0.80 (0.64, 0.99)  |   |
| p-value <sup>3</sup>  | 0.0384 <sup>4</sup>  |   |
| <b>Progression-Free Survival<sup>6</sup></b>  | n=453  | n=228   |
| Number of events (%)  | 330 (73%)  | 177 (78%)   |
| Median, months  | 7.2  | 6.5   |
| (95% CI)  | (6.7, 8.3)   | (5.6, 7.4)  |
| Hazard ratio <sup>2</sup> (95% CI)  | 0.75 (0.63, 0.91)  |   |
| p-value <sup>3</sup>  | 0.0024 <sup>5</sup>  |   |
| <b>Overall Response Rate<sup>6,7</sup></b>  | n=453  | n=228   |
| Number of responders (%)  | 207 (46%)  | 74 (32%)  |
| (95% CI)  | (41, 50)   | (26, 39)  |
| Complete Response   | 22 (5%)  | 2 (1%)  |
| Partial Response  | 185 (41%)  | 72 (32%)  |
| <b>Duration of Response<sup>6,7</sup></b>   | n=207  | n=74  |
| Median, months  | 10.8   | 7.8   |
| (95% CI)  | (9.0, 14.4)  | (6.8, 10.9)   |
| <sup>1</sup> Based on OS interim analysis   |  |   |
| <sup>2</sup> Stratified by sex and PD-L1 tumor expression on tumor cells (TC) and tumor infiltrating cells (IC) |  |   |
| <sup>3</sup> Based on the stratified log-rank test  |  |   |
| <sup>4</sup> Compared to the allocated $\alpha=0.0428$ (two sided) for this interim analysis                    |  |   |
| <sup>5</sup> Compared to the allocated $\alpha=0.006$ (two sided) for the final PFS analysis                    |  |   |

<sup>6</sup>As determined by independent review facility (IRF) per RECIST v1.1 (Response Evaluation Criteria in Solid Tumors v1.1)  
<sup>7</sup>Confirmed response  
 CI=confidence interval

**Figure 3: Kaplan-Meier Curves for Overall Survival in IMpower130**



### Previously Treated Metastatic NSCLC

The efficacy of TECENTRIQ was evaluated in a multicenter, international, randomized (1:1), open-label study (OAK; NCT02008227) conducted in patients with locally advanced or metastatic NSCLC whose disease progressed during or following a platinum-containing regimen. Patients with a history of autoimmune disease, symptomatic or corticosteroid-dependent brain metastases, or requiring systemic immunosuppression within 2 weeks prior to enrollment were ineligible. Randomization was stratified by PD-L1 expression tumor-infiltrating immune cells (IC), the number of prior chemotherapy regimens (1 vs. 2), and histology (squamous vs. non-squamous).

Patients were randomized to receive TECENTRIQ 1200 mg intravenously every 3 weeks until unacceptable toxicity, radiographic progression, or clinical progression or docetaxel 75 mg/m<sup>2</sup> intravenously every 3 weeks until unacceptable toxicity or disease progression. Tumor assessments were conducted every 6 weeks for the first 36 weeks and every 9 weeks thereafter. Major efficacy outcome measure was overall survival (OS) in the first 850 randomized patients and OS in the subgroup of patients with PD-L1-expressing tumors (defined as  $\geq 1\%$  PD-L1 expression on tumor cells [TC] or immune cells [IC]). Additional efficacy outcome measures were OS in all randomized patients (n = 1225), OS in subgroups based on PD-L1 expression, overall response rate (ORR), and progression free survival as assessed by the investigator per RECIST v.1.1.

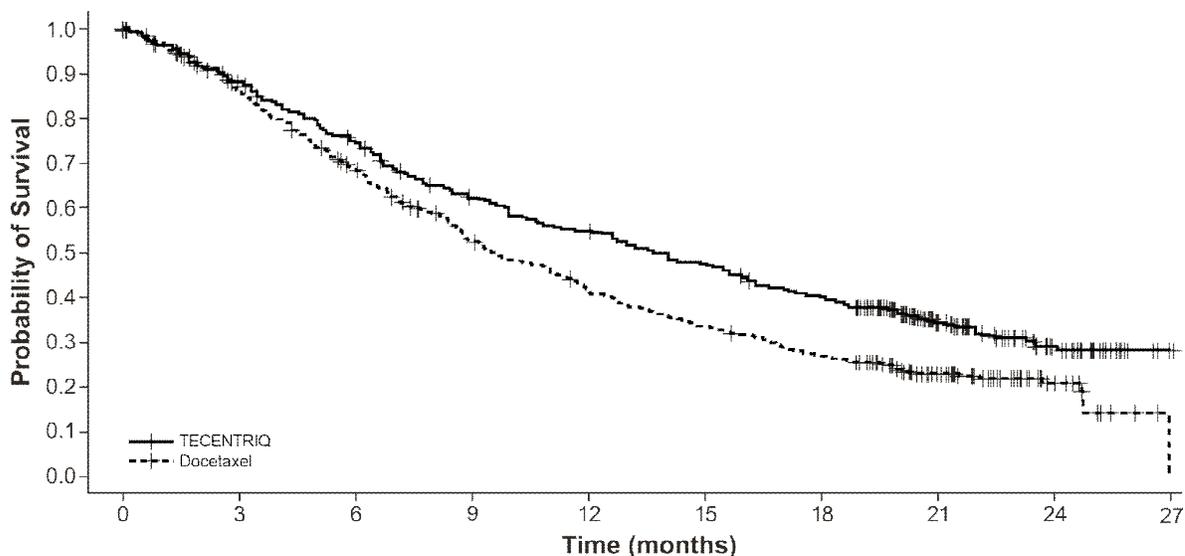
Among the first 850 randomized patients, the median age was 64 years (33 to 85 years) and 47% were  $\geq 65$  years old; 61% were male; 70% were White and 21% were Asian; 15% were current smokers and 67% were former smokers; and 37% had baseline ECOG PS of 0 and 63% had a baseline ECOG PS of 1. Nearly all (94%) had metastatic disease, 74% had non-squamous histology, 75% had received only one prior platinum-based chemotherapy regimen, and 55% of patients had PD-L1-expressing tumors.

Efficacy results are presented in Table 27 and Figure 4.

**Table 27: Efficacy Results in OAK**

|  | <b>TECENTRIQ</b>    | <b>Docetaxel</b> |
|--|---------------------|------------------|
| <b>Overall Survival in first 850 patients</b>  |                     |                  |
| Number of patients   | N=425               | N=425            |
| Deaths (%)   | 271 (64%)           | 298 (70%)        |
| Median, months   | 13.8                | 9.6              |
| (95% CI)   | (11.8, 15.7)        | (8.6, 11.2)      |
| Hazard ratio <sup>1</sup> (95% CI)   | 0.74 (0.63, 0.87)   |                  |
| p-value <sup>2</sup>   | 0.0004 <sup>3</sup> |                  |
| <b>Progression-Free Survival</b>   |                     |                  |
| Number of Patients   | N=425               | N=425            |
| Events (%)   | 380 (89%)           | 375 (88%)        |
| Progression (%)  | 332 (78%)           | 290 (68%)        |
| Deaths (%)   | 48 (11%)            | 85 (20%)         |
| Median, months   | 2.8                 | 4.0              |
| (95% CI)   | (2.6, 3.0)          | (3.3, 4.2)       |
| Hazard ratio <sup>1</sup> (95% CI)   | 0.95 (0.82, 1.10)   |                  |
| <b>Overall Response Rate<sup>4</sup></b>   |                     |                  |
| Number of Patients   | N=425               | N=425            |
| ORR, n (%)   | 58 (14%)            | 57 (13%)         |
| (95% CI)   | (11%, 17%)          | (10%, 17%)       |
| Complete Response  | 6 (1%)              | 1 (0.2%)         |
| Partial Response   | 52 (12%)            | 56 (13%)         |
| <b>Duration of Response<sup>3</sup></b>  |                     |                  |
|  | N=58                | N=57             |
| Median, months   | 16.3                | 6.2              |
| (95% CI)   | (10.0, NE)          | (4.9, 7.6)       |
| <b>Overall Survival in all 1225 patients</b>   |                     |                  |
| Number of patients   | N=613               | N=612            |
| Deaths (%)   | 384 (63%)           | 409 (67%)        |
| Median, months   | 13.3                | 9.8              |
| (95% CI)   | (11.3, 14.9)        | (8.9, 11.3)      |
| Hazard ratio <sup>1</sup> (95% CI)   | 0.79 (0.69, 0.91)   |                  |
| p-value <sup>2</sup>   | 0.0013 <sup>5</sup> |                  |
| <sup>1</sup> Stratified by PD-L1 expression in tumor infiltrating immune cells, the number of prior chemotherapy regimens, and histology<br><sup>2</sup> Based on the stratified log-rank test<br><sup>3</sup> Compared to the pre-specified allocated $\alpha$ of 0.03 for this analysis<br><sup>4</sup> Per RECIST v1.1 (Response Evaluation Criteria in Solid Tumors v1.1)<br><sup>5</sup> Compared to the allocated $\alpha$ of 0.0177 for this interim analysis based on 86% information using O'Brien-Fleming boundary<br>CI=confidence interval; NE=not estimable |                     |                  |

**Figure 4: Kaplan-Meier Curves of Overall Survival in the First 850 Patients Randomized in OAK**



| No. Patients at Risk | 0   | 3   | 6   | 9   | 12  | 15  | 18  | 21  | 24  | 27  |     |     |     |     |     |     |     |     |     |     |     |    |    |    |    |    |   |   |
|----------------------|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|----|----|----|----|----|---|---|
| TECENTRIQ            | 425 | 407 | 382 | 363 | 342 | 326 | 305 | 279 | 260 | 248 | 234 | 223 | 218 | 205 | 198 | 188 | 175 | 163 | 157 | 141 | 116 | 74 | 54 | 41 | 28 | 15 | 4 | 1 |
| Docetaxel            | 425 | 390 | 365 | 336 | 311 | 286 | 263 | 236 | 219 | 195 | 179 | 168 | 151 | 140 | 132 | 123 | 116 | 104 | 98  | 90  | 70  | 51 | 37 | 28 | 16 | 6  | 3 |   |

Tumor specimens were evaluated prospectively using the VENTANA PD-L1 (SP142) Assay at a central laboratory and the results were used to define the PD-L1 expression subgroups for pre-specified analyses. Of the 850 patients, 16% were classified as having high PD-L1 expression, defined as having PD-L1 expression on  $\geq 50\%$  of TC or  $\geq 10\%$  of IC. In an exploratory efficacy subgroup analysis of OS based on PD-L1 expression, the hazard ratio was 0.41 (95% CI: 0.27, 0.64) in the high PD-L1 expression subgroup and 0.82 (95% CI: 0.68, 0.98) in patients who did not have high PD-L1 expression.

Exploratory analyses showed that the subset of patients who were ADA positive by week 4 (21%) appeared to have less efficacy (effect on overall survival) as compared to patients who tested negative for treatment-emergent ADA by week 4 (79%) [see *Adverse Reactions (6.2)*, *Clinical Pharmacology (12.3)*]. ADA positive patients by week 4 appeared to have similar OS compared to docetaxel-treated patients. In an exploratory analysis, propensity score matching was conducted to compare ADA positive patients in the atezolizumab arm with a matched population in the docetaxel arm and ADA negative patients in the atezolizumab arm with a matched population in the docetaxel arm. Propensity score matching factors were: baseline sum of longest tumor size (BSLD), baseline ECOG, histology (squamous vs. non-squamous), baseline albumin, baseline LDH, gender, tobacco history, metastases status (advanced or local), metastatic site, TC level, and IC level. The hazard ratio comparing the ADA positive subgroup with its matched control was 0.89 (95% CI: 0.61, 1.3). The hazard ratio comparing the ADA negative subgroup with its matched control was 0.68 (95% CI: 0.55, 0.83).

### 14.3 Locally Advanced or Metastatic Triple-Negative Breast Cancer

The efficacy of TECENTRIQ in combination with paclitaxel protein-bound was investigated in IMpassion130 (NCT02425891), a multicenter, international, double-blinded, placebo-controlled, randomized (1:1) trial that included 902 unresectable locally advanced or metastatic triple-negative breast cancer patients that had not received prior chemotherapy for metastatic disease. The trial excluded patients with a history of autoimmune disease, administration of a live attenuated vaccine within 4 weeks prior to randomization, administration of systemic immunostimulatory agents within 4 weeks or systemic immunosuppressive medications within 2 weeks prior to randomization; or untreated or corticosteroid-dependent brain metastases. Randomization was stratified by presence of liver metastases, prior taxane treatment, and by PD-

L1 expression status in tumor infiltrating immune cells (IC) (PD-L1 stained tumor-infiltrating immune cells [IC] <1% of tumor area vs.  $\geq$  1% of the tumor area) by the VENTANA PD-L1 (SP142) Assay. Of the 902 patients in the intent to treat population (ITT), 41% (369 patients) were classified as PD-L1 expression  $\geq$  1%. Patients were randomized to receive TECENTRIQ 840 mg or placebo intravenously on Days 1 and 15 of every 28-day cycle with paclitaxel protein-bound 100 mg/m<sup>2</sup> intravenously on Days 1, 8 and 15 of every 28-day cycle. Patients received treatment until radiographic disease progression per RECIST v1.1, or unacceptable toxicity. Tumor assessments were performed every 8 weeks ( $\pm$  1 week) for the first 12 months after Cycle 1, day 1 and every 12 weeks ( $\pm$  1 week) thereafter. Major efficacy outcomes were investigator-assessed progression free survival (PFS) (per RECIST v1.1) and overall survival (OS) in the ITT population and PD-L1 expressing patient population.

In IMpassion130, the median age was 55 years (range: 20-86). Overall, most patients were women (99.6%) and the majority of patients were white (68%), Asian (18%), Black or African American (7%), and American Indian or Alaskan Native (4.4%). The demographic and baseline disease characteristics of the study population were well balanced between the treatment arms. Baseline ECOG performance status was 0 (58%) or 1 (41%). Overall, 41% of enrolled patients had PD-L1 expression  $\geq$  1%, 27% had liver metastases and 7% brain metastases at baseline. Approximately half the patients had received a taxane (51%) or anthracycline (54%) in the (neo)adjuvant setting. Patient demographics and baseline tumor disease in the PD-L1 expressing population were generally representative of the broader study population.

Tumor specimens (archival or fresh) were evaluated prospectively using the VENTANA PD-L1 (SP142) Assay at a central laboratory and the results were used as a stratification factor for randomization and to define the PD-L1 expression subgroups for pre-specified analyses. .

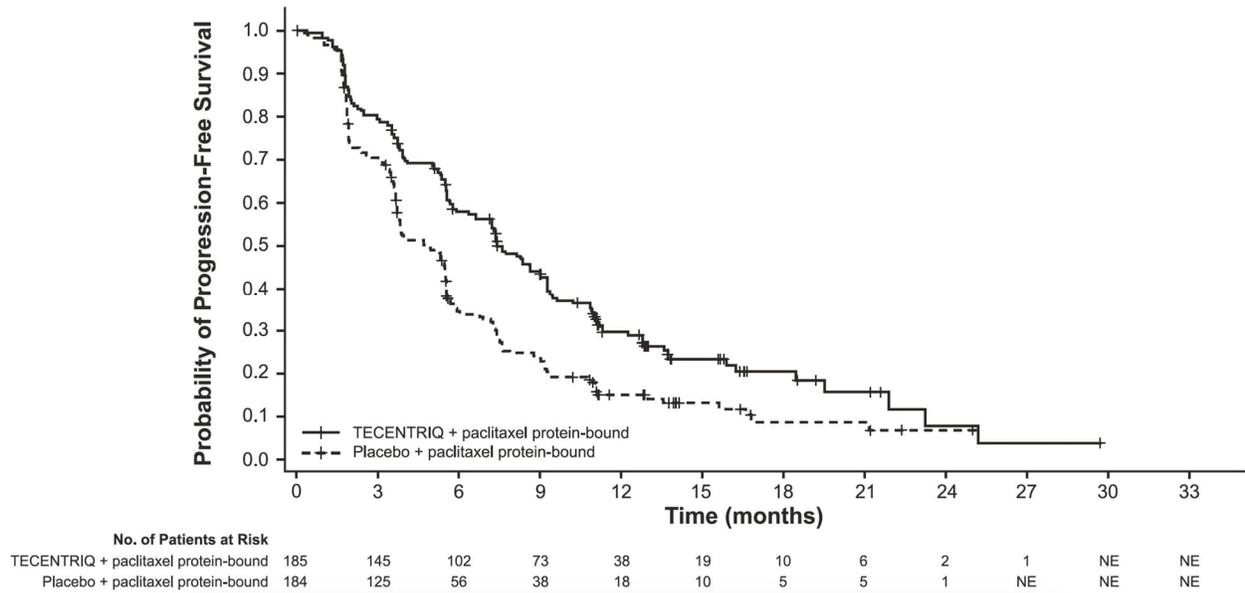
The efficacy results of IMpassion130 for the patient population with PD-L1 expression  $\geq$  1% are presented in Table 28 and Figure 5.

**Table 28: Efficacy Results from IMpassion130 in Patients with PD-L1 Expression  $\geq$  1%**

|   | PD-L1 Expression $\geq$ 1% <sup>1</sup>                |  |
|---|--|--|
|   | TECENTRIQ in combination with paclitaxel protein-bound | Placebo in combination with paclitaxel protein-bound |
| <b>Overall Survival</b> <sup>2</sup>              | n=185  | n=184  |
| Deaths (%)  | 120 (65%)  | 139 (76%)  |
| Median, months (95% CI)                           | 25.4 (19.6, 30.7)                                      | 17.9 (13.6, 20.3)                                    |
| Hazard ratio (95% CI)                             | 0.67 (0.53, 0.86)                                      |  |
| <b>Progression-Free Survival</b> <sup>3,4</sup>   | (n=185)  | (n=184)  |
| Events (%)  | 136 (74)   | 151 (82)   |
| Median, months (95% CI)                           | 7.4 (6.6, 9.2)   | 4.8 (3.8, 5.5)                                       |
| Stratified Hazard ratio (95% CI) <sup>5</sup>     | 0.60 (0.48, 0.77)                                      |  |
| p-value   | <0.0001  |  |
| <b>Objective Response Rate</b> <sup>3,4,6,7</sup> | n=185  | n=183  |
| Number of responders (%)                          | 98 (53)  | 60 (33)  |
| (95% CI)  | (45.5, 60.3)   | (26.0, 40.1)   |
| Complete Response (%)                             | 17 (9)   | 1 (<1)   |
| Partial Response (%)                              | 81 (44)  | 59 (32)  |
| <b>Duration of Response</b> <sup>3,4,7</sup>      | n=98   | n=60   |
| Median, months                                    | 9.2  | 6.2  |

|   |             |            |
|---|-------------|------------|
| (95% CI)  | (7.5, 11.9) | (5.5, 8.8) |
| <sup>1</sup> PD-L1 expression in tumor-infiltrating immune cells (IC)<br><sup>2</sup> OS in the PD-L1 expression $\geq$ 1% population was a major efficacy endpoint, but there remained no alpha to evaluate this endpoint. These results should be considered not statistically significant.<br><sup>3</sup> As determined by investigator assessment<br><sup>4</sup> per RECIST v1.1 (Response Evaluation Criteria in Solid Tumors v1.1)<br><sup>5</sup> Stratified by presence of liver metastases, and by prior taxane treatment<br><sup>6</sup> Patients with measurable disease at baseline<br><sup>7</sup> Confirmed responses<br>PFS=Progression-Free Survival; CI=Confidence Interval; ORR=Objective Response Rate; DOR=Duration of Response; NE=Not Estimable |             |            |

**Figure 5: Kaplan-Meier Plot of Progression-Free-Survival in IMpassion130 in Patients with PD-L1 Expression  $\geq$ 1%**



#### 14.4 Small Cell Lung Cancer

The efficacy of TECENTRIQ with carboplatin and etoposide was investigated in IMpower133 (NCT02763579), a randomized (1:1), multicenter, double-blind, placebo-controlled trial in 403 patients with ES-SCLC. IMpower133 enrolled patients with ES-SCLC who had received no prior chemotherapy for extensive stage disease and ECOG performance status 0 or 1. The trial excluded patients with active or untreated CNS metastases, history of autoimmune disease, administration of a live, attenuated vaccine within 4 weeks prior to randomization, or administration of systemic immunosuppressive medications within 1 week prior to randomization. Randomization was stratified by sex, ECOG performance status, and presence of brain metastases. Patients were randomized to receive one of the following two treatment arms:

- TECENTRIQ 1200 mg and carboplatin AUC 5 mg/mL/min on Day 1 and etoposide 100 mg/m<sup>2</sup> intravenously on Days 1, 2 and 3 of each 21-day cycle for a maximum of 4 cycles followed by TECENTRIQ 1200 mg once every 3 weeks until disease progression or unacceptable toxicity, or
- placebo and carboplatin AUC 5 mg/mL/min on Day 1 and etoposide 100 mg/m<sup>2</sup> intravenously on Days 1, 2, and 3 of each 21-day cycle for a maximum of 4 cycles followed by placebo once every 3 weeks until disease progression or unacceptable toxicity.

Administration of TECENTRIQ was permitted beyond RECIST-defined disease progression. Tumor assessments were conducted every 6 weeks for the first 48 weeks following Cycle 1, Day

1 and then every 9 weeks thereafter. Patients treated beyond disease progression had tumor assessment conducted every 6 weeks until treatment discontinuation.

Major efficacy outcome measures were OS and PFS as assessed by investigator per RECIST v1.1 in the intent-to-treat population. Additional efficacy outcome measures included ORR and DoR as assessed by investigator per RECIST v1.1.

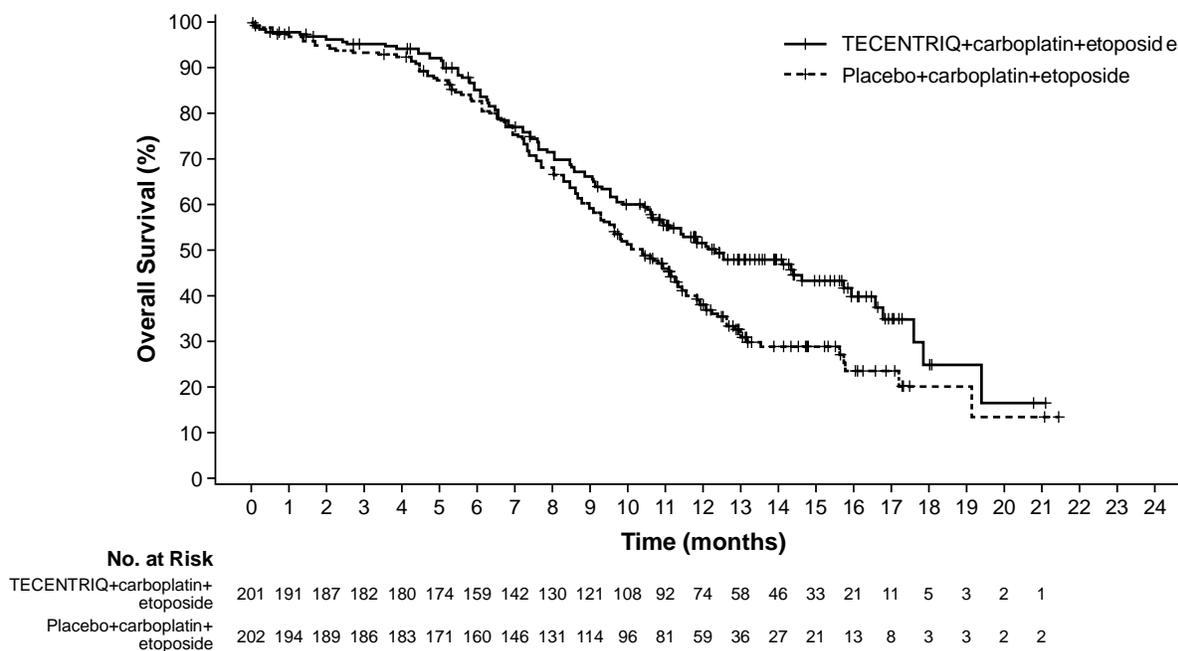
A total of 403 patients were randomized, including 201 to the TECENTRIQ arm and 202 to the chemotherapy alone arm. The median age was 64 years (range 26 to 90) and 65% were male. The majority of patients were White (80%); 17% were Asian, 4% were Hispanic and 1% were Black. Baseline ECOG performance status was 0 (35%) or 1 (65%); 9% of patients had a history of brain metastases, and 97% were current or previous smokers.

Efficacy results are presented in Table 29 and Figure 6.

**Table 29: Efficacy Results from IMpower133**

|   | <b>TECENTRIQ with Carboplatin and Etoposide</b> | <b>Placebo with Carboplatin and Etoposide</b> |
|---|---|---|
| <b>Overall Survival</b>   | N=201   | N=202   |
| Deaths (%)  | 104 (52%)                                       | 134 (66%)                                     |
| Median, months  | 12.3  | 10.3  |
| (95% CI)  | (10.8, 15.9)                                    | (9.3, 11.3)                                   |
| Hazard ratio <sup>3</sup> (95% CI)  | 0.70 (0.54, 0.91)                               |   |
| p-value <sup>4,5</sup>  | 0.0069  |   |
| <b>Progression-Free Survival<sup>1,2</sup></b>  | N=201   | N=202   |
| Number of events (%)  | 171 (85%)                                       | 189 (94%)                                     |
| Median, months  | 5.2   | 4.3   |
| (95% CI)  | (4.4, 5.6)                                      | (4.2, 4.5)                                    |
| Hazard ratio <sup>3</sup> (95% CI)  | 0.77 (0.62, 0.96)                               |   |
| p-value <sup>4,6</sup>  | 0.0170  |   |
| <b>Objective Response Rate<sup>1,2,7</sup></b>  | N=201   | N=202   |
| Number of responders (%)  | 121 (60%)                                       | 130 (64%)                                     |
| (95% CI)  | (53, 67)  | (57, 71)                                      |
| Complete Response (%)   | 5 (2%)  | 2 (1%)  |
| Partial Response (%)  | 116 (58%)                                       | 128 (63%)                                     |
| <b>Duration of Response<sup>1,2,7</sup></b>   | N=121   | N=130   |
| Median, months  | 4.2   | 3.9   |
| (95% CI)  | (4.1, 4.5)                                      | (3.1, 4.2)                                    |
| <sup>1</sup> As determined by investigator assessment<br><sup>2</sup> per RECIST v1.1 (Response Evaluation Criteria in Solid Tumors v1.1)<br><sup>3</sup> Stratified by sex and ECOG performance status<br><sup>4</sup> Based on the stratified log-rank test<br><sup>5</sup> Compared to the allocated $\alpha$ of 0.0193 for this interim analysis based on 78% information using O'Brien-Fleming boundary<br><sup>6</sup> Compared to the allocated $\alpha$ of 0.05 for this analysis.<br><sup>7</sup> Confirmed response<br>CI=confidence interval |   |   |

**Figure 6: Kaplan-Meier Plot of Overall Survival in IMpower133**



## 14.5 Hepatocellular Carcinoma

The efficacy of TECENTRIQ in combination with bevacizumab was investigated in IMbrave150 (NCT03434379), a multicenter, international, open-label, randomized trial in patients with locally advanced unresectable and/or metastatic hepatocellular carcinoma who have not received prior systemic therapy. Randomization was stratified by geographic region (Asia excluding Japan vs. rest of world), macrovascular invasion and/or extrahepatic spread (presence vs. absence), baseline AFP (<400 vs.  $\geq$ 400 ng/mL), and by ECOG performance status (0 vs. 1).

A total of 501 patients were randomized (2:1) to receive either TECENTRIQ as an intravenous infusion of 1200 mg, followed by 15 mg/kg bevacizumab, on the same day every 3 weeks or sorafenib 400 mg given orally twice daily, until disease progression or unacceptable toxicity. Patients could discontinue either TECENTRIQ or bevacizumab (e.g., due to adverse events) and continue on single-agent therapy until disease progression or unacceptable toxicity associated with the single-agent.

The study enrolled patients who were ECOG performance score 0 or 1 and who had not received prior systemic treatment. Patients were required to be evaluated for the presence of varices within 6 months prior to treatment, and were excluded if they had variceal bleeding within 6 months prior to treatment, untreated or incompletely treated varices with bleeding, or high risk of bleeding. Patients with Child-Pugh B or C cirrhosis, moderate or severe ascites; history of hepatic encephalopathy; a history of autoimmune disease; administration of a live, attenuated vaccine within 4 weeks prior to randomization; administration of systemic immunostimulatory agents within 4 weeks or systemic immunosuppressive medications within 2 weeks prior to randomization; or untreated or corticosteroid-dependent brain metastases were excluded. Tumor assessments were performed every 6 weeks for the first 54 weeks and every 9 weeks thereafter.

The demographics and baseline disease characteristics of the study population were balanced between the treatment arms. The median age was 65 years (range: 26 to 88) and 83% of patients were male. The majority of patients were Asian (57%) or White (35%); 40% were from Asia (excluding Japan). Approximately 75% of patients presented with macrovascular invasion and/or extrahepatic spread and 37% had a baseline AFP  $\geq$ 400 ng/mL. Baseline ECOG performance status was 0 (62%) or 1 (38%). HCC risk factors were Hepatitis B in 48% of patients, Hepatitis C in 22%, and 31% of patients had non-viral liver disease. The majority of patients had BCLC stage C (82%) disease at baseline, while 16% had stage B, and 3% had stage A.

The major efficacy outcome measures were overall survival (OS) and independent review facility (IRF)-assessed progression free survival (PFS) per RECIST v1.1. Additional efficacy outcome measures were IRF-assessed overall response rate (ORR) per RECIST and mRECIST.

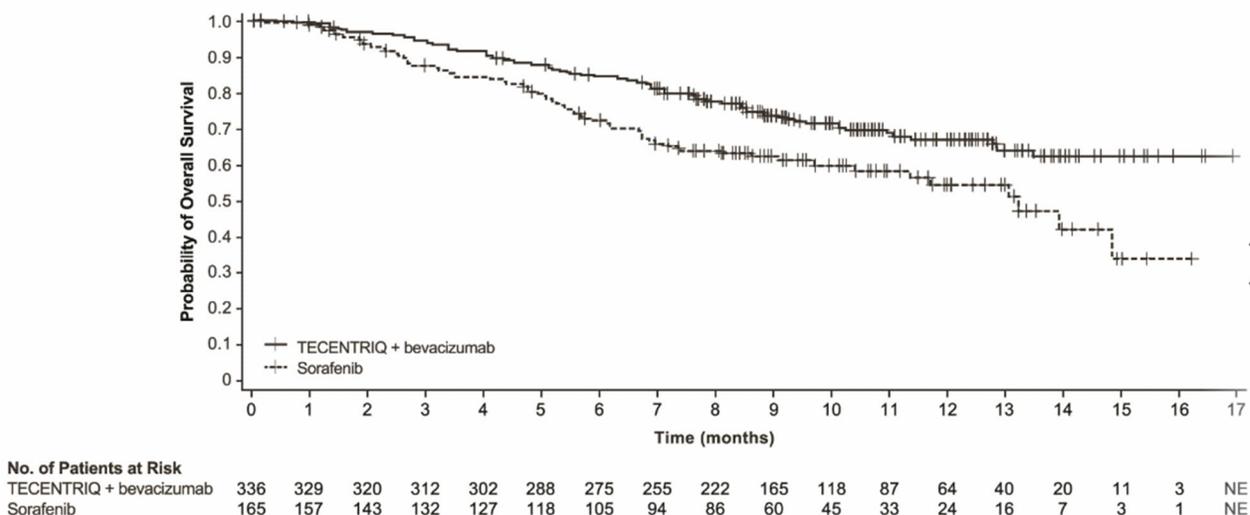
Efficacy results are presented in Table 30 and Figure 7.

**Table 30: Efficacy Results from IMbrave150**

|   | <b>TECENTRIQ in combination with<br/>Bevacizumab<br/>(N= 336)</b> | <b>Sorafenib<br/>(N=165)</b> |
|---|---|------------------------------|
| <b>Overall Survival</b>   |   |                              |
| Number of deaths (%)  | 96 (29)   | 65 (39)                      |
| Median OS in months<br>(95% CI)   | NE<br>(NE, NE)  | 13.2<br>(10.4, NE)           |
| Hazard ratio <sup>1</sup> (95% CI)  | 0.58 (0.42, 0.79)   |                              |
| p-value <sup>2</sup>  | 0.0006 <sup>2</sup>   |                              |
| <b>Progression-Free Survival<sup>3</sup></b>  |   |                              |
| Number of events(%)   | 197 (59)  | 109 (66)                     |
| Median PFS in months (95% CI)   | 6.8 (5.8, 8.3)  | 4.3 (4.0, 5.6)               |
| Hazard ratio <sup>1</sup> (95% CI)  | 0.59 (0.47, 0.76)   |                              |
| p-value   | <0.0001   |                              |
| <b>Overall Response Rate<sup>3,5</sup>(ORR), RECIST 1.1</b>   |   |                              |
| Number of responders (%)  | 93 (28)   | 19 (12)                      |
| (95% CI)  | (23, 33)  | (7,17)                       |
| p-value <sup>4</sup>  | <0.0001   |                              |
| Complete responses, n (%)   | 22 (7)  | 0                            |
| Partial responses, n (%)  | 71 (21)   | 19 (12)                      |
| <b>Duration of Response<sup>3,5</sup> (DOR) RECIST 1.1</b>  |   |                              |
|   | (n=93)  | (n=19)                       |
| Median DOR in months<br>(95% CI)  | NE<br>(NE, NE)  | 6.3<br>(4.7, NE)             |
| Range (months)  | (1.3+, 13.4+)   | (1.4+, 9.1+)                 |
| <b>Overall Response Rate<sup>3,5</sup> (ORR), HCC mRECIST</b>   |   |                              |
| Number of responders (%)  | 112 (33)  | 21 (13)                      |
| (95% CI)  | (28, 39)  | (8, 19)                      |
| p-value <sup>4</sup>  | <0.0001   |                              |
| Complete responses, n (%)   | 37 (11)   | 3 (1.8)                      |
| Partial responses, n (%)  | 75 (22)   | 18 (11)                      |
| <b>Duration of Response<sup>3,5</sup> (DOR) HCC mRECIST</b>   |   |                              |
|   | (n=112)   | (n=21)                       |
| Median DOR in months<br>(95% CI)  | NE<br>(NE, NE)  | 6.3<br>(4.9, NE)             |
| Range (months)  | (1.3+, 13.4+)   | (1.4+, 9.1+)                 |
| <sup>1</sup> Stratified by geographic region (Asia excluding Japan vs. rest of world), macrovascular invasion and/or extrahepatic spread (presence vs. absence), and baseline AFP (<400 vs. $\geq$ 400 ng/mL) |   |                              |
| <sup>2</sup> Based on two-sided stratified log-rank test; as compared to significance level 0.004 (2-sided) based on 161/312=52% information using the OBF method   |   |                              |

<sup>3</sup> Per independent radiology review  
<sup>4</sup> Based on two-sided Cochran-Mantel-Haenszel test  
<sup>5</sup> Confirmed responses  
+ Denotes a censored value  
CI=confidence interval; HCC mRECIST= Modified RECIST Assessment for Hepatocellular Carcinoma; NE=not estimable; RECIST 1.1= Response Evaluation Criteria in Solid Tumors v1.1

**Figure 7: Kaplan-Meier Plot of Overall Survival in IMbrave150**



Exploratory analyses showed that the subset of patients (20%) who were ADA-positive by week 6 appeared to have reduced efficacy (effect on OS) as compared to patients (80%) who tested negative for treatment-emergent ADA by week 6 [see Adverse Reactions (6.2), Clinical Pharmacology (12.3)]. ADA-positive patients by week 6 appeared to have similar overall survival compared to sorafenib-treated patients. In an exploratory analysis, inverse probability weighting was conducted to compare ADA-positive patients and ADA-negative patients in the TECENTRIQ and bevacizumab arm to the sorafenib arm. Inverse probability weighting factors were: baseline sum of longest tumor size (BSLD), baseline ECOG, baseline albumin, baseline LDH, sex, age, race, geographic region, weight, neutrophil-to-lymphocyte ratio, AFP (<400 ng/mL vs  $\geq$ 400 ng/mL), number of metastatic sites, MVI and/or EHS present at study entry, etiology (HBV vs. HCV vs. non-viral) and Child-Pugh Score (A5 VS. A6). The OS hazard ratio comparing the ADA-positive subgroup of the TECENTRIQ and bevacizumab arm to sorafenib was 0.93 (95% CI: 0.57, 1.53). The OS hazard ratio comparing the ADA-negative subgroup to sorafenib was 0.39 (95% CI: 0.26, 0.60).

## 14.6 Melanoma

The efficacy of TECENTRIQ in combination with cobimetinib and vemurafenib was evaluated in a double-blind, randomized (1:1), placebo-controlled, multicenter trial (IMspire150; NCT02908672) conducted in 514 patients. Randomization was stratified by geographic location (North America vs. Europe vs. Australia, New Zealand, and others) and baseline lactate dehydrogenase (LDH) [less than or equal to upper limit of normal (ULN) vs. greater than ULN]. Eligible patients were required to have previously untreated unresectable or metastatic BRAF V600 mutation-positive melanoma as detected by a locally available test and centrally confirmed with the FoundationOne™ assay. Patients were excluded if they had history of autoimmune disease; administration of a live, attenuated vaccine within 28 days prior to randomization; administration of systemic immunostimulatory agents within 4 weeks or systemic

immunosuppressive medications within 2 weeks prior to randomization; and active or untreated CNS metastases.

TECENTRIQ was initiated after patients received a 28-day treatment cycle of cobimetinib 60 mg orally once daily (21 days on / 7 days off) and vemurafenib 960 mg orally twice daily Days 1-21 and 720 mg orally twice daily Days 22-28. Patients received TECENTRIQ 840 mg intravenous infusion over 60 minutes every 2 weeks in combination with cobimetinib 60 mg orally once daily and vemurafenib 720 mg orally twice daily, or placebo in combination with cobimetinib 60 mg orally once daily and vemurafenib 960 mg orally twice daily. Treatment continued until disease progression or unacceptable toxicity. There was no crossover at the time of disease progression. Tumor assessments were performed every 8 weeks ( $\pm$  1 week) for the first 24 months and every 12 weeks ( $\pm$  1 week) thereafter.

The major efficacy outcome measure was investigator-assessed progression-free survival (PFS) per RECIST v1.1. Additional efficacy outcomes included PFS assessed by an independent central review, investigator-assessed ORR, OS, and DOR.

The median age of the study population was 54 years (range: 22-88), 58% of patients were male, 95% were White, a baseline ECOG performance status of 0 (77%) or 1 (23%), 33% had elevated LDH, 94% had metastatic disease, 60% were Stage IV (M1C), 56% had less than three metastatic sites at baseline, 3% had prior treatment for brain metastases, 30% had liver metastases at baseline, and 14% had received prior adjuvant systemic therapy. Based on central testing, 74% were identified as having a V600E mutation, 11% as having V600K mutation, and 1% as having V600D or V600R mutations.

Efficacy results are summarized in Table 31 and Figure 8. Patients had a median survival follow up time of 18.9 months.

**Table 31 Efficacy Results from IMspire150**

|  | <b>Tecentriq + Cobimetinib +<br/>Vemurafenib</b><br>N=256 | <b>Placebo + Cobimetinib +<br/>Vemurafenib</b><br>N=258 |
|--|---|---|
| <b>Progression-Free Survival<sup>1</sup></b> |   |   |
| Number of events (%)                         | 148 (58)  | 179 (69)  |
| Median, months                               | 15.1  | 10.6  |
| (95% CI)                                     | (11.4, 18.4)  | (9.3, 12.7)   |
| Hazard ratio <sup>2</sup> (95% CI)           | 0.78 (0.63, 0.97)   |   |
| p-value <sup>3</sup>                         | 0.0249  |   |
| <b>Overall Response Rate<sup>1,4</sup></b>   |   |   |
| Number of responders (%)                     | 170 (66)  | 168 (65)  |
| (95% CI)                                     | (60, 72)  | (59, 71)  |
| Complete responses, n (%)                    | 41 (16)   | 46 (18)   |
| Partial response, n (%)                      | 129 (50)  | 122 (47)  |
| <b>Duration of Response<sup>1,4</sup></b>    |   |   |
|  | n=170   | n=168   |
| Median, months                               | 20.4  | 12.5  |
| (95% CI)                                     | (15.1, NE)  | (10.7, 16.6)  |

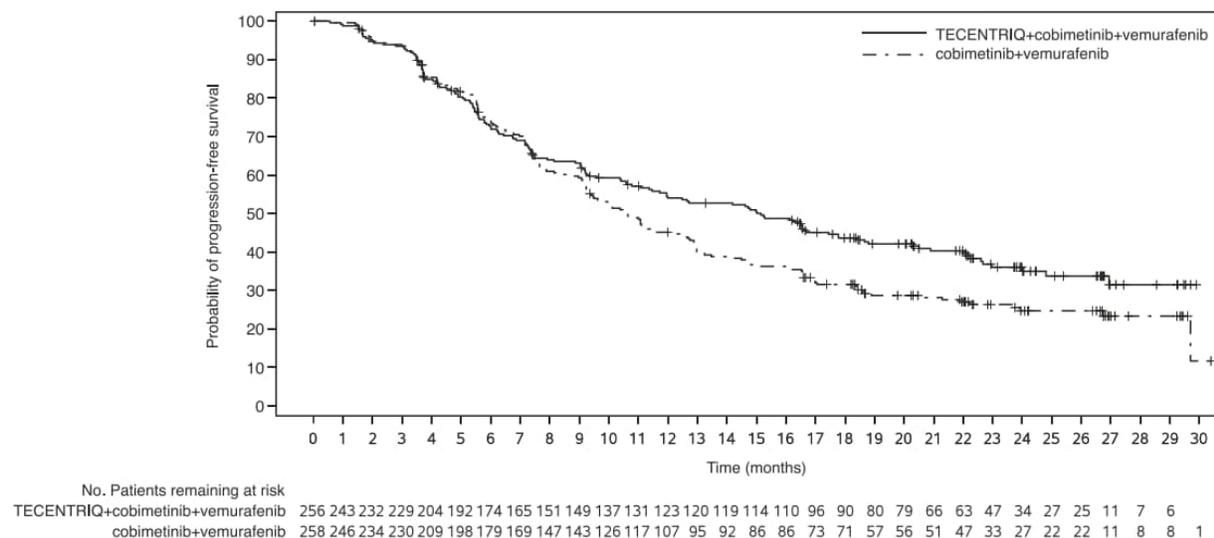
<sup>1</sup> As determined by investigator assessment with Response Evaluation Criteria in Solid Tumors v1.1.; CI=confidence interval;

<sup>2</sup> Stratified by baseline LDH

<sup>3</sup>Based on the stratified log-rank test

<sup>4</sup>Confirmed Responses

**Figure 8: Kaplan-Meier Plot for Progression-Free Survival in IMspire150**



At a pre-specified analysis at the time of the primary analysis of PFS, the OS data were not mature. The median OS was 28.8 months with 93 (36%) deaths in the TECENTRIQ plus cobimetinib and vemurafenib arm, and 25.1 months with 112 (43%) deaths in the placebo plus cobimetinib and vemurafenib arm. The hazard ratio for OS was 0.85 (95% CI: 0.64, 1.11) and the p-value was 0.2310.

## 16 HOW SUPPLIED/STORAGE AND HANDLING

TECENTRIQ injection is a sterile, preservative-free, and colorless to slightly yellow solution for intravenous infusion supplied as a carton containing one 840 mg/14 mL single-dose vial or 1,200 mg/20 mL single-dose vial

Store vials in a refrigerator at 2°C to 8°C in the outer carton in order to protect from light. Do not freeze. Do not shake.

Shelf-life

The expiry date of the product is indicated on the packaging materials.

**17    MARKETING AUTHORISATION HOLDER**

Roche Pharmaceuticals (Israel) Ltd. P.O.B. 6391, Hod Hasharon, 4524079.

**18    MARKETING AUTHORISATION NUMBER(S):**

158.66.34982

**19    MANUFACTURER**

Hoffmann-La Roche ltd, Basel, Switzerland

Revised on June 2022 according to MOHs guidelines