



# TECVAYLI™ (teclistamab)▼ Patient Card

Important Safety Information for Patient Receiving TECVAYLI®(teclistamab)

Carry this card with you at all times. **SHOW THIS CARD** to any healthcare professional involved in your care and if you go to the hospital

This card has been reviewed and approved by The Saudi Food and Drug Authority (SFDA).

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**TECVAYLI can cause side effects such as cytokine release syndrome (CRS).** Cytokine release syndrome is a serious immune reaction that can be triggered by a variety of factors, including a range of drugs.<sup>1</sup>

PATIENT'S NAME:

FOLD HERE This card has been reviewed and approved by The Saudi Food and Drug Authority (SFDA).

# Important Safety Information for Patients

Get medical help straight away if you experience any of the following:<sup>2</sup>

- Fever (38°C or higher)
- Chills
- Fast heartbeat

- Nausea
- Headache
- Feeling dizzy
- Difficulty breathing

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**IMPORTANT TO REMEMBER:** Stay close to the location where you received your TECVAYLI therapy for at least 2 days for daily monitoring after administration of your first three doses (usually two step-up doses and first maintenance dose).<sup>2</sup> If you have <u>any</u> of the symptoms listed on this card call your doctor or seek emergency medical attention right away! These are not all the possible side effects of TECVAYLI. Tell your doctor if you have any side effect that bothers you or does not go away.

### **Treating Physician**

#### TREATING PHYSICIAN'S NAME:

TREATING PHYSICIAN'S PHONE NUMBER:

# HOSPITAL NAME AND ADDRESS:

#### PHONE NUMBER:

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### Information for Healthcare Team to Fill In

Please give this card to your healthcare team to fill in the information and return to you.

Dates of TECVAYLI injections (step-up dosing schedule):

#### **STEP-UP DOSE 1**

#### **STEP-UP DOSE 2**

### FIRST MAINTENANCE DOSE\*

\*This is the first full treatment dose (1.5 mg/kg)<sup>2</sup>

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#### Important Safety Information for Healthcare Professionals

CRS, including life-threatening or fatal reactions, may occur in patients receiving TECVAYLI.<sup>2</sup> The majority of CRS events observed following TECVAYLI administration were Grade 1 and 2.<sup>2</sup> CRS may involve multiple organ systems. Assess the patient for signs and symptoms of CRS. If your patient reports any signs or symptoms as referenced on this card, please contact the patient's treating physician immediately for further information.

See Summary of Product Characteristics for full details.<sup>2</sup>

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 Adverse event: reporting guidance: Beldnah hemecoviplance Center (NFC) Cail Center: 1999 Wabile: https://doc.idla.gov.al Fortup Perschlag laformation, please refer to the datasheet or contect Jahnson & Johnson Middle Barl F2-Lit (Byodh) Office lafe 00%4 - 11.439/13 Mindle 200%4 - 11.439/13 

Hofline: 00966540015811

This medicinal product is subject to additional monitoring. This will allow quick identification of new safety information.

1. Shimabukuro-Vornhagen A et al. J Immunother Cancer 2018;6(1):56.

2. TECVAYLI™ SmPC (EU), March 2023.

Please refer to the full patient leaflet for more information on Tecvayli

