HIGHLIGHTS OF PRESCRIBING INFORMATION
These highlights do not include all the information needed to use BREVANZI safely and effectively. See full prescribing information for BREVANZI.

BREVANZI® (lisocabtagene maraleucel) suspension for intravenous infusion
Initial U.S. Approval: 2021

WARNING: CYTOKINE RELEASE SYNDROME and NEUROLOGIC TOXICITIES
See full prescribing information for complete boxed warning.

- Cytokine Release Syndrome (CRS), including fatal or life-threatening reactions, occurred in patients receiving BREVANZI. Do not administer BREVANZI to patients with active infection or inflammatory disorders. Treat severe or life-threatening CRS with tocilizumab with or without corticosteroids (2.2, 2.3, 5.1).
- Neurologic toxicities, including fatal or life-threatening reactions, occurred in patients receiving BREVANZI, including concurrently with CRS, as CRS resolution, or in the absence of CRS. Monitor for neurologic events after treatment with BREVANZI. Provide supportive care and/or corticosteroids as needed (2.2, 2.3, 5.2).
- BREVANZI is available only through a restricted program under a Risk Evaluation and Mitigation Strategy (REMS) called the BREVANZI REMS (5.3).

Recent Major Changes
- Dosing of BREVANZI is based on the number of chimeric antigen receptor (CAR)-positive viable T cells (2.1).

Dosage and Administration
- For LBCI after one line of therapy, the dose is 90 to 110 × 10^6 CAR-positive viable T cells (2.1).
- For LBCI after two or more lines of therapy, the dose is 50 to 110 × 10^6 CAR-positive viable T cells (2.1).
- Administer BREVANZI in a REMS-certified healthcare facility (2.2, 5.1, 5.2, 5.3).

Indications and Usage
- BREVANZI is a CD19-directed genetically modified autologous T cell immunotherapy indicated for the treatment of adult patients with large B-cell lymphoma (LBCL), including diffuse large B-cell lymphoma (DLBCL) not otherwise specified (including DLBCL arising from indolent lymphoma), high-grade B-cell lymphoma, primary mediastinal large B-cell lymphoma, and follicular lymphoma grade 3B, who have:
  - refractory disease to first-line chemoimmunotherapy or relapse within 12 months of first-line chemoimmunotherapy (1); or
  - refractory disease to first-line chemoimmunotherapy or relapse after first-line chemoimmunotherapy and are not eligible for hematopoietic stem cell transplantation (HSCT) due to comorbidities or age (1); or
  - relapsed or refractory disease after two or more lines of systemic therapy (1).

Limitations of Use: BREVANZI is not indicated for the treatment of patients with primary central nervous system lymphoma (1, 14).

Dose and Administration
- Indications and Usage (1), LBCI after one line of therapy 6/2022
- Dosage and Administration (2.1) 6/2022
- Warnings and Precautions (5.1, 5.2, 5.5, 5.6, 5.9) 6/2022

Indications and Usage

- Do NOT use a leukodepleting filter (2.2).
- Administer a lymphodepleting regimen of fludarabine and cyclophosphamide before infusion of BREVANZI (lisocabtagene maraleucel) (2.2).
- Premedicate with acetaminophen and an H1 antihistamine (2.2).
- Confirm availability of tocilizumab prior to infusion (2.2, 5.1).
- Dosing of BREVANZI is based on the number of chimeric antigen receptor (CAR)-positive viable T cells (2.1).
- For LBCL after one line of therapy, the dose is 90 to 110 × 10^6 CAR-positive viable T cells (2.1).
- For LBCL after two or more lines of therapy, the dose is 50 to 110 × 10^6 CAR-positive viable T cells (2.1).
- Administer BREVANZI in a REMS-certified healthcare facility (2.2, 5.1, 5.2, 5.3).

Contraindications

- None (4).

Warnings and Precautions

- Hypersensitivity Reactions: Monitor for hypersensitivity reactions during infusion (5.4).
- Serious Infections: Monitor patients for signs and symptoms of infection; treat appropriately (5.9).
- Prolonged Cytopenias: Patients may exhibit Grade 3 or higher cytopenias for several weeks following BREVANZI infusion. Monitor complete blood counts (5.6).
- Hypogammaglobulinemia: Monitor and consider immunoglobulin replacement therapy (5.7).
- Secondary Malignancies: In the event that a secondary malignancy occurs after treatment with BREVANZI, contact Bristol-Myers Squibb at 1-888-805-4555 (5.8).
- Effects on Ability to Drive and Use Machines: Advise patients to refrain from driving and engaging in hazardous occupations or activities, such as operating heavy or potentially dangerous machinery for at least 8 weeks after BREVANZI administration (5.9).

Adverse Reactions

- The most common nonlaboratory adverse reactions (incidence ≤30%) are fever, cytokine release syndrome, fatigue, musculoskeletal pain, and nausea. The most common Grade 3 laboratory abnormalities (≤30%) include lymphocyte count decrease, neutrophil count decrease, platelet count decrease, and hemoglobin decrease (6.1).

To report SUSPECTED ADVERSE REACTIONS, contact Bristol-Myers Squibb at 1-800-721-5072 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

See 17 for PATIENT COUNSELING INFORMATION and Medication Guide

Revised: 6/2023
**Receipt of BREYANZI**

- BREYANZI is shipped directly to the cell-associated lab or clinical pharmacy associated with the infusion center in the vapor phase of a liquid nitrogen shipper.
- Confirm the patient's identity with the patient identifiers on the shipper.
- If the patient is not expected to be ready for administration before the shipper expires and the infusion site is qualified for onsite storage, transfer BREYANZI to onsite vapor phase of liquid nitrogen storage prior to preparation.
- If the patient is not expected to be ready for administration before the shipper expires and the infusion site is not qualified for onsite storage, contact Bristol-Myers Squibb at 1-888-805-4555 to arrange for return shipment.

**Preparing BREYANZI**

**Before thawing the vials**

- Confirm the patient's identity with the patient identifiers on the RFI Certificate.
- Read the RFI Certificate (affixed inside the shipper) for information on the number of syringes you will need to administer the CD8 and CD4 components (syringe labels are provided with the RFI Certificate). There is a separate RFI Certificate for each cell component.
- Confirm tocilizumab and emergency equipment are available prior to infusion and during the recovery period.
- Confirm the infusion time in advance and adjust the start time of BREYANZI thaw such that it will be available for infusion when the patient is ready.

**Thawing the vials**

1. Confirm the patient’s identity with the patient identifiers on the outer carton and on the syringes labels. Once the vials of CAR-positive viable T cells (CD8 component and CD4 component) are removed from frozen storage, the thaw must be carried to completion and the cells administered within 2 hours.
2. Remove the CD8 component carton and CD4 component carton from the outer carton.
3. Confirm the patient’s identity with the patient identifiers on the inner carton.
4. Open each inner carton and visually inspect the vial(s) for damage. If the vials are intact, proceed to prepare the syringes.
5. Confirm the patient’s identity with the patient identifiers on the vials.
6. Carefully remove the vials from the cartons, place vials on a protective barrier pad, and thaw at room temperature until there is no visible ice in the vials. Thaw all of the vials at the same time. Keep the CD8 and CD4 components separate.

**Dose preparation**

- Prepare BREYANZI using sterile technique.
- Based on the concentration of CAR-positive viable T cells for each component, more than one vial of each of the CD8 and CD4 components may be required to complete a dose. A separate syringe should be prepared for each CD8 or CD4 component vial received.

**Note:** The volume to be drawn up and infused may differ for each component as indicated on the RFI Certificate. Do NOT draw up excess volume into the syringe.

- Each vial contains 5 mL with a total extractable volume of 4.6 mL of CD8 or CD4 component T cells. The RFI certificate for each component indicates the volume (mL) of cells to be drawn up into each syringe. Use the smallest Luer-lock tip syringe necessary (1, 3, or 5 mL) to draw up the specified volume from each vial. A 5 mL syringe should not be used for volumes less than 3 mL.

**7. Prepare the syringe(s) of the CD8 component first.** Affix the CD8 syringe labels to the syringe(s) prior to pulling the required volume into the syringe(s).

**Note:** It is important to confirm that the volume drawn up for each component matches the volume specified in the respective RFI Certificate. Do NOT draw up excess volume into the syringe.

**Withdrawal of the required volume of cells from each vial into a separate syringe should be carried out using the following instructions:**

- Hold the thawed vial(s) upright and gently invert the vial(s) 5 times to mix the cell product. If any clumping is apparent, continue to invert the vial(s) until clumps have dispersed and cells appear to be evenly suspended.

---

**1 INDICATIONS AND USAGE**

BREYANZI is a CD19-directed genetically modified autologous T cell immunotherapy indicated for the treatment of adult patients with large B-cell lymphoma (LBCL), including diffuse large B-cell lymphoma (DLBCL) not otherwise specified (including DLBCL arising from indolent lymphoma), high-grade B-cell lymphoma, primary mediastinal large B-cell lymphoma, and follicular lymphoma grade 3B who have:

- refractory disease to first-line chemoimmunotherapy or relapse within 12 months of first-line chemoimmunotherapy; or
- refractory disease to first-line chemoimmunotherapy or relapse after first-line chemoimmunotherapy and are not eligible for hematopoietic stem cell transplantation (HSCT) due to comorbidities or age; or
- relapsed or refractory disease after two or more lines of systemic therapy.

**Limitations of Use:** BREYANZI is not indicated for the treatment of patients with primary central nervous system (CNS) lymphoma [see Clinical Studies (14)].

**2 DOSAGE AND ADMINISTRATION**

**For autologous use only. For intravenous use only.**

**2.1 Dose**

See the respective Certificate of Release for Infusion (RFI Certificate) for each component, for the actual cell counts and volumes to be infused [see Dosage and Administration (2.2) and Dosage Forms and Strengths (3)].

**Relapsed or Refractory LBCL After One Line of Therapy**

A single dose of BREYANZI contains 90 to 110 × 10⁶ CAR-positive viable T cells (consisting of 1:1 CAR-positive viable T cells of the CD8 and CD4 components), with each component supplied separately in one to four single-dose vials.

**Relapsed or Refractory LBCL After Two or More Lines of Therapy**

A single dose of BREYANZI contains 50 to 110 × 10⁶ CAR-positive viable T cells (consisting of 1:1 CAR-positive viable T cells of the CD8 and CD4 components), with each component supplied separately in one to four single-dose vials.

**2.2 Administration**

BREYANZI is for autologous use only. The patient’s identity must match the patient identifiers on the BREYANZI cartons, vials and syringe labels. Do not confuse BREYANZI if the information on the patient-specific labels does not match the intended patient.

**Preparing the Patient for BREYANZI**

Confirm the availability of BREYANZI before starting lymphodepleting chemotherapy.

**Preparation**

Administer the lymphodepleting chemotherapy regimen before infusion of BREYANZI: fludarabine 30 mg/m²/day intravenously (IV), and cyclophosphamide 300 mg/m²/day IV for 3 days. See the prescribing information for fludarabine and cyclophosphamide for information on dose adjustment in renal impairment.

Infuse BREYANZI 2 to 7 days after completion of lymphodepleting chemotherapy.

Delay the infusion of BREYANZI if the patient has unresolved serious adverse events from preceding chemotherapies, active uncontrolled infection, or active graft-versus-host disease (GVHD).

**Premedication**

To minimize the risk of infusion reactions, premedicate the patient with acetaminophen (650 mg orally) and diphenhydramine (25-50 mg, IV or orally), or another H1-antihistamine, 30 to 60 minutes prior to treatment with BREYANZI.

Avoid prophylactic use of systemic corticosteroids, as they may interfere with the activity of BREYANZI.
13. WITHOUT drawing air into the syringe, slowly withdraw the target volume (as specified in the RFI Certificate). Carefully inspect the syringe for signs of debris prior to proceeding. If there is debris, contact Bristol-Myers Squibb at 1-888-805-4555. The liquid in the vials should be slightly opaque to opaque, colorless to yellow or brownish-yellow.

10. Remove the polyaluminum cover (if present) from the bottom of the vial and swab the septum with an alcohol wipe. Allow to air dry before proceeding. 

**NOTE:** The absence of the polyaluminum cover does not impact the sterility of the vial.

11. Keeping the vial(s) upright, cut the seal on the tubing line on the top of the vial immediately above the filter to open the air vent on the vial.

**NOTE:** Be careful to select the correct tubing line with the filter. Cut ONLY the tubing immediately above the filter to open the air vent on the vial.

12. Hold a 20-gauge, 1-1 ½ inch needle, with the opening of the needle tip away from the retrieval port septum. 

a. Insert the needle into the septum at a 45°-60° angle to puncture the retrieval port septum.

b. Increase the angle of the needle gradually as the needle enters the vial.

14. Verify that the volume of CD8/CD4 component matches the volume specified for the relevant component in the RFI Certificate. Once the volume is verified, remove the syringe/needle from the vial, carefully detach the needle from the syringe and cap the syringe.

15. Continue to keep the vial horizontal and return it to the carton to avoid leaking from the vial.

16. Dispose of any unused portion of BREYANZI (according to local biosecurity guidelines).

17. Repeat the process steps 7-16 for the CD4 Component.

18. Transport the labeled CD8 and CD4 syringes to the bedside by placing with protective barrier pad inside an insulated room temperature container.

**BREYANZI Administration**

- **Do NOT** use a leukodepleting filter.
- Ensure tocilizumab and emergency equipment are available prior to infusion and during the recovery period.
- Confirm the patient’s identity matches the patient identifiers on the syringe label.
- Once BREYANZI has been drawn into syringes, proceed with administration as soon as possible. The total time from removal from frozen storage to patient administration should not exceed 2 hours as indicated by the time entered on the syringe label.

1. Use intravenous normal saline to flush all the infusion tubing prior to and after each CD8 or CD4 component administration.

2. Administer the entire volume of the CD8 component intravenously at an infusion rate of approximately 0.5 mL/minute, using the closest port or Y-arm.

**NOTE:** The time for infusion will vary but will usually be less than 15 minutes for each component.

3. If more than one syringe is required for a full cell dose of the CD8 component, administer the volume in each syringe consecutively without any time between administering the contents of the syringes (unless there is a clinical reason (e.g., infusion reaction) to hold the dose).

4. After the CD8 component has been administered, flush the tubing with normal saline, using enough volume to clear the tubing and the length of the IV catheter.

5. Administer the CD4 component second, immediately after administration of the CD8 component is complete, using steps 1-4, as described for the CD8 component. Following administration of the CD4 component, flush the tubing with normal saline, using enough volume to clear the tubing and the length of the IV catheter.

**BREYANZI** contains human blood cells that are genetically modified with replication-incompetent, self-inactivating lentiviral vector. Follow universal precautions and local biosecurity guidelines applicable for the handling and disposal, to avoid potential transmission of infectious diseases.

**Monitoring**

- Administer BREYANZI at a REMS-certified healthcare facility.
- Monitor patients daily for at least 7 days following BREYANZI infusion at a REMS-certified healthcare facility for signs and symptoms of CRS and neurologic toxicities.
- Instruct patients to remain within proximity of the certified healthcare facility for at least 4 weeks following infusion.
- Instruct patients to refrain from driving or hazardous activities for at least 8 weeks following infusion.

2.3 **Management of Severe Adverse Reactions**

**Cytokine Release Syndrome**

Identify cytokine release syndrome (CRS) based on clinical presentation (see Warnings and Precautions [5.1]). Evaluate for and treat other causes of fever, hypoxia, and hypotension. If CRS is suspected, manage according to the recommendations in Table 1. Patients who experience Grade 2 or higher CRS (e.g., hypotension not responsive to fluids, or hypoxia requiring supplemental oxygenation) should be monitored with continuous cardiac telemetry and pulse oximetry. For patients experiencing severe CRS, consider performing an echocardiogram to assess cardiac function. For severe or life-threatening CRS, consider intensive-care supportive therapy.

If concurrent neurologic toxicity is suspected during CRS, administer:

- Corticosteroids according to the more aggressive intervention based on the CRS and neurologic toxicity grades in Tables 1 and 2.
- Tocilizumab according to the CRS grade in Table 1.
- Antiseizure medication according to the neurologic toxicity in Table 2.
Table 1: CRS Grading and Management Guidance

<table>
<thead>
<tr>
<th>CRS Grade</th>
<th>Tocilizumab</th>
<th>Corticosteroids</th>
</tr>
</thead>
<tbody>
<tr>
<td>Grade 1</td>
<td>If less than 72 hours after infusion, consider tocilizumab 8 mg/kg IV over 1 hour (not to exceed 800 mg). If 72 hours or more after infusion, treat symptomatically.</td>
<td>If less than 72 hours after infusion, consider dexamethasone 10 mg IV every 24 hours. If 72 hours or more after infusion, treat symptomatically.</td>
</tr>
<tr>
<td>Grade 2</td>
<td>Administer tocilizumab 8 mg/kg IV over 1 hour (not to exceed 800 mg). Repeat tocilizumab every 8 hours as needed if not responsive to intravenous fluids or increasing supplemental oxygen. Limit to a maximum of 3 doses in a 24-hour period; maximum total of 4 doses.</td>
<td>If less than 72 hours after infusion, administer dexamethasone 10 mg IV every 12-24 hours. If 72 hours or more after infusion, consider dexamethasone 10 mg IV every 12-24 hours.</td>
</tr>
<tr>
<td>Grade 3</td>
<td>Per Grade 2.</td>
<td>Administer dexamethasone 10 mg IV every 12 hours.</td>
</tr>
<tr>
<td>Grade 4</td>
<td>Per Grade 2.</td>
<td>Administer dexamethasone 20 mg IV every 6 hours.</td>
</tr>
</tbody>
</table>

Table 2: Neurologic Toxicity (NT) Grading and Management Guidance

<table>
<thead>
<tr>
<th>NT Grade</th>
<th>Corticosteroids and Antiseizure Medication</th>
</tr>
</thead>
<tbody>
<tr>
<td>Grade 1</td>
<td>Start non-sedating, antiseizure medicines (e.g., levetiracetam) for seizure prophylaxis. If 72 hours or more after infusion, observe. If less than 72 hours after infusion, consider dexamethasone 10 mg IV every 12 to 24 hours for 2 to 3 days.</td>
</tr>
<tr>
<td>Grade 2</td>
<td>Start non-sedating, antiseizure medicines (e.g., levetiracetam) for seizure prophylaxis. Dexamethasone 10 mg IV every 12 hours for 2-3 days, or longer for persistent symptoms. Consider taper for a total steroid exposure of greater than 3 days. If no improvement after 24 hours or worsening of neurologic toxicity, increase the dose and/or frequency of dexamethasone up to a maximum of 20 mg IV every 6 hours. If no improvement after another 24 hours, rapidly progressing symptoms, or life-threatening complications arise, give methylprednisolone (2 mg/kg) loading dose, followed by 2 mg/kg divided 4 times a day; taper within 7 days.</td>
</tr>
<tr>
<td>Grade 3</td>
<td>Start non-sedating, antiseizure medicines (e.g., levetiracetam) for seizure prophylaxis. Dexamethasone 10 to 20 mg IV every 8 to 12 hours. Steroids are not recommended for isolated Grade 3 headaches. If no improvement after 24 hours or worsening of neurologic toxicity, escalate to methylprednisolone (dose and frequency as per Grade 2). If cerebral edema is suspected, consider hyperhydration and hyperosmolar therapy. Give high-dose methylprednisolone (1-2 g, repeat every 24 hours if needed; taper as clinically indicated) and cyclophosphamide 1.5 g/m².</td>
</tr>
<tr>
<td>Grade 4</td>
<td>Start non-sedating, antiseizure medicines (e.g., levetiracetam) for seizure prophylaxis. Dexamethasone 20 mg IV every 6 hours. If no improvement after 24 hours or worsening of neurologic toxicity, escalate to methylprednisolone (dose and frequency as per Grade 2). If cerebral edema is suspected, consider hyperhydration and hyperosmolar therapy. Give high-dose methylprednisolone (1-2 g, repeat every 24 hours if needed; taper as clinically indicated), and cyclophosphamide 1.5 g/m².</td>
</tr>
</tbody>
</table>

Neurologic Toxicity

Monitor patients for signs and symptoms of neurologic toxicities (Table 2). Rule out other causes of neurologic symptoms. Provide intensive care supportive therapy for severe or life-threatening neurologic toxicities. If neurologic toxicity is suspected, manage according to the recommendations in Table 2.

If concurrent CRS is suspected during neurologic toxicity, administer:
- Corticosteroids according to the more aggressive intervention based on the CRS and neurologic toxicity grades in Tables 1 and 2
- Tocilizumab according to the CRS grade in Table 1
- Antiseizure medication according to the neurologic toxicity in Table 2
5 Warnings and Precautions

5.1 Cytokine Release Syndrome
Cytokine release syndrome (CRS), including fatal or life-threatening reactions, occurred following treatment with BREYANZI. Among patients receiving BREYANZI for LBL (N=418), CRS occurred in 46% (190/418), including ≥Grade 3 CRS (Lee grading system) in 3.1% of patients. In patients receiving BREYANZI after two or more lines of therapy for LBL, CRS occurred in 46% (122/268), including ≥Grade 3 CRS in 4.1% of patients. One patient had fatal CRS and 2 had ongoing CRS at time of death. The median time to onset was 5 days (range: 1 to 15 days). CRS resolved in 98% with a median duration of 5 days (range: 1 to 17 days).

In patients receiving BREYANZI after one line of therapy for LBL, CRS occurred in 45% (68/150), including Grade 3 CRS in 1.3% of patients. The median time to onset was 4 days (range: 1 to 63 days). CRS resolved in all patients with a median duration of 4 days (range: 1 to 16 days).

The most common manifestations of CRS (≥10%) included fever (94%), hypotension (42%), tachycardia (28%), chilling (23%), hypoxia (16%), and headache (12%) [see Adverse Reactions (6.1)].

Serious events that may be associated with CRS include cardiac arrhythmias (including atrial fibrillation and ventricular tachycardia), cardiac arrest, cardiac failure, diffuse alveolar damage, renal insufficiency, capillary leak syndrome, hypotension, hypoxia, and hemophagocytic lymphohistocytosis/macrophage activation syndrome (HLH/MAS) [see Adverse Reactions (6.1)].

Ensure that 2 doses of tocilizumab are available prior to infusion of BREYANZI.

Of the 418 patients who received BREYANZI for LBL, 23% received tocilizumab and/or a corticosteroid for CRS, including 10% who received tocilizumab only and 2.2% who received corticosteroids only.

Monitor patients daily for at least 7 days following BREYANZI infusion at a REMS-certified healthcare facility for signs and symptoms of CRS. Monitor patients for signs or symptoms of CRS for at least 4 weeks after infusion. At the first sign of CRS, institute treatment with supportive care, tocilizumab, or tocilizumab and corticosteroids as indicated [see Dosage and Administration (2.3)].

Counsel patients to seek immediate medical attention should signs or symptoms of CRS occur at any time [see Patient Counseling Information (17)].

5.2 Neurologic Toxicities
Neurologic toxicities that were fatal or life-threatening, including immune effector cell-associated neurotoxicity syndrome (IGNS), occurred following treatment with BREYANZI. Serious events including cerebral edema and seizures occurred with BREYANZI. Fatal and serious cases of leukoencephalopathy, some attributable to fludarabine, also occurred.

In patients receiving BREYANZI after two or more lines of therapy for LBL, CAR T cell-associated neurologic toxicities occurred in 35% (85/268), including ≥Grade 3 cases in 12% of patients. Three patients had fatal neurologic toxicity and 7 had ongoing neurologic toxicity at time of death. The median time to onset of neurotoxicity was 8 days (range: 1 to 46 days). Neurologic toxicities resolved in 85% with a median duration of 12 days (range: 1 to 87 days).

In patients receiving BREYANZI after one line of therapy for LBL, CAR T cell-associated neurologic toxicities occurred in 27% (41/150) of patients, including Grade 3 cases in 7% of patients. The median time to onset of neurologic toxicity was 8 days (range: 1 to 63 days). The median duration of neurologic toxicity was 6 days (range: 1 to 119 days).

In all patients combined receiving BREYANZI for LBL, neurologic toxicities occurred in 33% (136/418), including ≥Grade 3 cases in 10% of patients. The median time to onset was 8 days (range: 1 to 63), with 87% of cases developing by 16 days. Neurologic toxicities resolved in 85% of patients with a median duration of 11 days (range: 1 to 119 days).

Of patients developing neurotoxicity, 77% (105/136) also developed CRS.

The most common neurologic toxicities (≥5%) included encephalopathy (20%), tremor (13%), aphasia (8%), headache (6%), diziness (6%), and delirium (5%).

Monitor patients daily at least for at least 7 days following BREYANZI infusion at a REMS-certified healthcare facility for signs and symptoms of neurologic toxicities and assess for other causes of neurological symptoms. Monitor patients for signs or symptoms of neurologic toxicities for at least 4 weeks after infusion and treat promptly. Manage neurologic toxicity with supportive care and/or corticosteroids as needed [see Dosage and Administration (2.3)].

Counsel patients to seek immediate medical attention should signs or symptoms of neurologic toxicity occur at any time [see Patient Counseling Information (17)].

5.3 BREYANZI REMS
Because of the risk of CRS and neurologic toxicities, BREYANZI is available only through a restricted program under a Risk Evaluation and Mitigation Strategy (REMS) called the BREYANZI REMS [see Boxed Warning and Warnings and Precautions (5.1, 5.2)]. The required components of the BREYANZI REMS are:

- Healthcare facilities that dispense and administer BREYANZI must be enrolled and comply with the REMS requirements.

- Certified healthcare facilities must have on-site, immediate access to tocilizumab.

- Ensure that a minimum of 2 doses of tocilizumab are available for each patient for infusion within 2 hours after BREYANZI infusion, if needed for treatment of CRS.

- Certified healthcare facilities must ensure that healthcare providers who prescribe, dispense, or administer BREYANZI are trained on the management of CRS and neurologic toxicities.

Further information is available at www.BreyanziREMS.com, or contact Bristol-Myers Squibb at 1-888-423-5436.

5.4 Hypersensitivity Reactions
Allergic reactions may occur with the infusion of BREYANZI. Serious hypersensitivity reactions, including anaphylaxis, may be due to dimethyl sulfoxide (DMSO).

5.5 Serious Infections
Severe infections, including life-threatening or fatal infections, have occurred in patients after BREYANZI infusion.

In patients receiving BREYANZI for LBL, infections of any grade occurred in 36%, with Grade 3 or higher infections occurring in 12% of all patients. Grade 3 or higher infections with an unspecified pathogen occurred in 7%, bacterial infections in 4.3%, viral infections in 1.9%, and fungal infections in 0.5%.

Febrile neutropenia developed after BREYANZI infusion in 8% of patients with LBL. Febrile neutropenia may be concurrent with CRS. In the event of febrile neutropenia, evaluate for infection and manage with broad-spectrum antibiotics, fluids, and other supportive care as medically indicated.

Monitor patients for signs and symptoms of infection before and after BREYANZI administration and treat appropriately. Administer prophylactic antimicrobials according to standard institutional guidelines.

Avoid administration of BREYANZI in patients with clinically significant, active systemic infections.

Viral Reactivation
Hepatitis B virus (HBV) reactivation, in some cases resulting in fulminating hepatitis, hepatic failure, and death, can occur in patients treated with drugs directed against B cells.

In patients who received BREYANZI for LBL, 15 of the 16 patients with a prior history of HBV were treated with concurrent antiviral suppressive therapy. Performance screening for HBV, HCV, and HIV in accordance with clinical guidelines before collection of cells for manufacturing. In patients with prior history of HBV, consider concurrent antiviral suppressive therapy to prevent HBV reactivation per standard guidelines.

5.6 Prolonged Cytopenias
Patients may exhibit cytopenias not resolved for several weeks following lymphodepleting chemotherapy and BREYANZI infusion.

Grade 3 or higher cytopenias persisted at Day 29 following BREYANZI infusion in 36% of patients with LBL and included thrombocytopenia in 28%, neutropenia in 21%, and anemia in 6%.

Monitor complete blood counts prior to and after BREYANZI administration.

5.7 Hypogammaglobulinemia
B-cell aplasia and hypogammaglobulinemia can occur in patients receiving BREYANZI.

In patients receiving BREYANZI for LBL, hypogammaglobulinemia was reported as an adverse reaction in 11% of patients. Hypogammaglobulinemia, either as an adverse reaction or laboratory IgG level below 500 mg/dL after infusion, was reported in 28% of patients.

Monitor immunoglobulin levels after treatment with BREYANZI and manage using infection precautions, antibiotic prophylaxis, and immunoglobulin replacement as clinically indicated.

Live Vaccines
The safety of immunization with live viral vaccines during or following BREYANZI treatment has not been studied. Vaccination with live virus vaccines is not recommended for at least 6 weeks prior to the start of lymphodepleting chemotherapy during BREYANZI treatment, and until immune recovery following treatment with BREYANZI.

5.8 Secondary Malignancies
Patients treated with BREYANZI may develop secondary malignancies. Monitor lifelong for secondary malignancies. In the event that a secondary malignancy occurs, contact Bristol-Myers Squibb at 1-888-905-4553 for reporting and to obtain instructions on collection of patient samples for testing.

5.9 Effects on Ability to Drive and Use Machines
Due to the potential for neurologic events, including altered mental status or seizures, patients receiving BREYANZI are at risk for developing altered or decreased consciousness or impaired coordination in the 8 weeks following BREYANZI administration. Advise patients to refrain from driving and engaging in hazardous occupations or activities, such as operating heavy or potentially dangerous machinery, for at least 8 weeks.
6. ADVERSE REACTIONS

The following adverse reactions are described elsewhere in the labeling:

- Cytokine Release Syndrome [see Warnings and Precautions (5.1, 5.3)]
- Neurologic Toxicities [see Warnings and Precautions (5.2, 5.3)]
- Hypersensitivity Reactions [see Warnings and Precautions (5.4)]
- Serious Infections [see Warnings and Precautions (5.5)]
- Prolonged Cytophenias [see Warnings and Precautions (5.6)]
- Hypogammaglobulinemia [see Warnings and Precautions (5.7)]

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 safety data described in the WARNINGS and PRECAUTIONS reflect exposure to a single dose of BREYANZI in 418 patients with relapsed or refractory R/R LBLCL: one randomized, open-label study with 89 patients (TRANSFORM) and two open-label, single-arm studies with 61 patients (PILOT) and 298 patients (TRANSCEND).

Relapsed or Refractory LBLCL After One Line of Therapy

TRANSFORM

The safety of BREYANZI was evaluated in the TRANSFORM, a randomized, open-label, multicenter study, in which patients with primary refractory LBLCL or relapse within 1 year of first-line chemotherapy received BREYANZI (N=89) or standard therapy (N=31) [see Clinical Studies (14)]. Patients had not yet received treatment for relapsed or refractory lymphoma and were potential candidates for autologous HSCT. The trial excluded patients who were ineligible for transplant or who had age >75 years, ECOG performance status >1, history of central nervous system (CNS) disorders (such as seizures or cerebrovascular ischemia), uncontrolled infection, CrCl <45 mL/min, ALT >5 times the upper limit of normal (ULN), LVEF <40%, or absolute neutrophil count (ANC) <1.0 × 10⁹ cells/L or platelets <50 × 10⁹ cells/L in the absence of bone marrow involvement.

The planned dose of BREYANZI was 100 × 10⁹ CAR-positive viable T cells. The median age of the BREYANZI-treated population was 59 years (range: 20 to 74 years); 47% were male; 58% were white, 11% were Asian, and 5% were black.

Serious adverse reactions occurred in 38% of patients. The most common nonlaboratory, serious adverse reactions (>2%) were CRS, sepsis, fever, febrile neutropenia, headache, aphasia, COVID-19 infection, and pulmonary embolism.

Table 3 presents selected nonlaboratory adverse reactions in patients treated with BREYANZI, and Table 4 describes selected new or worsening Grade 3 or 4 laboratory abnormalities.

The most common nonlaboratory adverse reactions (≥20%) were fever, CRS, musculoskeletal pain, headache, fatigue, nausea, constipation, and dizziness.

### Table 3: Adverse Reactions in ≥10% of Patients Treated with BREYANZI in the TRANSFORM Study (N=89)

<table>
<thead>
<tr>
<th>Adverse Reaction</th>
<th>Any Grade (%)</th>
<th>Grade 3 or Higher (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Blood and lymphatic system disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Febrile neutropenia</td>
<td>10</td>
<td>10</td>
</tr>
<tr>
<td>Cardiac disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Tachycardia[2]</td>
<td>15</td>
<td>1.1</td>
</tr>
<tr>
<td>Gastrointestinal disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Nausea</td>
<td>24</td>
<td>0</td>
</tr>
<tr>
<td>Constipation</td>
<td>20</td>
<td>2.2</td>
</tr>
<tr>
<td>Abdominal pain[3]</td>
<td>18</td>
<td>0</td>
</tr>
<tr>
<td>Abdominal pain[3]</td>
<td>13</td>
<td>2.2</td>
</tr>
<tr>
<td>Vomiting</td>
<td>11</td>
<td>0</td>
</tr>
<tr>
<td>General disorders and administration site conditions</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fever</td>
<td>55</td>
<td>3.4</td>
</tr>
<tr>
<td>Fatigue[2]</td>
<td>28</td>
<td>1.1</td>
</tr>
<tr>
<td>Edema[2]</td>
<td>13</td>
<td>0</td>
</tr>
<tr>
<td>Immune system disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cytokine release syndrome</td>
<td>49</td>
<td>1.1</td>
</tr>
<tr>
<td>Infections and infestations[4]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bacterial infectious disorders</td>
<td>12</td>
<td>6</td>
</tr>
<tr>
<td>Infections with pathogen unspecified</td>
<td>12</td>
<td>6</td>
</tr>
<tr>
<td>Sepsis[7]</td>
<td>10</td>
<td>7</td>
</tr>
</tbody>
</table>

(Continued)
The safety of BREYANZI was evaluated in the PILOT study, a single-arm open-label study in transplant-ineligible patients with R/R LBCL after one line of chemomunotherapy. The study enrolled patients who were ineligible for high-dose therapy and autologous HSCT due to organ function or age, but who had adequate organ function for CAR-T cell therapy. Patients with a history of relevant CNS disorders (such as seizures or cerebrovascular ischemia), ECOG performance status >2, or uncontrolled systemic immunosuppression were ineligible. The median age of the study population was 74 years (range: 53 to 84 years), 90% were age ≥65 years, 61% were male. The ECOG performance status was 0 or 1 in 74% of patients and 2 in 26% of patients; 25% had CrCl <60 mL/min; 20% had a baseline ANC <1000/µL.

Serious adverse reactions occurred in 33% of patients. The most common nonlaboratory, serious adverse reactions (≥2%) were CRS, confusion, state, gastrointestinal hemorrhage, muscular weakness, musculoskeletal pain, pulmonary embolism, and sepsis. Table 5 presents selected nonlaboratory adverse reactions, and Table 6 describes selected new or worsening Grade 3 or 4 laboratory abnormalities.

The most common nonlaboratory adverse reactions (≥20%) were fatigue, CRS, fever, nausea, encephalopathy, hypotension, musculoskeletal pain, and edema.

Table 5: Adverse Reactions in ≥10% of Patients Treated with BREYANZI in the PILOT Study (N=61)

<table>
<thead>
<tr>
<th>Adverse Reaction</th>
<th>Any Grade (%)</th>
<th>Grade 3 or Higher (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cardiac disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Tachycardia</td>
<td>10</td>
<td>0</td>
</tr>
<tr>
<td>Gastrointestinal disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Nausea</td>
<td>25</td>
<td>1.6</td>
</tr>
<tr>
<td>Diarrhea</td>
<td>15</td>
<td>0</td>
</tr>
<tr>
<td>Constipation</td>
<td>11</td>
<td>0</td>
</tr>
<tr>
<td>General disorders and administration site conditions</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fatigue</td>
<td>44</td>
<td>1.6</td>
</tr>
<tr>
<td>Fever</td>
<td>38</td>
<td>1.6</td>
</tr>
<tr>
<td>Edema</td>
<td>20</td>
<td>0</td>
</tr>
<tr>
<td>Immune system disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cytokine release syndrome</td>
<td>39</td>
<td>1.6</td>
</tr>
<tr>
<td>Infections and infestations</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Infections with pathogen unspecified</td>
<td>13</td>
<td>4.9</td>
</tr>
<tr>
<td>Upper respiratory tract infection</td>
<td>13</td>
<td>0</td>
</tr>
<tr>
<td>Bacterial infectious disorders</td>
<td>10</td>
<td>3.3</td>
</tr>
<tr>
<td>Metabolism and nutrition disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Decreased appetite</td>
<td>13</td>
<td>1.6</td>
</tr>
<tr>
<td>Musculoskeletal and connective tissue disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Musculoskeletal pain</td>
<td>23</td>
<td>4.9</td>
</tr>
<tr>
<td>Nervous system disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Encephalopathy</td>
<td>23</td>
<td>4.9</td>
</tr>
<tr>
<td>Dizziness</td>
<td>16</td>
<td>1.6</td>
</tr>
<tr>
<td>Tremor</td>
<td>16</td>
<td>0</td>
</tr>
<tr>
<td>Headache</td>
<td>11</td>
<td>1.6</td>
</tr>
</tbody>
</table>

(Continued)
Table 7 presents selected nonlaboratory adverse reactions reported in patients treated with BREYANZI, and Table 8 describes selected new or worsening Grade 3 or 4 laboratory abnormalities.

The most common nonlaboratory adverse reactions (≥20%) were fatigue, CRS, musculoskeletal pain, nausea, headache, encephalopathy, infections (pathogen unspecified), decreased appetite, diarrhea, hypotension, tachycardia, dizziness, cough, constipation, abdominal pain, vomiting, and edema.

Table 7: Adverse Reactions in ≥10% of Patients Treated with BREYANZI in the TRANSCEND Study (N=268)

<table>
<thead>
<tr>
<th>Adverse Reaction</th>
<th>Any Grade (%)</th>
<th>Grade 3 or Higher (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Cardiac disorders</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Tachycardia^a</td>
<td>25</td>
<td>0</td>
</tr>
<tr>
<td><strong>Gastrointestinal disorders</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Nausea</td>
<td>33</td>
<td>1.5</td>
</tr>
<tr>
<td>Diarrhea</td>
<td>26</td>
<td>0.4</td>
</tr>
<tr>
<td>Constipation</td>
<td>23</td>
<td>0</td>
</tr>
<tr>
<td>Abdominal pain^b</td>
<td>21</td>
<td>3.0</td>
</tr>
<tr>
<td>Vomiting</td>
<td>21</td>
<td>0.4</td>
</tr>
<tr>
<td><strong>General disorders and administration site conditions</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fatigue</td>
<td>48</td>
<td>3.4</td>
</tr>
<tr>
<td>Edema</td>
<td>21</td>
<td>1.1</td>
</tr>
<tr>
<td>Fever</td>
<td>16</td>
<td>0</td>
</tr>
<tr>
<td>Chills</td>
<td>12</td>
<td>0</td>
</tr>
<tr>
<td><strong>Immune system disorders</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cytochrome release syndrome</td>
<td>46</td>
<td>4.1</td>
</tr>
<tr>
<td><strong>Infections and infestations</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Infection with pathogen unspecified</td>
<td>29</td>
<td>16</td>
</tr>
<tr>
<td>Bacterial infection</td>
<td>13</td>
<td>5</td>
</tr>
<tr>
<td>Upper respiratory tract infection</td>
<td>13</td>
<td>0.7</td>
</tr>
<tr>
<td>Viral infection</td>
<td>10</td>
<td>1.5</td>
</tr>
<tr>
<td><strong>Metabolism and nutrition disorders</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Decreased appetite</td>
<td>28</td>
<td>2.6</td>
</tr>
<tr>
<td><strong>Musculoskeletal and connective tissue disorders</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Musculoskeletal pain</td>
<td>37</td>
<td>2.2</td>
</tr>
<tr>
<td>Motor dysfunction</td>
<td>10</td>
<td>1.1</td>
</tr>
<tr>
<td><strong>Nervous system disorders</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Headache</td>
<td>30</td>
<td>1.1</td>
</tr>
<tr>
<td>Encephalopathy</td>
<td>29</td>
<td>9</td>
</tr>
<tr>
<td>Dizziness</td>
<td>24</td>
<td>2.6</td>
</tr>
<tr>
<td>Tremor</td>
<td>16</td>
<td>0</td>
</tr>
<tr>
<td>Peripheral neuropathy</td>
<td>11</td>
<td>0</td>
</tr>
<tr>
<td>Aphasia</td>
<td>10</td>
<td>2.2</td>
</tr>
<tr>
<td><strong>Psychiatric disorders</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Insomnia</td>
<td>14</td>
<td>0.4</td>
</tr>
<tr>
<td>Anxiety</td>
<td>10</td>
<td>0</td>
</tr>
<tr>
<td>Delirium</td>
<td>10</td>
<td>2.2</td>
</tr>
<tr>
<td><strong>Renal and urinary disorders</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Renal failure</td>
<td>11</td>
<td>3.0</td>
</tr>
<tr>
<td><strong>Respiratory, thoracic, and mediastinal disorders</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cough</td>
<td>23</td>
<td>0</td>
</tr>
<tr>
<td>Dyspnea</td>
<td>16</td>
<td>2.6</td>
</tr>
<tr>
<td><strong>Skin and subcutaneous tissue disorders</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rash</td>
<td>13</td>
<td>0.4</td>
</tr>
</tbody>
</table>

(Continued)
8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary
There are no available data with BREYANZI use in pregnant women. No animal reproductive and developmental toxicity studies have been conducted with BREYANZI to assess whether it can cause fetal harm when administered to a pregnant woman.

It is not known if BREYANZI has the potential to be transferred to the fetus. Based on the mechanism of action, if the transduced cells cross the placenta, they may cause fatal toxicity, including B-cell lymphocytopenia and hypergammaglobulinemia. Therefore, BREYANZI is not recommended for women who are pregnant, and pregnancy after BREYANZI infusion should be discussed with the treating physician.

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.

8.2 Lactation

Risk Summary
There is no information regarding the presence of BREYANZI in human milk, the effect on the breastfed infant, and the effects on milk production. The developmental and health benefits of breastfeeding should be considered along with the mother’s clinical need for BREYANZI and any potential adverse effects on the breastfed infant from BREYANZI or from the underlying maternal condition.

8.3 Females and Males of Reproductive Potential

Pregnancy Testing
Pregnancy status of females with reproductive potential should be verified. Sexually active females of reproductive potential should have a pregnancy test prior to starting treatment with BREYANZI.

Contraception
See the prescribing information for fludarabine and cyclophosphamide for information on the need for effective contraception in patients who receive lymphodepleting chemotherapy.

There are insufficient exposure data to provide a recommendation concerning duration of contraception following treatment with BREYANZI.

Infertility
There are no data on the effects of BREYANZI on fertility.

8.4 Pediatric Use
The safety and efficacy of BREYANZI have not been established in pediatric patients.
evidence for transformation and/or immortalization and no preferential integration near genes associated with oncogenic transformation.

14 CLINICAL STUDIES

Relapsed or Refractory LBCL After One Line of Therapy

TRANSFORM

A randomized, open-label, multicenter trial evaluated the efficacy of BREYANZI in adult patients with relapsed or refractory LBCL after first-line chemoimmunotherapy (TRANSFORM; NCT03575351). Patients had not yet received treatment for relapsed or refractory lymphoma, were potential candidates for autologous HSCT, and were required to have primary refractory disease or relapse within 12 months from complete response (CR) to initial chemoimmunotherapy. Eligibility criteria required adequate organ function and blood counts for HSCT.

In total, 184 patients were randomized in a 1:1 ratio to receive a single infusion of BREYANZI (planned dose, 100 × 10⁶ CAR-positive viable T cells) or to receive standard therapy consisting of 3 cycles of chemoimmunotherapy followed by high-dose therapy and autologous HSCT in patients who attained CR or PR. All patients underwent leukapheresis prior to randomization.

Patients randomized to BREYANZI were to receive lymphodepleting chemotherapy consisting of fludarabine 30 mg/m²/day and cyclophosphamide 300 mg/m²/day concurrently for 3 days followed by BREYANZI infusion 2 to 7 days after completion of lymphodepleting chemotherapy. Bridging chemotherapy was permitted between leukapheresis and the start of lymphodepleting chemotherapy. BREYANZI was administered in the inpatient (79%) and outpatient (21%) setting.

In the overall study population, the median age was 59 years (range: 20 to 75 years), 57% were male, 59% were white, 10% were Asian, and 4% were black. Diagnoses included de novo DLBCL NOS (55%), high-grade B-cell lymphoma (23%), primary mediastinal large B-cell lymphoma (10%), and DLBCL arising from indolent lymphoma (8%). Of these patients, 73% had primary refractory disease to last therapy and 27% had relapsed disease within 12 months of achieving CR to first-line therapy.

Of the 92 patients randomized to receive BREYANZI, 89 (97%) received BREYANZI. The median time from leukapheresis to product availability was 26 days (range: 19 to 84 days), and the median time from leukapheresis to product infusion was 36 days (range: 25 to 91 days). Fifty-eight (63%) patients received bridging therapy. One patient received a nonconforming product (manufacturing failure; 1%).

Of the 92 patients randomized to receive standard therapy, 91 started treatment and 43 (47%) received high-dose therapy and HSCT. The most common reason for not receiving HSCT was lack of efficacy of the salvage chemotherapy.

The primary efficacy measure was event-free survival (EFS) as determined by an independent review committee (IRC). Other efficacy measures included progression-free survival. Efficacy is summarized in Table 9 and Figure 1. The estimated 1-year EFS was 45% (95% CI: 29, 59) in the BREYANZI arm and 24% (95% CI: 14, 35) in the standard therapy arm.

Of the 92 patients in the BREYANZI arm, the estimated median DOR was not reached (95% CI: 7.9 months, NR) in patients who received CR (N=61) and 2.3 months (95% CI: 2.1, NR) in patients who achieved a best response of PR (N=18).

An interim analysis of overall survival was conducted at the time of the primary EFS analysis. The interim analysis of overall survival did not meet the criteria for statistical significance. Forty-six patients randomized to the standard therapy arm (50%) subsequently received BREYANZI on protocol.

### Table 9: Efficacy Results in Patients with Relapsed or Refractory LBCL (TRANSFORM Study)

<table>
<thead>
<tr>
<th>Outcome</th>
<th>BREYANZI Arm (N=92)</th>
<th>Standard Therapy Arm (N=92)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Event-Free Survival</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Number of events, n (%)</td>
<td>35 (38)</td>
<td>63 (69)</td>
</tr>
<tr>
<td>Median, months (95% CI)</td>
<td>10.1 [6.1, NR]</td>
<td>2.3 [2.2, 4.3]</td>
</tr>
<tr>
<td>Hazard ratio (95% CI)</td>
<td>0.34 [0.22, 0.52]</td>
<td></td>
</tr>
<tr>
<td>One-sided p-value</td>
<td>&lt;0.0001</td>
<td></td>
</tr>
<tr>
<td>Complete Response Rate, % (95% CI)</td>
<td>66 [56, 76]</td>
<td>39 [23, 50]</td>
</tr>
<tr>
<td>Difference in CR rate, % (95% CI)</td>
<td>27 [12, 41]</td>
<td></td>
</tr>
<tr>
<td>One-sided p-value</td>
<td>&lt;0.0001</td>
<td></td>
</tr>
<tr>
<td>Progression-Free Survival</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Number of events, n (%)</td>
<td>28 (30)</td>
<td>43 (47)</td>
</tr>
<tr>
<td>Median, months (95% CI)</td>
<td>14.8 [6.6, NR]</td>
<td>5.7 [3.9, 9.4]</td>
</tr>
<tr>
<td>Hazard ratio (95% CI)</td>
<td>0.41 [0.25, 0.66]</td>
<td></td>
</tr>
<tr>
<td>One-sided p-value</td>
<td>0.0001</td>
<td></td>
</tr>
</tbody>
</table>

NR = not reached; CI = confidence interval.

1. Per the Lugano criteria, as assessed by an IRC.
2. EFS is defined as time from randomization to the earliest date of disease progression or relapse, death from any cause, failure to achieve CR or PR by 9 weeks post-randomization, or start of new lymphoma therapy due to efficacy concerns.
4. Based on a stratified Cox proportional hazards model. For all stratified analyses, stratification was based on response to first-line therapy (primary refractory vs relapsed) and second-line age-adjusted International Prognostic Index.
6. p-value is compared with 0.012 of the allocated alpha for this pre-specified interim analysis.

### Figure 1: Kaplan-Meier Plot of IRC-Assessed Event-Free Survival

(Independence-to-Treat Analysis)

<table>
<thead>
<tr>
<th>Time (Months) From Randomization</th>
<th>Event-Free Survival Probability</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>BREYANZI</td>
</tr>
<tr>
<td>1</td>
<td>Standard Therapy</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Number at Risk</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Standard Therapy</td>
<td>92 83 66 35 32 23 21 16 12 11 10 6 4 4 4 2 2 0</td>
</tr>
<tr>
<td>BREYANZI</td>
<td>92 89 66 66 62 43 36 27 26 21 19 17 9 7 9 7 6 6 4 0</td>
</tr>
</tbody>
</table>

### PILOT

The efficacy of BREYANZI was evaluated in a single-arm, open-label, multicenter trial (PILOT; NCT03483103) in transplant-ineligible patients with relapsed or refractory LBCL after one line of chemoimmunotherapy. The study enrolled patients who were not eligible for high-dose therapy and autologous HSCT due to organ function or age, while also having adequate organ function for CAR-T cell therapy. The study required at least one of the following criteria: age ≤70 years, adjusted diffusion capacity of the lung for carbon monoxide (DLCO) ≤60%; LV EF <50%; creatinine clearance <60mL/min; AST or ALT greater than 2 × ULN, or ECOG performance status of 2. The planned dose of BREYANZI was 100 × 10⁶ CAR-positive viable T cells. Bridging therapy for disease control was permitted between leukapheresis and the start of lymphodepleting chemotherapy. Of the 61 patients treated with BREYANZI, 32 (53%) received bridging therapy.

BREYANZI was administered two to seven days following completion of lymphodepleting chemotherapy. The lymphodepleting chemotherapy regimen consisted of fludarabine 30 mg/m²/day and cyclophosphamide 300 mg/m²/day concurrently for 3 days. BREYANZI was administered in the inpatient (67%) and outpatient (33%) setting.
Of 74 patients who underwent leukapheresis, 61 (82%) received BREYANZI and comprise the main efficacy population; 1 (1.4%) received CAR-positive T cells that did not meet the product specifications for BREYANZI (manufacturing failure); and 12 (16%) did not receive CAR-positive T cells for other reasons.

Of the 61 patients who received BREYANZI, the median age was 74 years (range: 53 to 84 years); 61% were male, 89% were white, 3% were Asian, and 2% were black. Diagnoses included de novo DBCL NOS (51%), high-grade B-cell lymphoma (33%), and DLBCL arising from follicular lymphoma (15%). Of these patients, 53% had primary refractory disease, 23% had relapse within 12 months of completing first-line therapy, and 25% had relapse >12 months after first-line therapy.

Efficacy was based on complete response (CR) rate and duration of response (DOR), as determined by an independent review committee (IRC) using 2014 Lugano criteria (Tables 10 and 11). The median time to CR was 1 month (range 0.8 to 6.9 months).

Table 10: Response Rate in Relapsed or Refractory LBCL (PILOT Study)

<table>
<thead>
<tr>
<th>Outcome</th>
<th>BREYANZI-Treated (N=61)</th>
<th>All Leukapheresed Patients (N=74)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Overall Response, n (%)</td>
<td>49 (80%) [68%, 89%]</td>
<td>50 (88%) [56%, 78%]</td>
</tr>
<tr>
<td>Complete Response</td>
<td>33 (64%) [41%, 67%]</td>
<td>34 (46%) [34%, 58%]</td>
</tr>
<tr>
<td>Partial Response</td>
<td>16 (26%) [16%, 39%]</td>
<td>16 (22%) [13%, 33%]</td>
</tr>
</tbody>
</table>

CI=confidence interval; NR=Not reached

Table 11: Duration of Response in the PILOT Study

<table>
<thead>
<tr>
<th>Outcome</th>
<th>BREYANZI-Treated (N=61)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of Responders</td>
<td>49</td>
</tr>
<tr>
<td>DOR</td>
<td>11.2 [5.1, NR]</td>
</tr>
<tr>
<td>Range, months†</td>
<td>0.0+ to 22.8+</td>
</tr>
<tr>
<td>DOR if Best Response is CR</td>
<td>83% [64, 93]</td>
</tr>
<tr>
<td>Range, months</td>
<td>2.0+ to 22.8+</td>
</tr>
<tr>
<td>Rate at 6 months [95% CI]</td>
<td>68% [45, 83]</td>
</tr>
<tr>
<td>Rate at 12 months [95% CI]</td>
<td>82.8% [40.5, 100]</td>
</tr>
</tbody>
</table>

DOR=duration of response; CI=confidence interval; CR=complete response; PR=partial response; NR=not reached.

Table 12: Response Rate in Main Efficacy Population (TRANSCEND Study)

<table>
<thead>
<tr>
<th>Outcome</th>
<th>BREYANZI-Treated (N=192)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Overall Response Rate, n (%)</td>
<td>141 (73%) [67%, 80%]</td>
</tr>
<tr>
<td>Complete Response, n</td>
<td>104 (54%) [47%, 61%]</td>
</tr>
<tr>
<td>Partial Response, n</td>
<td>37 (19%) [14%, 26%]</td>
</tr>
</tbody>
</table>

CI=confidence interval.

Table 13: Duration of Response in the TRANSCEND Study

<table>
<thead>
<tr>
<th>Outcome</th>
<th>BREYANZI-Treated (N=192)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of Responders</td>
<td>141</td>
</tr>
<tr>
<td>DOR (Months)</td>
<td>16.7 [5.3, NR]</td>
</tr>
<tr>
<td>Range‡</td>
<td>0.0+ to 23.5+</td>
</tr>
<tr>
<td>DOR if Best Response is CR (Months)</td>
<td>1.4 [1.1, 2.2]</td>
</tr>
<tr>
<td>Range</td>
<td>0.0+ to 22.6+</td>
</tr>
</tbody>
</table>

DOR=duration of response; CI=confidence interval; CR=complete response; PR=partial response; NR=not reached.

Response durations were longer in patients who achieved a CR, as compared to patients with a best response of PR (Table 13). Of the 104 patients who achieved CR, 68 (65%) had remission lasting at least 6 months and 64 (62%) had remission lasting at least 9 months. Of the 287 patients who underwent leukapheresis and had radiographically evaluable disease, 27 additional patients achieved a response, apart from the responses noted in Table 12. The IRC-assessed overall response rate in the leukapheresed population (n=287) was 59% (95% CI: 53, 64), with a CR rate of 43% (95% CI: 37, 49) and PR rate of 15% (95% CI: 11, 20). These efficacy results include responses that may have been contributed solely by bridging therapy, responses after receipt of product outside of the intended dose range, and responses to product that did not meet release specifications.

15 REFERENCES

16 HOW SUPPLIED/STORAGE AND HANDLING

BREYANZI consists of genetically modified autologous T cells, supplied in vials as separate frozen suspensions of each CD8 component (NDC 73153-901-08) and CD4 component (NDC 73153-902-04). Each CD8 or CD4 component is packed in a carton containing up to 4 vials, depending upon the concentration of the cryopreserved drug product CAR-positive viable T cells. The cartons for each CD8 component and CD4 component are in an outer carton (NDC 73153-900-01). BREYANZI is shipped directly to the cell lab or clinical pharmacy associated with the infusion center in the vapor phase of a liquid nitrogen shipper. A Release for Infusion (RFI) Certificate for each component and patient-specific syringe labels are affixed inside the shipper.

- Confirm patient identity upon receipt.
- Store vials in the vapor phase of liquid nitrogen (less than or equal to minus 130°C) in a temperature-monitored system.
- Thaw BREYANZI prior to infusion [see Dosage and Administration (2.2)].

17 PATIENT COUNSELING INFORMATION

Advise the patient to read the FDA-approved patient labeling (Medication Guide).

Ensure that patients understand the risk (11%) of manufacturing failure. In case of a manufacturing failure, a second manufacturing of BREYANZI may be attempted. While the patient awaits the product, additional bridging therapy (not the lymphodepletion) may be necessary. This bridging therapy may be associated with adverse events during the pre-infusion period, which could delay or prevent the administration of BREYANZI.

Advise patients that they will be monitored daily for at least 7 days following the BREYANZI infusion at a REMS-certified healthcare facility and instruct patients to remain within 2 hours of a REMS-certified healthcare facility for at least 4 weeks following the infusion.

Prior to infusion, advise patients of the following risks:

- Cytokine Release Syndrome (CRS) – Signs and symptoms of CRS (fever, chills, hypotension, tachycardia, hypoxia, and fatigue). Counsel patients to seek immediate medical attention should signs or symptoms of CRS occur at any time [see Warnings and Precautions (5.1) and Adverse Reactions (6.1)].

- Neurologic Toxicities – Signs or symptoms associated with neurologic events including encephalopathy, confusion, decreased consciousness, speech disorders, tremor, and seizures. Counsel patients to seek immediate medical attention should signs or symptoms of neurologic toxicity occur at any time [see Warnings and Precautions (5.2) and Adverse Reactions (6.1)].

- Serious Infections – Signs or symptoms associated with infection [see Warnings and Precautions (5.5) and Adverse Reactions (6.1)].

- Prolonged Cytopenias – Signs or symptoms associated with bone marrow suppression including neutropenia, anemia, thrombocytopenia, or febrile neutropenia [see Warnings and Precautions (5.6) and Adverse Reactions (6.1)].

Advise patients of the need to:

- Contact Bristol-Myers Squibb at 1-888-805-4555 if they are diagnosed with a secondary malignancy [see Warnings and Precautions (5.8)].

- Refrain from driving or operating heavy or potentially dangerous machines until at least 8 weeks after BREYANZI administration [see Warnings and Precautions (5.9)].

Manufactured by:

Juno Therapeutics Inc., a Bristol-Myers Squibb Company, Bothell, WA 98021.
US License No.: 2156.

Celgene Corporation, a Bristol-Myers Squibb Company, Summit, NJ 07901.
US License No.: 2252.

Bristol-Myers Squibb Company, Devens, MA 01434.
US License No.: 1713.

BREYANZI® is a trademark of Juno Therapeutics, Inc., a Bristol-Myers Squibb Company.
BREPI.004/MG.004
MEDICATION GUIDE
BREYANZI® (pronounced braye an' zee)
(lisocabtagene maraleucel)

Read this Medication Guide before you start your BREYANZI treatment. The more you know about your treatment, the more active you can be in your care. Talk with your healthcare provider if you have questions about your health condition or treatment. Reading this Medication Guide does not take the place of talking with your healthcare provider about your treatment.

What is the most important information I should know about BREYANZI?

BREYANZI may cause side effects that are life-threatening and can lead to death. Call your healthcare provider or get emergency help right away if you get any of the following:
- difficulty breathing
- fever (100.4°F/38°C or higher)
- chills/shaking chills
- confusion
- severe nausea, vomiting, diarrhea
- fast or irregular heartbeat
- dizziness/lightheadedness
- severe fatigue or weakness

It is important that you tell your healthcare providers that you have received BREYANZI and to show them your BREYANZI Patient Wallet Card. Your healthcare provider may give you other medicines to treat your side effects.

What is BREYANZI?

BREYANZI is a prescription medicine used to treat large B cell lymphoma, a type of non-Hodgkin lymphoma, when:
- your first treatment has not worked or your cancer returned within a year of your first treatment OR
- your first treatment has not worked or your cancer returned after the first treatment, and you are not eligible for hematopoietic stem cell transplantation because of medical conditions or age OR
- two or more kinds of treatment have not worked or stopped working.

BREYANZI is different than other cancer medicines because it is made from your own white blood cells, which have been genetically modified to recognize and attack your lymphoma cells.

Before getting BREYANZI, tell your healthcare provider about all your medical problems, including if you have or have had:
- Neurologic problems (such as seizures, stroke, or memory loss)
- Lung or breathing problems
- Heart problems
- Liver problems
- Kidney problems
- A recent or active infection

Tell your healthcare provider about all the medications you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements.

How will I receive BREYANZI?

- BREYANZI is made from your own white blood cells, so your blood will be collected by a process called “leukapheresis” (LOO-kuh-feh-REE-sis).
- It takes about 3-4 weeks from the time your cells are received at the manufacturing site and are available to be shipped back to your healthcare provider, but the time may vary.
- Before you get BREYANZI, you will get 3 days of chemotherapy to prepare your body.
- When your BREYANZI is ready, your healthcare provider will give it to you through a catheter (tube) placed into your vein (intravenous infusion). BREYANZI is given as infusions of 2 different cell types.
  - You will receive infusions of one cell type, immediately followed by the other cell type.
  - The time for infusion will vary but will usually be less than 15 minutes for each of the 2 cell types.
- During the first week, you will be monitored daily by the facility where you received your treatment.
BREYANZI® (lisocabtagene maraleucel)

- You should plan to stay within 2 hours of the location where you received your treatment for at least 4 weeks after getting BREYANZI. Your healthcare provider will check to see that your treatment is working and help you with any side effects that may occur.
- You may be hospitalized for side effects and your healthcare provider will discharge you if your side effects are under control, and it is safe for you to leave the hospital.
- Your healthcare provider will want to do blood tests to follow your progress. It is important that you do have your blood tested. If you miss an appointment, call your healthcare provider as soon as possible to reschedule.

**What should I avoid after receiving BREYANZI?**

- Do not drive, operate heavy machinery, or do other activities that could be dangerous if you are not mentally alert, for at least 8 weeks after you get BREYANZI. This is because the treatment can cause temporary memory and coordination problems, including sleepiness, confusion, dizziness, and seizures.
- Do not donate blood, organs, tissues, or cells for transplantation.

**What are the possible or reasonably likely side effects of BREYANZI?**

The most common side effects of BREYANZI are:

- fatigue
- difficulty breathing
- fever (100.4°F/38°C or higher)
- chills/shaking chills
- confusion
- difficulty speaking or slurred speech
- severe nausea, vomiting, diarrhea
- headache
- dizziness/lightheadedness
- fast or irregular heartbeat
- swelling

BREYANZI can increase the risk of life-threatening infections that may lead to death. Tell your healthcare provider right away if you develop fever, chills, or any signs or symptoms of an infection.

BREYANZI can lower one or more types of your blood cells (red blood cells, white blood cells, or platelets). After treatment, your healthcare provider will test your blood to check for this. Tell your healthcare provider right away if you get a fever, are feeling tired, or have bruising or bleeding.

Having BREYANZI in your blood may cause a false-positive HIV test result by some commercial tests.

These are not all the possible side effects of BREYANZI. Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

**General information about the safe and effective use of BREYANZI**

Medicines are sometimes prescribed for purposes other than those listed in a Medication Guide. If you would like more information about BREYANZI, talk with your healthcare provider. You can ask your healthcare provider for information about BREYANZI that is written for health professionals.

For more information, go to BREYANZI.com or call 1-888-805-4555.

Manufactured by:
Juno Therapeutics Inc., a Bristol-Myers Squibb Company, Bothell, WA 98021. US License No.: 2156.
Celgene Corporation, a Bristol-Myers Squibb Company, Summit, NJ 07901. US License No.: 2252.
Bristol-Myers Squibb Company, Devens, MA 01434. US License No.: 1713.

BREYANZI® is a trademark of Juno Therapeutics, Inc., a Bristol-Myers Squibb Company.
BREM.004 06/2023

This Medication Guide has been approved by the U.S. Food and Drug Administration.