POWERFUL MIGRAINE PREVENTION STARTS TODAY

Help your patients meet their migraine-free potential today with one 30-minute treatment every three months.1,2

Discussion guide
Use this guide to help support conversations with your patients about VYEPTI.

VYEPTI is indicated for the preventive treatment of migraine in adults.

IMPORTANT SAFETY INFORMATION
CONTRAINDICATIONS
VYEPTI is contraindicated in patients with serious hypersensitivity to eptinezumab-jjmr or to any of the excipients. Reactions have included angioedema.

Please see additional Important Safety Information throughout this document. For more information, see the accompanying full Prescribing Information, including Patient Information.
ABOUT VYEPTI (eptinezumab-jjmr)
The first and only CGRP blocker given as an intravenous (IV) treatment

“What results can I expect with VYEPTI”

In adults with 4-14 headache days per month, during Months 1-3:

VYEPTI 100 mg and 300 mg reduced average monthly migraine days by 3.9 and 4.3, respectively (vs. 3.2 with placebo).

Information is based on a 12-month clinical study of 665 adults with a history of episodic migraine.

In adults with 15-26 headache days per month, during Months 1-3:

VYEPTI 100 mg and 300 mg reduced average monthly migraine days by 7.7 and 8.2, respectively (vs. 5.6 with placebo).

Information is based on a 6-month clinical study of 1,072 adults with a history of chronic migraine.

EARLY BENEFIT SEEN WITHIN THE FIRST 7 DAYS, AND AS SOON AS DAY 1 POST-INFUSION, VS. PLACEBO

VYEPTI 100 mg is the recommended dose.
Some patients may benefit from the available and approved 300 mg dose.

APPROVED USE
VYEPTI is a prescription medicine used for the preventive treatment of migraine in adults.

IMPORTANT SAFETY INFORMATION
Do not receive VYEPTI if you are allergic to eptinezumab-jjmr or its ingredients. See the end of the accompanying Patient Information leaflet for a complete list of ingredients in VYEPTI.

VYEPTI may cause allergic reactions. Call your healthcare provider or get emergency medical help right away if you have any symptoms of an allergic reaction: rash, swelling of your face, lips, tongue, or throat, if you have trouble breathing, hives or redness in your face.

Please see additional Important Safety Information throughout this document. For more information, see the accompanying full Prescribing Information, including Patient Information.
“How much of a reduction in migraine days might I experience?”¹,³,⁴

In adults with 4-14 headache days per month, during Months 1-3:

**22% of people treated with VYEPTI 100 mg experienced ≥75% reduction in migraine days** (vs. 16% with placebo).*

* This was not a significant difference.

In adults with 4-14 headache days per month, during the 12-month study period:

**63% of people treated with VYEPTI 100 mg were migraine-free for a month or more** (vs. 48% with placebo).

In adults with 15-26 headache days per month, during Months 1-3:

**27% of people treated with VYEPTI 100 mg experienced ≥75% reduction in migraine days** (vs. 15% with placebo).

In adults with 15-26 headache days per month, during the 6-month study period:

**35% of people treated with VYEPTI 100 mg were migraine-free for a month or more** (vs. 22% with placebo).

† A migraine-free month was defined as a consecutive 28-day period in which the patient experienced no migraine. 100% responder rate was a prespecified secondary endpoint, but not a prespecified analysis. Therefore, no definitive conclusions should be drawn.

**TREATMENT BENEFIT WAS MAINTAINED THROUGH MONTH 6 VS. PLACEBO**

**IMPORTANT SAFETY INFORMATION (continued)**

Before starting VYEPTI, tell your healthcare provider about all your medical conditions, including if you are pregnant or plan to become pregnant, or you are breastfeeding or plan to breastfeed.

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

Please see additional Important Safety Information throughout this document.

For more information, see the accompanying full Prescribing Information, including Patient Information.
“How does VYEPTI work?”

VYEPTI is designed to work fast. After each 30-minute infusion, 100% of the medication is available in the bloodstream.

VYEPTI is a CGRP (calcitonin gene-related peptide), blocker. This means it binds to a protein called CGRP, preventing it from activating CGRP receptors. The activation of CGRP receptors is thought to play a large role in causing migraine.

The relationship between the body’s response to the medication and the way VYEPTI prevents migraine is unknown.

Safety results from two clinical trials of ~1,700 people with migraine

- 6% of people treated with VYEPTI 100 mg, and 8% treated with VYEPTI 300 mg, experienced a stuffy nose and scratchy throat, compared to 6% treated with placebo.
- 1% of people treated with VYEPTI 100 mg, and 2% treated with VYEPTI 300 mg, experienced allergic reactions, compared to 0% treated with placebo.
- 1.9% discontinued VYEPTI due to side effects.

Patient counseling information

- **Hypersensitivity reactions:** Inform patients that hypersensitivity reactions, including angioedema, urticaria, facial flushing, and rash, can occur. Advise patients to contact you immediately if signs or symptoms of hypersensitivity reactions occur.

- **Pregnancy:** Advise patients to notify you if they become pregnant during treatment or plan to become pregnant.

- **Lactation:** Inform patients to notify you if they are breastfeeding or plan to breastfeed.

Please see additional Important Safety Information throughout this document. For more information, see the accompanying full Prescribing Information, including Patient Information.
“Is there a support program available for VYEPTI?”*

VYEPTI GO is a personalized patient support program designed to assist you during your VYEPTI treatment journey.

With VYEPTI GO, you will have access to:

- Patient Navigators available by phone 6 days per week to answer questions and direct you to helpful resources, such as the VYEPTI Copay Assistance Program for commercial patients.*
- A registered nurse available by phone to help you better understand migraine, and provide treatment tips and VYEPTI education.
- Text and email communications to assist you with appointment scheduling and reminders.

Get all the details at vyepti.com or call 1-833-4-VYEPTI (1-833-489-3784), and press option 2 for VYEPTI GO.

* Terms and conditions apply. Visit vyepti.com to learn more.

The VYEPTI Copay Assistance Program

Commercial patients may pay as little as $5 for VYEPTI treatments with the VYEPTI Copay Assistance Program. The VYEPTI Copay Assistance Program helps eligible patients with their payments for VYEPTI. A maximum annual limit of $4,000 and other restrictions may apply.

To be eligible, patients must:
- Be aged 17 years or older.
- Have a valid prescription for VYEPTI that is not eligible for reimbursement through any state or federal healthcare programs.
- Meet all other eligibility requirements set forth in the VYEPTI Copay Assistance Program terms and conditions.

Visit vyepti.com for full terms and conditions.

IMPORTANT SAFETY INFORMATION (continued)

The most common side effects of VYEPTI include stuffy nose and scratchy throat, and allergic reactions. These are not all the possible side effects of VYEPTI. Tell your healthcare provider if you have any side effect that bothers you or that does not go away.

You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch or call 1-800-FDA-1088.

Please see additional Important Safety Information throughout this document.

For more information, see the accompanying full Prescribing Information, including Patient Information.
IMPORTANT SAFETY INFORMATION (continued)

WARNINGS AND PRECAUTIONS

Hypersensitivity reactions: Hypersensitivity reactions, including angioedema, urticaria, facial flushing, and rash, have occurred with VYEPTI in clinical trials. Most hypersensitivity reactions occurred during infusion and were not serious, but often led to discontinuation or required treatment. Serious hypersensitivity reactions may occur. If a hypersensitivity reaction occurs, consider discontinuing VYEPTI, and institute appropriate therapy.

ADVERSE REACTIONS

The most common adverse reactions (≥2% and at least 2% or greater than placebo) in the clinical trials for the preventive treatment of migraine were nasopharyngitis and hypersensitivity.

For more information, see the accompanying full Prescribing Information, including Patient Information.


©2020 Lundbeck. All rights reserved.
VYEPTI and VYEPTI GO are trademarks of Lundbeck Seattle BioPharmaceuticals, Inc. EPT-B-100241
HIGHLIGHTS OF PRESCRIBING INFORMATION
These highlights do not include all the information needed to use VYEPTI safely and effectively. See full prescribing information for VYEPTI.

VYEPTI™ (eptinezumab-jjmr) injection, for intravenous use
Initial U.S. Approval: 2020

------------------INDICATIONS AND USAGE-------------------
VYEPTI is a calcitonin gene-related peptide antagonist indicated for the preventive treatment of migraine in adults (1)

--------------DOSAGE AND ADMINISTRATION--------------
• Must dilute before use. For intravenous infusion only (2.1, 2.2)
• Recommended dosage is 100 mg as an intravenous infusion over approximately 30 minutes every 3 months. Some patients may benefit from a dosage of 300 mg (2.1, 2.3)
• Dilute only in 100 mL of 0.9% Sodium Chloride Injection, USP (2.2)

------------DOSAGE FORMS AND STRENGTHS------------
Injection: 100 mg/mL solution in a single-dose vial (3)

---------------------CONTRAINDICATIONS----------------------
VYEPTI is contraindicated in patients with serious hypersensitivity to eptinezumab-jjmr or to any of the excipients (4)

--------------WARNINGS AND PRECAUTIONS--------------
Hypersensitivity Reactions: Reactions have included angioedema, urticaria, facial flushing, and rash. If a hypersensitivity reaction occurs, consider discontinuing VYEPTI and initiate appropriate therapy (5.1)

--------------ADVERSE REACTIONS--------------
The most common adverse reactions (≥2% and 2% or greater than placebo) were nasopharyngitis and hypersensitivity (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Lundbeck at 1-800-455-1141 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

See 17 for PATIENT COUNSELING INFORMATION and FDA-approved patient labeling.

Revised: 2/2020

FULL PRESCRIBING INFORMATION: CONTENTS*

1 INDICATIONS AND USAGE
2 DOSAGE AND ADMINISTRATION
  2.1 Recommended Dosing
  2.2 Dilution Instructions
  2.3 Infusion Administration Instructions
3 DOSAGE FORMS AND STRENGTHS
4 CONTRAINDICATIONS
5 WARNINGS AND PRECAUTIONS
  5.1 Hypersensitivity Reactions
6 ADVERSE REACTIONS
  6.1 Clinical Trials Experience
  6.2 Immunogenicity
8 USE IN SPECIFIC POPULATIONS
  8.1 Pregnancy
  8.2 Lactation
  8.4 Pediatric Use
  8.5 Geriatric Use

11 DESCRIPTION
12 CLINICAL PHARMACOLOGY
  12.1 Mechanism of Action
  12.2 Pharmacodynamics
  12.3 Pharmacokinetics
13 NONCLINICAL TOXICOLOGY
  13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility
14 CLINICAL STUDIES
16 HOW SUPPLIED/STORAGE AND HANDLING
  16.1 How Supplied
  16.2 Storage and Handling
17 PATIENT COUNSELING INFORMATION

*Sections or subsections omitted from the full prescribing information are not listed.
FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

VYEPTI is indicated for the preventive treatment of migraine in adults.

2 DOSAGE AND ADMINISTRATION

2.1 Recommended Dosing
The recommended dosage is 100 mg administered by intravenous infusion every 3 months. Some patients may benefit from a dosage of 300 mg administered by intravenous infusion every 3 months.

2.2 Dilution Instructions
VYEPTI requires dilution prior to administration. Dilute only in 100 mL 0.9% Sodium Chloride Injection, USP. The infusion bags must be made of polyvinyl chloride (PVC), polyethylene (PE), or polyolefin (PO). Use appropriate aseptic technique when preparing VYEPTI solution for intravenous infusion. VYEPTI single-dose vials contain no preservative; discard unused portion remaining in the vial.

Dilution
100 mg dose:
To prepare the solution, withdraw 1 mL of VYEPTI from a single-dose vial using a sterile needle and syringe. Inject the 1 mL content into a 100 mL bag of 0.9% Sodium Chloride Injection, USP.

300 mg dose:
To prepare the solution, withdraw 1 mL of VYEPTI from each of 3 single-dose vials using a sterile needle and syringe. Inject the resulting 3 mL content into a 100 mL bag of 0.9% Sodium Chloride Injection, USP.

Storage and Handling of Diluted Product
Gently invert the VYEPTI solution to mix completely. Do not shake. Following dilution, VYEPTI solution must be infused within 8 hours. During this time, VYEPTI solution should be stored at room temperature, 20°C to 25°C (68°F to 77°F). Do not freeze.

2.3 Infusion Administration Instructions
Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit. Do not use if the liquid contains visible particulate matter or is cloudy or discolored [see Dosage Forms and Strengths (3)].

No other medications should be administered through the infusion set or mixed with VYEPTI. VYEPTI is for intravenous infusion only; infuse over approximately 30 minutes. Do not administer VYEPTI as an intravenous push or bolus injection. Use an intravenous infusion set with a 0.2 micron or 0.22 micron in-line or add-on sterile filter. After the infusion is complete, flush the line with 20 mL of 0.9% Sodium Chloride Injection, USP.

3 DOSAGE FORMS AND STRENGTHS

VYEPTI is a clear to slightly opalescent, colorless to brownish-yellow solution available as follows:

- Injection: 100 mg/mL in a single-dose vial
4 CONTRAINDICATIONS

VYEPTI is contraindicated in patients with serious hypersensitivity to eptinezumab-jjmr or to any of the excipients in VYEPTI. Reactions have included angioedema [see Warnings and Precautions (5.1)].

5 WARNINGS AND PRECAUTIONS

5.1 Hypersensitivity Reactions

Hypersensitivity reactions, including angioedema, urticaria, facial flushing, and rash, have occurred with VYEPTI in clinical trials. Most hypersensitivity reactions occurred during infusion and were not serious, but often led to discontinuation or required treatment. Serious hypersensitivity reactions may occur. If a hypersensitivity reaction occurs, consider discontinuing VYEPTI, and institute appropriate therapy [see Contraindications (4) and Patient Counseling Information (17)].

6 ADVERSE REACTIONS

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

• Hypersensitivity Reactions [see Warnings and Precautions (5.1)].

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 clinical practice.

The safety of VYEPTI was evaluated in 2076 patients with migraine who received at least one dose of VYEPTI, representing 1615 patient-years of exposure; of these, 1524 patients were exposed to 100 mg or 300 mg. Across all doses, 1872 patients were exposed for at least 6 months and 991 patients were exposed for 12 months. In the placebo-controlled clinical studies (Study 1 and Study 2) of 1372 patients, 579 patients received at least one dose of VYEPTI 100 mg, 574 patients received at least one dose of VYEPTI 300 mg, and 588 patients received placebo [see Clinical Studies (14)]. Approximately 86% were female, 89% were white, and the mean age was 40.4 years at study entry.

The most common (incidence at least 2% and at least 2% greater than placebo) adverse reactions in the clinical trials for the preventive treatment of migraine were nasopharyngitis and hypersensitivity.

Table 1 summarizes the adverse reactions that occurred during Study 1 and Study 2.

Table 1. Adverse Reactions Occurring with an Incidence of at Least 2% for VYEPTI and at Least 2% Greater than Placebo in Studies 1 and 2

<table>
<thead>
<tr>
<th>Adverse Reactions</th>
<th>VYEPTI 100 mg N=579 %</th>
<th>VYEPTI 300 mg N=574 %</th>
<th>Placebo N=588 %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nasopharyngitis</td>
<td>6</td>
<td>8</td>
<td>6</td>
</tr>
<tr>
<td>Hypersensitivity reactions*</td>
<td>1</td>
<td>2</td>
<td>0</td>
</tr>
</tbody>
</table>

* Hypersensitivity reactions includes multiple related adverse event terms, such as hypersensitivity, pruritus, and flushing/hot flush that occurred on the day of dosing.

In Study 1 and Study 2, 1.9% of patients treated with VYEPTI discontinued treatment because of adverse reactions [see Warnings and Precautions (5.1)].
6.2 Immunogenicity

As with all therapeutic proteins, there is 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 eptinezumab-jjmr in the studies described below with the incidence of antibodies in other studies or to other products may be misleading.

In patients receiving VYEPTI 100 mg or 300 mg every 3 months, the incidence of anti-eptinezumab-jjmr antibody development in Study 1 (up to 56 weeks) was 20.6% (92/447), and 41.3% (38/92) of those patients developed anti-eptinezumab-jjmr neutralizing antibodies. In Study 2 (up to 32 weeks), the incidence of anti-eptinezumab-jjmr antibody development was 18.3% (129/706), and 34.9% (45/129) of those patients developed anti-eptinezumab-jjmr neutralizing antibodies. In an open-label study with 84 weeks of treatment, 18% (23/128) of patients developed anti-eptinezumab-jjmr antibodies, and 39% (9/23) of those patients developed anti-eptinezumab-jjmr neutralizing antibodies.

Although the results from both studies showed no clear evidence of an impact from development of anti-eptinezumab-jjmr antibodies, including neutralizing antibodies, on the safety and efficacy profiles of VYEPTI, the available data are too limited to make definitive conclusions.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

There are no adequate data on developmental risks associated with the use of VYEPTI in pregnant women.

No adverse developmental effects were observed following administration of eptinezumab-jjmr to pregnant animals at doses greater than those used clinically [see Data].

In the U.S. general population, the estimated background risk of major birth defects and miscarriages in clinically recognized pregnancies is 2%-4% and 15%-20%, respectively. The estimated rate of major birth defects (2.2%-2.9%) and miscarriage (17%) among deliveries to women with migraine are similar to rates reported in women without migraine.

Clinical Considerations

Disease-Associated Maternal and/or Embryo/Fetal Risk

Published data have suggested that women with migraine may be at increased risk of preeclampsia and gestational hypertension during pregnancy.

Data

Animal Data

When eptinezumab-jjmr (0, 75, or 150 mg/kg) was administered weekly to female rats and rabbits by intravenous injection throughout organogenesis, no adverse effects on embryofetal development were observed. The higher dose tested (150 mg/kg) is 30 times the maximum recommended human dose (MRHD) of 300 mg, on a body weight basis (mg/kg).

When eptinezumab-jjmr (0, 75, or 150 mg/kg) was administered weekly to female rats throughout pregnancy and lactation, no adverse effects on pre- and postnatal development were observed. The higher dose tested (150 mg/kg) is 30 times the MRHD, on a mg/kg basis.
8.2 Lactation

Risk Summary

There are no data on the presence of eptinezumab-jjmr in human milk, the effects on the breastfed infant, or the effects on milk production. The developmental and health benefits of breastfeeding should be considered along with the mother’s clinical need for VYEPTI and any potential adverse effects on the breastfed infant from VYEPTI or from the underlying maternal condition.

8.4 Pediatric Use

Safety and effectiveness in pediatric patients have not been established.

8.5 Geriatric Use

Clinical studies of VYEPTI did not include sufficient numbers of patients aged 65 and over to determine whether they respond differently from younger patients.

11 DESCRIPTION

Eptinezumab-jjmr is a humanized immunoglobulin G1 (IgG1) monoclonal antibody specific for calcitonin gene-related peptide (CGRP) ligand. Eptinezumab-jjmr has an approximate molecular weight of 143 kD. Eptinezumab-jjmr is produced in Pichia pastoris yeast cells by recombinant DNA technology.

VYEPTI (eptinezumab-jjmr) injection is a sterile, preservative-free, clear to slightly opalescent, colorless to brownish-yellow solution, for intravenous infusion. VYEPTI is supplied as a 100 mg/mL single-dose vial. Each mL contains 100 mg eptinezumab-jjmr formulated in L-histidine (1 mg), L-histidine hydrochloride monohydrate (2.8 mg), polysorbate 80 (0.15 mg), sorbitol (40.5 mg), and Water for Injection, USP, at a pH of 5.8.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Eptinezumab-jjmr is a humanized monoclonal antibody that binds to calcitonin gene-related peptide (CGRP) ligand and blocks its binding to the receptor.

12.2 Pharmacodynamics

The relationship between the pharmacodynamic activity and the mechanism(s) by which eptinezumab-jjmr exerts its clinical effects is unknown.

12.3 Pharmacokinetics

Eptinezumab-jjmr exhibits linear pharmacokinetics and exposure increases proportionally with doses from 100 mg to 300 mg after intravenous administration. Steady-state plasma concentration is attained after the first dose with a once every 3-month dosing schedule.

Distribution

The central volume of distribution (Vc) for eptinezumab-jjmr is approximately 3.7 liters.

Metabolism & Elimination

Eptinezumab-jjmr is expected to be degraded by proteolytic enzymes into small peptides and amino acids.

The apparent clearance of eptinezumab-jjmr was 0.006 L/h, and the terminal elimination half-life was approximately 27 days.
Specific Populations
A population pharmacokinetic analysis assessing the effects of age, race, sex, and body weight did not suggest any clinically significant impact of these covariates on eptinezumab exposures.

Patients with Renal or Hepatic Impairment
No dedicated studies were conducted to assess the effects of renal or hepatic impairment on the pharmacokinetics of eptinezumab-jjmr. However, hepatic or renal impairment is not expected to affect the pharmacokinetics of eptinezumab-jjmr. A population pharmacokinetic analysis of integrated data from eptinezumab-jjmr clinical studies did not reveal clinically significant impact on pharmacokinetics of patients with hepatic or renal impairment.

Drug Interaction Studies
P450 Enzymes
Eptinezumab-jjmr is not metabolized by cytochrome P450 enzymes; therefore, interactions with concomitant medications that are substrates, inducers, or inhibitors of cytochrome P450 enzymes are unlikely.

Sumatriptan
The co-administration of a single dose of 300 mg eptinezumab-jjmr administered as an intravenous infusion (over a period of 1 hour ± 15 min) with a single dose of 6 mg sumatriptan administered subcutaneously did not significantly influence the pharmacokinetics of eptinezumab-jjmr or sumatriptan.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Carcinogenesis
The carcinogenic potential of eptinezumab-jjmr has not been assessed.

Mutagenesis
Genetic toxicology studies of eptinezumab-jjmr have not been conducted.

Impairment of Fertility
When eptinezumab-jjmr (0, 75, or 150 mg/kg) was administered weekly by intravenous injection to male and female rats prior to and during mating and continuing in females to gestation day 3-4, no adverse effects on fertility were observed. The higher dose tested (150 mg/kg) is 30 times the maximum recommended human dose of 300 mg, on a body weight basis (mg/kg).

14 CLINICAL STUDIES
The efficacy of VYEPTI was evaluated as a preventive treatment of episodic and chronic migraine in two randomized, multicenter, placebo-controlled studies, both with 6-month double-blind periods: one study in patients with episodic migraine (Study 1) and one study in patients with chronic migraine (Study 2). VYEPTI was administered by intravenous infusion every 3 months in both studies; however, the primary endpoint was measured at 12 weeks.

Study 1: Episodic Migraine
Study 1 (NCT02559895) included adults with a history of episodic migraine (4 to 14 headache days per month, of which at least 4 were migraine days). A total of 665 patients were randomized to receive placebo (N=222), 100 mg VYEPTI (N=221), or 300 mg VYEPTI (N=222) every 3 months for 12 months. Patients were allowed to use concurrent acute migraine or headache medications, including migraine-specific medications (i.e., triptans, ergotamine derivatives), during the trial.

The study excluded patients with a history of cardiovascular disease (hypertension, ischemic heart disease), neurological disease, or cerebrovascular disease.

The primary efficacy endpoint was the change from baseline in mean monthly migraine days over Months 1-3. Secondary endpoints included the percentages of patients with 50% or greater, and 75% or greater reductions from baseline in monthly migraine days over Months 1-3.

Patients had a median age of 39 years (range: 18 to 71 years), 84% were female, and 84% were white. The mean migraine frequency at baseline was approximately 8.6 migraine days per month and was similar across treatment groups.

VYEPTI treatment demonstrated statistically significant improvements compared to placebo for the primary efficacy endpoint, as shown in Table 2; secondary endpoints are also summarized in Table 2.

Table 2. Efficacy Endpoint Results in Study 1

<table>
<thead>
<tr>
<th></th>
<th>VYEPTI 100 mg</th>
<th>VYEPTI 300 mg</th>
<th>Placebo</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>N=221</td>
<td>N=222</td>
<td>N=222</td>
</tr>
<tr>
<td><strong>Monthly Migraine Days (MMD) – Months 1-3</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Change from baseline</td>
<td>-3.9</td>
<td>-4.3</td>
<td>-3.2</td>
</tr>
<tr>
<td>Difference from placebo</td>
<td>-0.7</td>
<td>-1.1</td>
<td></td>
</tr>
<tr>
<td>p-value</td>
<td>0.018</td>
<td>&lt;0.001</td>
<td></td>
</tr>
<tr>
<td><strong>≥50% MMD responders – Months 1-3</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>% Responders</td>
<td>49.8%</td>
<td>56.3%</td>
<td>37.4%</td>
</tr>
<tr>
<td>Difference from placebo</td>
<td>12.4%</td>
<td>18.9%</td>
<td></td>
</tr>
<tr>
<td>p-value</td>
<td>0.009*</td>
<td>&lt;0.001</td>
<td></td>
</tr>
<tr>
<td><strong>≥75% MMD responders – Months 1-3</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>% Responders</td>
<td>22.2%</td>
<td>29.7%</td>
<td>16.2%</td>
</tr>
<tr>
<td>Difference from placebo</td>
<td>6.0%</td>
<td>13.5%</td>
<td></td>
</tr>
<tr>
<td>p-value</td>
<td>NS**</td>
<td>&lt;0.001</td>
<td></td>
</tr>
</tbody>
</table>

* Nominal statistical significance
** NS = Not statistically significant

Figure 1 shows the mean change from baseline in average monthly migraine days in Study 1. Patients treated with VYEPTI at both doses had greater mean decreases from baseline in mean monthly migraine days over Months 1-3 compared to placebo-treated patients.
**Figure 1. Change from Baseline in Monthly Migraine Days in Study 1**

![Graph showing change from baseline in monthly migraine days over months 1 to 6 by treatment group.](image)

Figure 1 shows the change from baseline in monthly migraine days over months 1 to 6 by treatment group.

**Figure 2. Distribution of Change from Baseline in Mean Monthly Migraine Days over Months 1 to 3 by Treatment Group in Study 1**

![Bar chart showing distribution of change from baseline in mean monthly migraine days over months 1 to 3 by treatment group.](image)

Figure 2 shows the distribution of change from baseline in mean monthly migraine days through Month 3 by treatment group in 2-day increments.
Figure 3 demonstrates that greater percentages of placebo-treated patients had migraines on most days during the first 7 days of treatment compared to VYEPTI-treated patients in Study 1.

**Figure 3. Percentage of Patients with a Migraine from Day -1 (Day Prior to Infusion) to Day 7 in Study 1**

---

**Study 2: Chronic Migraine**

Study 2 (NCT02974153) included adults with a history of chronic migraine (15 to 26 headache days per month, of which at least 8 were migraine days). A total of 1072 patients were randomized and received placebo (N=366), 100 mg VYEPTI (N=356), or 300 mg VYEPTI (N=350) every 3 months for 6 months. Patients were allowed to use and to continue an established stable regimen of acute migraine or headache preventive medication (except onabotulinumtoxinA). Patients with a dual diagnosis of chronic migraine and medication overuse headache attributable to acute medication overuse (triptans, ergotamine, or combination analgesics greater than 10 days per month) were included in the study population. Patients using opioids or butalbital-containing products greater than 4 days per month were not allowed.

The study excluded patients with a history of cardiovascular disease (hypertension, ischemic heart disease), neurological disease, or cerebrovascular disease.

The primary efficacy endpoint was the change from baseline in mean monthly migraine days over Months 1-3. Secondary endpoints included the percentages of patients with 50% or greater and 75% or greater reductions from baseline in monthly migraine days over Months 1-3.

Patients had a median age of 41 years (range: 18 to 65 years), 88% were female, and 91% were white. Forty-one percent of patients were taking concomitant preventive medication for migraine. The mean migraine frequency at baseline was approximately 16.1 migraine days per month and was similar across treatment groups.

VYEPTI treatment demonstrated statistically significant improvements compared to placebo for the primary efficacy endpoint, as shown in Table 3; secondary endpoints are also summarized in Table 3.
<table>
<thead>
<tr>
<th>Table 3. Efficacy Endpoint Results in Study 2</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
</tr>
<tr>
<td>VYEPTI</td>
</tr>
<tr>
<td>100 mg</td>
</tr>
<tr>
<td>N=356</td>
</tr>
<tr>
<td>VYEPTI</td>
</tr>
<tr>
<td>300 mg</td>
</tr>
<tr>
<td>N=350</td>
</tr>
<tr>
<td>Placebo</td>
</tr>
<tr>
<td>N=366</td>
</tr>
<tr>
<td>Monthly Migraine Days (MMD) – Months 1-3</td>
</tr>
<tr>
<td>Change from baseline</td>
</tr>
<tr>
<td>Difference from placebo</td>
</tr>
<tr>
<td>p-value</td>
</tr>
<tr>
<td>≥50% MMD responders – Months 1-3</td>
</tr>
<tr>
<td>% Responders</td>
</tr>
<tr>
<td>Difference from placebo</td>
</tr>
<tr>
<td>p-value</td>
</tr>
<tr>
<td>≥75% MMD responders – Months 1-3</td>
</tr>
<tr>
<td>% Responders</td>
</tr>
<tr>
<td>Difference from placebo</td>
</tr>
<tr>
<td>p-value</td>
</tr>
</tbody>
</table>

Figure 4 shows the mean change from baseline in average monthly migraine days for Study 2. Patients treated with VYEPTI at both doses had greater mean decreases from baseline in mean monthly migraine days over Month 1-3 compared to placebo-treated patients.

**Figure 4. Change from Baseline in Monthly Migraine Days in Study 2**

Figure 5 shows the distribution of change from baseline in mean monthly migraine days through Month 3 by treatment group in 3-day increments.
Figure 5. Distribution of Change from Baseline in Mean Monthly Migraine Days over Months 1-3 by Treatment Group in Study 2

Figure 6 demonstrates that greater percentages of placebo-treated patients had migraines on individual days during the first 7 days of treatment compared to VYEPTI-treated patients in Study 2.
16 HOW SUPPLIED/STORAGE AND HANDLING

16.1 How Supplied
VYEPTI (eptinezumab-jjmr) injection is a clear to slightly opalescent, colorless to brownish-yellow solution supplied as:

Carton containing one 100 mg/mL single-dose vial - NDC 67386-130-51.

16.2 Storage and Handling
Store refrigerated at 2°C to 8°C (36°F to 46°F) in the original carton to protect from light until time of use. Do not freeze or shake.

The vial stopper is not made with natural rubber latex.

17 PATIENT COUNSELING INFORMATION

Advise the patient to read the FDA-approved patient labeling (Patient Information).

Hypersensitivity Reactions
Inform patients that hypersensitivity reactions, including angioedema, urticaria, facial flushing, and rash, can occur. Advise patients to contact their healthcare provider immediately if signs or symptoms of hypersensitivity reactions occur [see Warnings and Precautions (5.1)].

Pregnancy
Advise patients to notify their healthcare provider if they become pregnant during treatment or plan to become pregnant [see Use in Specific Populations (8.1)].
Lactation
Inform patients to notify their healthcare provider if they are breastfeeding or plan to breastfeed [see Use in Specific Populations (8.2)].

Manufactured by:
Lundbeck Seattle BioPharmaceuticals, Inc.
11804 North Creek Parkway South
Bothell, WA 98011 USA
U.S. License No. 2097

Vyepti is a trademark of Lundbeck Seattle BioPharmaceuticals, Inc.
**PATIENT INFORMATION**

**VYEPTI™ (vye ep’ tee)**

(eptinezumab-jjmr)

injection, for intravenous use

---

**What is VYEPTI?**

VYEPTI is a prescription medicine used for the preventive treatment of migraine in adults. It is not known if VYEPTI is safe and effective in children.

**Do not receive VYEPTI** if you are allergic to eptinezumab-jjmr or any of the ingredients in VYEPTI. See the end of this Patient Information leaflet for a complete list of ingredients in VYEPTI.

**Before you receive VYEPTI, tell your healthcare provider about all of your medical conditions, including if you:**

- are pregnant or plan to become pregnant. It is not known if VYEPTI will harm your unborn baby.
- are breastfeeding or plan to breastfeed. It is not known if VYEPTI passes into your breast milk. Talk to your healthcare provider about the best way to feed your baby while using VYEPTI.

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

**How will I receive VYEPTI?**

- VYEPTI will be given by a healthcare provider in a healthcare setting.
- VYEPTI is given by intravenous (IV) infusion in your vein.
- VYEPTI will be given over 30 minutes every 3 months.

If you have questions about your infusion schedule, ask your healthcare provider.

**What are the possible side effects of VYEPTI?**

VYEPTI may cause serious side effects, including:

- **Allergic reactions.** Allergic reactions can happen after receiving VYEPTI. Call your healthcare provider or get emergency medical help right away if you have any of the following symptoms of an allergic reaction:
  - rash
  - swelling of your face, lips, or tongue or throat
  - trouble breathing
  - hives
  - redness in your face

The most common side effects of VYEPTI include:

- stuffy nose and scratchy throat
- allergic reactions

These are not all of the possible side effects of VYEPTI.

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 VYEPTI.**

Medicines are sometimes prescribed for purposes other than those listed in the Patient Information leaflet. You can ask your pharmacist or healthcare provider for information about VYEPTI that is written for health professionals.

**What are the ingredients in VYEPTI?**

**Active ingredient:** eptinezumab-jjmr

**Inactive ingredients:** L-histidine, L-histidine hydrochloride monohydrate, polysorbate 80, sorbitol, and Water for Injection.

The vial stopper is not made with natural rubber latex.

---

*Manufactured by: Lundbeck Seattle BioPharmaceuticals, Inc., 11804 North Creek Parkway South, Bothell, WA 98011
US License Number: 2097
Vyepti is a trademark of Lundbeck Seattle BioPharmaceuticals, Inc.
For more information, call 1-833-4-VYEPTI (833-489-3784) or go to www.Vyepti.com.

This Patient Information has been approved by the U.S. Food and Drug Administration. Revised: 2/2020*