

VYEPTI CONNECT® ENROLLMENT FORM

Complete all required (*) fields, sign where indicated, and fax to VYEPTI CONNECT at 866-868-7071.
For questions, call 833-4-VYEPTI (833-489-3784), Monday - Friday, 8 AM - 8 PM ET.



Patient and Prescriber Information	PATIENT INFORMATION	DOB (MM/DD/YYYY)*	Sex ☐ M ☐ F	PRESCRIBER INFORMATION
	First Name*	MI		First Name* Last Name*
	Last Name*			Specialty NPI*
	Address*			PTAN Tax ID*
	City* State* ZIP*			Office Name*
	Preferred Phone Number ☐ Cell ☐ Home			Office Address*
	Alternate Phone Number ☐ Cell ☐ Home			City* State* ZIP*
	Email			Office Contact Name Title
	Alternate Contact Name			Phone* Fax*
	Relationship to Patient			Email
	Alternate Contact Phone			Preferred Method of Contact ☐ Phone ☐ Fax ☐ Both (Note: If no option is selected, will default to both. Updates can be made anytime by calling VYEPTI CONNECT)

Insurance Information	Insurance Information
Please include front and back copy of insurance card(s) along with this form. If copies of insurance cards are provided, the fields below are not required.	
☐ Patient Does Not Have Insurance. (If this is selected, insurance information should be left blank.)	
MEDICAL INSURANCE	
Primary Insurance* Phone*	Secondary Insurance Phone
Member ID* Group ID*	Member ID Group ID
Policy Holder Full Name	Policy Holder Full Name
Relationship to Patient DOB (MM/DD/YYYY)	Relationship to Patient DOB (MM/DD/YYYY)
PHARMACY INSURANCE	
Insurance Name Policy ID Group ID Rx BIN Rx PCN	

Infusion Provider
Select Site of Infusion*: ☐ Physician Office ☐ Home Infusion ☐ Infusion Center ☐ Hospital Outpatient
Please complete below, if infusion provider is not the same as the prescriber address information above.
Infusion Provider Name*
Address* City* State* ZIP*
Contact Name* Phone* Fax* Tax ID*
If Rx is included below, it will be sent to the indicated infusion provider.

Clinical
Please fax all clinical documentation and labs to VYEPTI CONNECT for inclusion with the prescription when sending to the dispensing pharmacy or infusion provider.
VYEPTI Dosage*: ☐ 100 mg ☐ 300 mg ☐ Other
Primary Diagnosis (ICD-10 Code)* Patient Is New to VYEPTI ☐ Yes ☐ No Date Migraines Started
Headache Days Per Month Date of Last Treatment Anticipated Infusion Date
Allergies
Previous Migraine Medication

Prescription
Rx COMPLETING THE PRESCRIPTION IS ONLY REQUIRED WHEN SENDING TO SPECIALTY PHARMACY OR INFUSION PROVIDER LISTED ABOVE.
Rx*: ☐ VYEPTI 100 mg/mL Single-Use Vial
Quantity (Vials)* Refills*
SIG: Infuse* mg over ~30 min once every 3 months.
Please attach a separate prescription if this does not comply with your state's prescription laws.
If using specialty pharmacy , indicate preference ☐ Orsini Healthcare ☐ AllianceRx Walgreens Prime
☐ Rx has already been provided to specialty pharmacy or the Infusion Provider listed above.
Prescriber's Signature* (Sign either box A or B below.) (Physician attests that this is his/her legal signature. NO STAMPS.)
A. Dispense as written Date B. Generic substitution permitted Date

I elect to OPT-IN to the following VYEPTI CONNECT options (choose all that apply):
☐ Final Coverage Status (including PA/Appeal Info) ☐ Upcoming Infusion Reminders ☐ Apply these choices for all of the patients I enroll in VYEPTI CONNECT

PRESCRIBER CERTIFICATION AND AUTHORIZATION (Signature and Date Required)

I certify that I have obtained authorization from the patient named above to release their protected health information to Lundbeck LLC and its affiliates, agents and service providers ("Lundbeck"), so Lundbeck may use it to provide patient support through the VYEPTI program(s) indicated on this form ("Programs"), and for the purposes described in the Patient Authorization on page 2 of this form. I certify that the information contained in this form is complete and accurate to the best of my knowledge and I will notify VYEPTI CONNECT if I learn of any updates to the patient's insurance. I have obtained the patient's authorization for Lundbeck to contact the patient regarding the Patient Authorization section on page 2 of this form. Lundbeck may contact me, including via email, fax, and telephone, to seek additional information relating to the Programs, VYEPTI, or the information contained on this form.

I certify that VYEPTI is medically necessary for the patient above, and if completed, I authorize or have obtained authorization for Lundbeck to send the prescription above and all information on this form to the dispensing pharmacy or infusion provider on this form or later selected by the patient or based on the patient's insurance.

SIGN Prescriber Name (Print)* Prescriber Signature* Date*

Please see the Important Safety Information on page 4. For more information, see the Full Prescribing Information, including Patient Information, or go to vyeptihcp.com.

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PATIENT TO COMPLETE ALL REQUIRED FIELDS AND SIGN



Patient Authorization for Use and Disclosure of Personal Health Information

(Signature and Date Required)

By signing below, I authorize my healthcare providers (including pharmacies and infusion providers) and health plans (together, my "Health Team") to disclose my personal health information relevant to my treatment or potential treatment with VYEPTI® (eptinezumab-jjmr), including my contact and other information provided on this enrollment form (my "Information"), to Lundbeck LLC and its affiliates, agents, representatives, and service providers (collectively, "Lundbeck"), so that Lundbeck can determine if I am eligible for the VYEPTI Program(s) I am seeking to participate in (collectively, the "Programs"), enroll me in the Programs if I am eligible, and undertake the activities listed below that involve the use and disclosure of my Information.

I authorize Lundbeck to share my Information with my Health Team to communicate about benefit and coverage status, medical care, and payments for my treatment with VYEPTI and to use and disclose my information as needed to: (1) administer the Programs; (2) provide me with patient support, including facilitating the provision of VYEPTI to me; (3) provide me with informational and promotional materials relating to Lundbeck products and/or my condition or treatment; (4) contact me for Program or research purposes or to provide information about Lundbeck products and services, including by phone, email and/or text message following my preferences indicated on this form, and including through messages that disclose that I take or may take VYEPTI; and (5) allow Lundbeck to analyze the usage patterns and the effectiveness of Lundbeck products, support, and programs and help develop new products, support, and programs, and for other Lundbeck general business and administrative purposes. I understand that Lundbeck may compensate my pharmacy providers for communicating with me about the benefits of Lundbeck products or services, and/or disclosing PHI pursuant to this authorization. I authorize my pharmacy providers to make such communications.

I understand that I am not required to sign this Authorization in order to receive healthcare benefits or treatment, including with Lundbeck products. I also understand that once my Information has been disclosed to Lundbeck, federal privacy laws may no longer restrict its disclosure and it might legally be redisclosed to others.

I understand that I may cancel this authorization at any time by sending a written cancellation notice to 2250 Perimeter Park Drive, Suite 300, Morrisville, North Carolina 27560, attention, Privacy Office. I understand that if I were to cancel the authorization, it would be invalid for further uses and disclosures of my Information, but that my cancellation would not invalidate any uses and disclosures of my Information made prior to the Program's receipt of my notice of withdrawal. I understand that if I do not cancel this authorization, the authorization will expire ten years from the date of signature (or the maximum period allowed by applicable state law, if less than ten years). I understand that I am entitled to receive a copy of this authorization once it has been signed.

PRINT PATIENT NAME*: _____

PATIENT CELL PHONE: _____ **OK TO TEXT:** ☐ Yes ☐ No

PATIENT EMAIL: _____

PATIENT SIGNATURE*: _____ **DATE*:** _____

IF UTILIZING A PERSONAL REPRESENTATIVE, THE FOLLOWING FIELDS WILL BE REQUIRED INSTEAD OF PATIENT SIGNATURE:

PRINT PERSONAL REPRESENTATIVE (AUTHORIZED TO SIGN FOR PATIENT)*: _____

PERSONAL REPRESENTATIVE SIGNATURE*: _____ **DATE*:** _____

RELATIONSHIP TO PATIENT*: ☐ Parent ☐ Spouse ☐ Power of Attorney ☐ Other _____



VYEPTI CONNECT Nurse Support is Available

Once enrolled in VYEPTI CONNECT, eligible patient will automatically be enrolled to receive VYEPTI CONNECT Nurse Support that includes helpful information and resources to help you stay on track while you are taking VYEPTI.

Nurse support is available Monday-Friday 8:00 a.m. – 8:00 p.m. (ET)
Call 833-4-VYEPTI (833-489-3784), option 2

☐ I prefer not to receive nursing support from VYEPTI CONNECT.

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PATIENT TO COMPLETE THIS PAGE AND SIGN

Commercially Insured Patients Can Enroll Today for VYEPTI CONNECT Copay Assistance

Financial assistance is available for eligible commercially insured patients through the VYEPTI CONNECT Copay Assistance Program. Eligible commercially insured patients may pay as little as \$0 for each VYEPTI® (eptinezumab-jjmr) infusion. Includes up to \$200 in administration fee cost sharing per treatment; a maximum annual benefit limit and other restrictions apply. Please see Terms and Conditions below for full details.

To enroll in the VYEPTI CONNECT Copay Assistance Program: 1. Answer the following five questions; 2. Review the Terms and Conditions; 3. Check the authorization button; 4. Sign and date below.

ANSWER

- ☐ YES ☐ NO Are you taking or planning to take VYEPTI for an approved indication? For more information about VYEPTI's indication, please review the "What is VYEPTI" section of the Patient Information in the attached PI or at vyepti.com.
- ☐ YES ☐ NO Are you 17 years of age or older?
- ☐ YES ☐ NO Are you enrolled in a health plan where the federal or state government could pay anything for your prescription, either all of it or part of it? Examples of government programs that pay for medication are Medicare or Medicaid, Medigap, VA, DOD, or TRICARE.
- ☐ YES ☐ NO Are you Medicare-eligible and enrolled in an employer-sponsored retiree health plan or prescription drug benefit program?
- ☐ YES ☐ NO Are you self-pay, meaning you pay the entire cost of the prescription out of your own pocket without any assistance from your insurance plan?

CHECK

- ☐ I have read and agree to the complete Terms and Conditions for the VYEPTI CONNECT Copay Assistance Program ("Program") below. I authorize the Program to provide payment directly to my healthcare provider, and not to me, for my out-of-pocket drug costs for VYEPTI. I authorize my healthcare provider to contact the Program on my behalf to initiate payment for VYEPTI after my infusion has taken place. I understand that I will be responsible for any out-of-pocket expenses for VYEPTI if (1) my healthcare provider does not request payment within 180 days of my date of service, or (2) if the Program determines that I am not eligible to receive copay assistance through the Program. I understand that by participating in the Program, I authorize Lundbeck LLC and its affiliates, agents, representatives, and service providers (together "Lundbeck") to use, disclose, and/or transfer my personal information in order to administer the Program, including contacting me by email, mail, or telephone to request or provide further information about the Program; evaluate the effectiveness of the Program; and provide resources and information related to VYEPTI. If my insurance changes, including becoming eligible to participate in any federal healthcare program such as Medicare or Medicaid, I will call VYEPTI CONNECT so my eligibility can be re-evaluated.

SIGN

PATIENT/AUTHORIZED CONTACT SIGNATURE: _____ DATE: _____

Please make sure that you've reviewed and checked the button above.

Terms and Conditions for the VYEPTI CONNECT Copay Assistance Program (the "Program")

Terms and Conditions: Commercially insured patients aged 17 years and older whose insurance policy provides coverage for VYEPTI® (eptinezumab-jjmr) (or the "Product") and whose insurance company does not pay for the entire cost of their prescription, may be eligible to receive financial assistance with out-of-pocket expenses for VYEPTI medication costs and VYEPTI administration costs (the "Offer"). Patients are not eligible for the Offer:

- (1) If they are self-pay, meaning the patient pays the entire cost of the prescription out of their own pocket; or
- (2) If the patient is enrolled in a health plan in which the federal or state government could pay for their prescription, either all of it or part of it; examples of government programs that pay for medication are Medicare or Medicaid, Medigap, VA, DOD, or TRICARE; or
- (3) If they are Medicare-eligible and enrolled in an employer-sponsored retiree health plan or prescription drug benefit program.

In addition, patients residing in Massachusetts, Minnesota, and Rhode Island are eligible only for copay assistance for the out-of-pocket Product costs and are not eligible for copay assistance for Product administration costs.

Eligible patients may pay as little as \$0 per infusion of VYEPTI. This Offer is subject to a Program assistance cap of \$200 per infusion for eligible administration costs, and a calendar year maximum on all Program assistance for all out-of-pocket expenses for VYEPTI (i.e., medication and infusion expenses) (the "Cap"). If the patient's total out-of-pocket expenses for VYEPTI exceed the Cap established by Lundbeck, the patient will be responsible for the additional balance. Patients should confirm their out-of-pocket cost with their pharmacy, or with their healthcare provider ("HCP"), prior to treatment. Copay claims must be submitted to the Program within 180 days of the patient's date of service.

The Offer is valid for use only with a valid prescription for VYEPTI (up to 300 mg) for the preventive treatment of migraine, at the time the prescription is filled by the pharmacist, or at the time the HCP administers VYEPTI to the patient. The Offer applies only to prescriptions filled before the Program expires or

terminates. The Offer applies to the patient's out-of-pocket costs for the Product and the eligible out-of-pocket Product administration costs. Any other fees related to the Product infusion are the responsibility of the patient. The patient or patient's HCP shall not submit any prescription or administration copays for payment to any public third-party payer, including Medicaid or Medicare, or to any other similar federal or state healthcare program. Patients are responsible for complying with any obligations or requirements imposed by their commercial insurance plans.

The Offer is intended solely for the eligible patient's benefit and is not transferable to any other person. The selling, purchasing, trading, or counterfeiting of the Offer is prohibited by law. The Offer has no cash value and may not be used in combination with any other discount, coupon, rebate, free trial, or similar offer for the specified VYEPTI prescription (including any pharmacy benefit manager or insurer program that adjusts patients' out-of-pocket costs for their drugs, such as through "maximizers" or "accumulators").

Lundbeck reserves the right to rescind, revoke, terminate, or amend the Offer at any time without notice. The Offer is intended to comply with all applicable laws and regulations including, without limitation, the federal Anti-Kickback Statute, its implementing regulations, and related guidance interpreting the federal Anti-Kickback Statute. The Offer is not health insurance. The Offer is valid only in the USA and Puerto Rico where allowed by law. This Offer is not conditioned on any past, present, or future Product purchase requirement, including refills. Patients can discontinue participation in the Program at any time and their questions and requests can be directed to 833-4-VYEPTI (833-489-3784) Monday through Friday, 8 AM - 8 PM ET.

The Offer will automatically renew each calendar year. If the patient no longer wishes to participate in the Offer, he/she can call and cancel at any time. By participating in the VYEPTI CONNECT Copay Assistance Program, the patient acknowledges and agrees that he/she is eligible to participate pursuant to the rules stated in these VYEPTI CONNECT Copay Assistance Program Terms and Conditions and that he/she understands and agrees to comply with these VYEPTI CONNECT Copay Assistance Program Terms and Conditions.

Please see the Important Safety Information on page 4. For more information, talk with your healthcare provider, and see the Full Prescribing Information and Patient Information, or go to vyepti.com.

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INDICATION

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 anaphylaxis and angioedema.

WARNINGS AND PRECAUTIONS

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

Constipation with Serious Complications: Constipation with serious complications has been reported following the use of monoclonal antibody CGRP antagonists, including VYEPTI, in the postmarketing setting. There were cases with monoclonal antibody CGRP antagonists that required hospitalization, including cases where surgery was necessary. In a majority of these cases, the onset of constipation was reported after the first dose; however, patients have also presented with constipation later on in treatment. The monoclonal antibody CGRP antagonist was discontinued in many of the reported cases of constipation with serious complications.

Monitor patients treated with VYEPTI for severe constipation and manage as clinically appropriate. The concurrent use of medications that reduce gastrointestinal motility may increase the risk for more severe constipation and the potential for constipation-related complications.

Hypertension: Development of hypertension and worsening of pre-existing hypertension have been reported following the use of CGRP antagonists, including VYEPTI, in the postmarketing setting. Some of the patients who developed new-onset hypertension had risk factors for hypertension. There were cases requiring initiation of pharmacological treatment for hypertension, and in some cases hospitalization. Hypertension may occur at any time during treatment, but was most frequently reported within 7 days of therapy initiation. The CGRP antagonist was discontinued in many of the reported cases.

Monitor patients treated with VYEPTI for new-onset hypertension or worsening of pre-existing hypertension, and consider whether discontinuation of VYEPTI is warranted if evaluation fails to establish an alternative etiology or blood pressure is inadequately controlled.

Raynaud's Phenomenon: Development of Raynaud's phenomenon and recurrence or worsening of pre-existing Raynaud's phenomenon have been reported in the postmarketing setting following the use of CGRP antagonists. In reported cases with monoclonal antibody CGRP antagonists, symptom onset occurred a median of 71 days following dosing. Many of the cases reported serious outcomes, including hospitalizations and disability, generally related to debilitating pain. In most reported cases, discontinuation of the CGRP antagonist resulted in resolution of symptoms.

VYEPTI should be discontinued if signs or symptoms of Raynaud's phenomenon develop, and patients should be evaluated by a healthcare provider if symptoms do not resolve. Patients with a history of Raynaud's phenomenon should be monitored for, and informed about the possibility of, worsening or recurrence of signs and symptoms.

ADVERSE REACTIONS

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

Please see the accompanying Full Prescribing Information, including Patient Information, or go to vyeptihcp.com.

