

# MAYZENT® Prescription Start Form

FAX  
1-877-750-9068

ENROLL ONLINE  
CoverMyMeds.com

QUESTIONS? CALL  
1-877-MAYZENT (1-877-629-9368)

 **MAYZENT**  
(siponimod) tablets  
0.25 mg • 2 mg

New Patient    Restarting Treatment

## 1. PATIENT INFORMATION

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|                       |                            |           |
|-----------------------|----------------------------|-----------|
| First Name            | Middle Initial             | Last Name |
| Sex: M F              | Date of Birth (MM/DD/YYYY) |           |
| Address (no PO boxes) |                            |           |
| City                  | State                      | ZIP       |
| Home Phone            | Cell Phone                 |           |

### Insurance Information

(include a copy of both sides of all cards, including secondary insurance)

### Medical Insurance(s)

|                         |                 |
|-------------------------|-----------------|
| Cardholder Name(s)      |                 |
| Insurance Carrier(s)    | Phone Number(s) |
| Cardholder ID Number(s) | Group Number(s) |

### Prescription Insurance

|                                |              |
|--------------------------------|--------------|
| Cardholder Name                |              |
| Prescription Insurance Carrier | Phone Number |
| ID Number                      | Group Number |
| PCN Number                     | BIN Number   |

E-mail (To make more communications convenient and paperless)

### Contact Preferences

OK to leave MAYZENT voicemail

### Preferred Language

English    Spanish

### Caregiver Information

|                                  |                 |
|----------------------------------|-----------------|
| Name of Caregiver (First & Last) | Caregiver Phone |
| Caregiver E-mail                 |                 |

## 2. PATIENT AUTHORIZATIONS AND ADDITIONAL CONSENTS

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I have read and agree to the Patient Authorization, which includes sharing of genetic information (page 3)

→ X

\_\_\_\_\_  
Patient/Legal Guardian Signature

\_\_\_\_\_/\_\_\_\_\_/\_\_\_\_\_  
Date of Signature (MM/DD/YYYY)

MAYZENT Co-pay Program: I have read and agree to the Terms and Conditions for participation (please see page 4)  
Receiving text messages and calls: I have read and agree to receiving marketing texts and calls as explained in the Telephone Consumer Protection Act (TCPA) Consent (optional, please see page 3)  
I have read and agree to the Fair Credit Reporting Act (FCRA) Authorization (optional, please see page 4)

## FOR OFFICE USE ONLY

## 3. PRESCRIBING PHYSICIAN INFORMATION

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|            |           |     |
|------------|-----------|-----|
| First Name | Last Name |     |
| Address    |           |     |
| City       | State     | ZIP |

|                         |                      |
|-------------------------|----------------------|
| Phone                   | Fax                  |
| State Medical License # | NPI #                |
| Office Contact Name     | Office Contact Phone |
| E-mail Address          |                      |

## 4. ASSESSMENT SUPPORT

**Request Co-pay Support Only:** Select checkbox to skip remaining Section 4 if the MAYZENT prescription will be sent directly to the Specialty Pharmacy and assessment/FDO support from Alongside™ MAYZENT is not needed.

**Assessment assistance requested\*\*:** (check all that apply)

### Blood Tests:

- CBC
- LFTs (transaminase & bilirubin)
- VZV antibody serology
- Genotype CYP2C9<sup>†</sup>

### Cardiac Evaluation:

- Electrocardiogram (ECG)

### Eye Exam:

- Macular edema screening<sup>‡</sup>

OR

**No assessment assistance requested and I clear for therapy:**

The following tests are completed or not required for this patient: CBC, LFTs (transaminase & bilirubin), VZV antibody serology, genotype CYP2C9 (to determine dose),<sup>†</sup> ECG, and a macular edema screening

**First-Dose Observation (FDO) period:** (select one)

- FDO assistance requested for this patient<sup>†</sup>
- FDO not required for patient<sup>†</sup>
- FDO required for patient<sup>†</sup> and will be performed outside of the support program

**Ship treatment initiation product for FDO to:**

|                      |                      |     |
|----------------------|----------------------|-----|
| Prescribing office   |                      |     |
| FDO location below:  |                      |     |
| Health Care Provider | Office Contact Phone |     |
| Address              |                      |     |
| City                 | State                | ZIP |

**CONTINUED ON THE NEXT PAGE**

5. PRESCRIPTION INFORMATION

**A DOSE IS REQUIRED TO INITIATE COVERAGE SUPPORT. PRODUCT WILL NOT BE DISPENSED UNTIL YOU HAVE CLEARED THE PATIENT FOR THERAPY.**

The recommended maintenance dose is 2 mg<sup>#</sup> taken orally once daily. **You will be required to confirm or change the patient's dose before dispense.**

**Cannot process form without this field completed**

**Primary diagnosis: ICD-10:G35 (select one)**

**Patient information:**

Name (First & Last) \_\_\_\_\_ Date of Birth (MM/DD/YYYY) \_\_\_\_\_ Other: \_\_\_\_\_

Active SPMS  
CIS/RRMS

**Preferred Specialty Pharmacy:**

Specialty Pharmacy \_\_\_\_\_

Phone \_\_\_\_\_

Fax \_\_\_\_\_

**Additional Notes:**

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**Pharmacy Prescription:**

For **2-mg** maintenance dose patients:  
MAYZENT (siponimod) tablets  
0.25mg tablet - Starter Pack.....NDC 0078-0979-12  
2mg tablet, Bottle of 30 tablets.....NDC 0078-0986-15

**Starter Pack:**  
No, patient already on therapy  
Yes, 0.25mg Starter Pack  
SIG: twelve 0.25mg tab for a 5 day supply  
**Day 1:** 1 x 0.25 mg | **Day 4:** 3 x 0.25 mg  
**Day 2:** 1 x 0.25 mg | **Day 5:** 5 x 0.25 mg  
**Day 3:** 2 x 0.25 mg  
**Qty:** 1 Starter Pack, then 1 refill

**Maintenance Dose:**  
2-mg tablet  
**SIG:** 1 tablet taken orally once a day  
**Qty:** 1 bottle (30 tablets/bottle), then 11 refills,  
or \_\_\_\_\_ refills

For **1-mg** maintenance dose patients:  
MAYZENT (siponimod) tablets  
0.25mg tablet, Bottle of 28 tablets.....NDC 0078-0979-50

**Treatment Initiation:**  
No, patient already on therapy  
Yes, 1-mg, 4 x 0.25-mg tablets  
**SIG: Day 1:** 1 x 0.25 mg | **Day 4:** 3 x 0.25 mg  
**Day 2:** 1 x 0.25 mg | **Day 5 and every day after:** 4 x 0.25 mg  
**Day 3:** 2 x 0.25 mg  
**Qty:** 1 Starter Pack, then 1 refill

**Maintenance Dose:**  
1-mg tablet  
**SIG:** 4 x 0.25-mg tablets taken orally once a day  
**Qty:** 4 bottles (28 tablets/bottle), then 11 refills,  
or \_\_\_\_\_ refills (1 refill = 4 bottles)

**Bridge to Commercial Coverage of MAYZENT\*\*:** (optional for commercially insured patients only)

Dispensed directly from Homescripts™ at no cost to the patient (check only 1 box):

For **2-mg** maintenance dose patients:

**Starter Pack:**  
No, patient already on therapy  
Yes, 0.25mg Starter Pack  
SIG: twelve 0.25mg tab for a 5 day supply  
**Day 1:** 1 x 0.25 mg | **Day 4:** 3 x 0.25 mg  
**Day 2:** 1 x 0.25 mg | **Day 5:** 5 x 0.25 mg  
**Day 3:** 2 x 0.25 mg  
**Qty:** 1 Starter Pack, then 1 refill

**Maintenance Dose:**  
2-mg tablet  
**SIG:** 1 tablet taken orally once a day  
**Qty:** 1 bottle (30 tablets/bottle), then 11 refills,  
or \_\_\_\_\_ refills

For **1-mg** maintenance dose patients:

**Treatment Initiation:**  
No, patient already on therapy  
Yes, 1-mg, 4 x 0.25-mg tablets  
**SIG: Day 1:** 1 x 0.25 mg | **Day 4:** 3 x 0.25 mg  
**Day 2:** 1 x 0.25 mg | **Day 5 and every day after:** 4 x 0.25 mg  
**Day 3:** 2 x 0.25 mg  
**Qty:** 4 bottles (28 tablets/bottle). No refills

**Maintenance Dose:**  
1-mg, 4 x 0.25-mg tablets  
**SIG:** 4 x 0.25-mg tablets taken orally once a day  
**Qty:** 4 bottles (28 tablets/bottle), then 11 refills,  
or \_\_\_\_\_ refills (1 refill = 4 bottles)

6. SIGNATURE AND PHYSICIAN ATTESTATION

**Cannot process form without this field completed**

**You must authorize these instructions by signing at the end of this section. We cannot process this form without your signature.**

I certify the above therapy is medically necessary and this information is accurate to the best of my knowledge. I certify I am the physician who has prescribed MAYZENT to the previously identified patient and I provided the patient with a description of the MAYZENT Support Program. For the purposes of transmitting these prescriptions, I authorize NPAF, Novartis Pharmaceuticals Corporation, and its affiliates, business partners, and agents to forward as my agent for these limited purposes these prescriptions electronically, by facsimile, or by mail to the appropriate dispensing pharmacies. I will not attempt to seek reimbursement for free product provided to me for purposes of performing a first-dose observation. I have read and agree to the Prescriber Authorization for the NPAF on page 4.

→ X \_\_\_\_\_ / / \_\_\_\_\_

**Prescriber Signature (Dispense as Written)** (Substitution Permissible) **Date of Signature (MM/DD/YYYY)**

ATTN: New York and Iowa prescribers, please submit electronic prescription to Homescripts Pharmacy, NPI#1528362076.

\*A benefits investigation to determine eligibility for certain support services will be completed even if assistance for treatment-related assessments are not requested.

†Listed services provided via Program may be available for commercially insured and uninsured patients starting MAYZENT. Offer not valid for services (i) performed in RI, (ii) for which payment may be made in whole or in part under federal or state health care programs, including, but not limited to, Medicare or Medicaid, or (iii) where prohibited by law. No purchase required. This Program is subject to termination or modification at any time. For questions regarding covered services, please contact your Alongside Coordinator.

‡An FDA-cleared or -approved test for the detection of CYP2C9 variants to direct the use of siponimod is not currently available.

§An ophthalmic evaluation of the fundus, including the macula, is recommended in all patients before starting treatment.

¶FDO is recommended for patients with certain preexisting cardiac conditions, including sinus bradycardia, first- or second-degree [Mobitz type I] AV block, or a history of myocardial infarction or heart failure.

\*\*For patients with a CYP2C9\*1\*3 or \*2\*3 genotype, the recommended maintenance dose of MAYZENT is 1 mg once daily (4 x 0.25 mg). For treatment initiation in these patients, the first month's supply should be used. MAYZENT should not be used in patients with a CYP2C9\*3\*3 genotype. An FDA-cleared or -approved test for the detection of CYP2C9 variants to direct the use of siponimod is not currently available.

\*\*\*Eligible patients must have commercial insurance and a valid prescription for MAYZENT. By participating, patient acknowledges intent to pursue insurance coverage for MAYZENT with their health care provider. Program requires the submission of a request for coverage within 9 months post-Program initiation in order to remain eligible. Patients will receive their maintenance drug supply each month for up to 12 months or until they receive insurance coverage approval, whichever occurs earlier. Program is not available to patients who are uninsured or whose medications are reimbursed in whole or in part by Medicare, Medicaid, TRICARE, or any other federal or state program, or where prohibited by law. Patients may be asked to reverify insurance coverage status during the course of the Program. No purchase necessary. Program is not health insurance, nor is participation a guarantee of insurance coverage. Other limitations may apply. Novartis reserves the right to rescind, revoke, or amend the Program and discontinue support at any time without notice.

*Please read the following carefully, then check the box where indicated on the previous pages.*

**PATIENT AUTHORIZATION.** I authorize my healthcare providers, pharmacies and health insurers, and their service providers (“Providers”) to disclose information relating to my insurance benefits, medical condition, treatment, **genetic information, including the results of genetic testing** and prescription details (“Personal Information”) to Novartis Pharmaceuticals Corporation, its affiliates and service providers (“Novartis”) and the Novartis Patient Assistance Foundation, Inc., and its service providers (“NPAF”) so they can provide the following support services (the “Services”):

- Help coordinate insurance coverage for, access to, and receipt of my medication.
- Communicate with me about possible financial assistance, including Novartis copay or NPAF programs, and, if I am enrolled, administer my participation in those programs.
- Communicate with me about my medication and treatment, including reminders, health and lifestyle tips, and product and other related information. Communications may be customized based on Personal Information obtained from my Providers.
- Conduct quality assurance and other internal business activities and ask for feedback related to the Services or my treatment.

In delivering the Services, Novartis and NPAF may share my Personal Information with each other, with my Providers, or with government agencies or other financial assistance programs that might help me pay for my medication. They may combine information collected from me with information collected from other sources and use that information to administer the Services. My pharmacies or other healthcare providers may receive payment from Novartis or NPAF for providing certain Services, such as medication or refill reminders, based on my enrollment or participation. Once I authorize disclosure of my Personal Information, it may no longer be protected by federal health privacy law and applicable state laws.

I understand I do not have to sign this Authorization to get my medication or insurance coverage, that I have a right to a copy, and can cancel this Authorization at any time by calling 1-877-629-9368 or writing to:

PO Box 2971

850 Twin Rivers Dr

Columbus, OH 43216-9532

OR

Customer Interaction Center

Novartis Pharmaceuticals Corporation, One Health Plaza

East Hanover, NJ 07936-1080

This Authorization will expire 5 years after I sign it, or earlier if required by state law, unless I cancel it sooner. If I cancel it, I may no longer qualify for Services from Novartis or NPAF, but it will not impact my Provider’s treatment or my insurance benefits. I also understand that if a Provider is disclosing my Personal Information to Novartis or NPAF on an authorized, ongoing basis, my cancellation will be effective with respect to that Provider as soon as they receive notice of my cancellation. Cancellation will not affect prior uses or disclosures.

#### **Telephone Consumer Protection Act (TCPA) Consent**

I consent to receive marketing calls and texts from and on behalf of the Novartis Group and NPAF, made with an auto-dialer or prerecorded voice, at the phone number(s) provided. I understand that my consent is not required or a condition of purchase. Number of messages will vary based on your Program selections. Message and data rates may apply. I understand that I can read the full Novartis Pharmaceuticals Corporation Privacy Policy at [www.usprivacy.novartis.com](http://www.usprivacy.novartis.com). Text STOP to opt out and HELP for help.

## PATIENT AUTHORIZATION (cont)

### **MAYZENT Co-pay Program Terms and Conditions**

Limitations apply. Valid only for those with private insurance. The Program includes the Co-Pay Card, Payment Card (if applicable), and Rebate, with a combined annual limit of \$18,000. Patient is responsible for any costs once limit is reached in a calendar year. Program not valid (i) under Medicare, Medicaid, TRICARE, VA, DoD, or any other federal or state health care program, (ii) where patient is not using insurance coverage at all, or (iii) where the patient's insurance plan reimburses for the entire cost of the drug. The value of this Program is exclusively for the benefit of patients and is intended to be credited towards patient out-of-pocket obligations and maximums, including applicable co-payments, coinsurance, and deductibles. Program is not valid where prohibited by law. Patient may not seek reimbursement for the value received from this Program from other parties, including any health insurance program or plan, flexible spending account, or health care savings account. Patient is responsible for complying with any applicable limitations and requirements of their health plan related to the use of the Program. Valid only in the United States and Puerto Rico. This Program is not health insurance. Program may not be combined with any third-party rebate, coupon, or offer. Proof of purchase may be required. Novartis reserves the right to rescind, revoke, or amend the Program and discontinue support at any time without notice.

### **Fair Credit Reporting Act (FCRA) Authorization**

I understand that I am providing "written instructions" authorizing NPAF and its vendor, under the FCRA, to obtain information from my credit profile or other information from the vendor, solely for the purpose of determining financial qualifications for programs administered by NPAF. I understand that I must affirmatively agree to these terms in order to proceed in this financial screening process.

### **Prescriber Authorization for the Novartis Patient Assistance Foundation, Inc. (NPAF)**

I certify that any medication received will be used only for the patient named on this form and will not be offered for sale, trade, or barter. Furthermore, no claim for reimbursement will be submitted concerning this medication, nor will any medication be returned for credit. I acknowledge that NPAF is exclusively for purposes of patient care and not for remuneration of any sort. I understand that NPAF may revise, change, or terminate programs at any time.

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Alongside is a trademark of Novartis AG.  
All trademarks are the property of their respective owners.

 **MAYZENT**  
(siponimod) tablets  
0.25 mg • 2 mg



**Novartis Pharmaceuticals Corporation**  
East Hanover, New Jersey 07936-1080

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