# **ENROLLMENT FORM** Phone: 888-APELLIS (888-273-5547) • Fax: 888-405-6966

 $Please\ ensure\ you\ and\ your\ patient\ complete\ all\ required\ information\ on\ the\ form\ and\ sign\ where\ indicated.$ \*Required Field



*Section 1. Support Rec  Check here for all available serv OR choose individual services	• vices (Checking "	'all" allows for support	services throughou	ut the patient journey	v, but only when	needed)
☐ Benefits Investigation Only	☐ Prior Auth	norization Assistance	e Co-pay	Program ially insured patients)	☐ Patient A	Assistance Progra or underinsured patie
Section 2. Patient Inforr	nation					
*First Name:		Middle Initial	*Last Name	a.		
Gender: ☐ Male ☐ Female [						
*Preferred Phone:						
Address:						
Preferred Language: ☐ English						
Section 3. Caregiver Inf Does patient have a caregiver with	-		o share informatic	on? 🗌 Yes 🔲 No (If	yes, please co	mplete this secti
Caregiver First Name:		Last Nar	ne:			
Preferred Phone:						
What is the caregiver's relationsh						
3	1	5.				
Section 4. Patient Insura *Does patient have insurance?						
Primary Insurance ( I If copy of	card is attached	d, check here)	Secondary Insu	ırance(☐ If copy	of card is attac	ched, check here
*Payer Name and Payer ID:			*Payer Name ar	nd Payer ID:		
Phone:			Phone:			
Policyholder Name:			Policyholder N	lame:		
Policyholder DOB:			Policyholder D	OOB:		
*Policy Number:			*Policy Number	r:		
Employer/Group Number:			Employer/Gro	oup Number:		
(Optional Section) Pharmacy (PB						
PBM Group ID:	РВМ	BIN/PCN:		_ PBM Phone Nur	mber:	
Section 4.1 Financial How many people live in the	patient's house	ehold?				-
Total annual household inco			Security income;	disability income	; any otner inco	ome):
☐ Less than \$150,000 ☐ Supporting documentation			also ask for proo	f of income at any	time for audit,	/verification.
*Section 5. Patient Auth	norization T	his form cannot b	e processed w	rithout the patie	ent's signatu	re.
☐ I have read and agre	e to 🗀	] I have read a	nd agree to	□ I hav	e read and	d agree to
Section 10.1 Authori		Section 10.2				uthorization
to Share Personal		Enroll in Apel	lisAssist Pat	ient to Re	eceive Ma	rketina
Health Information (	required)	Support Pro				ons (optiona
<b>&gt;</b>					/	
Patient Signature					Date (MM	/DD/YYYY)

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SYFOVRE (pegcetacoplan injection)

*Patient First Name:	Middle Initia	l:*Last Name:		(pegcetacopian injection) 15mg/0.1mL
Section 6. Prescribing Phy	sician Informa	ition		
Site of Service: Physician Office *Practice/Facility Name:			ical Center 🔲 Other:	
*Physician Name:			Specialty:	
*Phone:				
Address:				
Practice Tax ID#:	•		Physician PTAN#	
*NPI:	Practice NP	Physician NPI		
Section 7. Office Contact	Information			
*Primary Office Contact Name:				
*Phone:				
*Section 8. Prescription Ir	nformation 🔲 B	uy and bill 🔲 Specialty pha	rmacy	
<b>Geographic Atrophy Diagnosis</b> S	elect one diagnosis as	primary. For additional diagno	oses, please use the "second	ary diagnosis" section below.
Nonexudative age-related macul	ar degeneration	RIGHT EYE	LEFT EYE	BILATERAL
Advanced atrophic <b>without</b> subfoveal involvement		□ H35.3113	☐ H35.3123	□ H35.3133
Advanced atrophic <b>with</b> subfoveal involvement		□ H35.3114	□ H35.3124	□ H35.3134
Secondary Diagnosis: .	Has patient s	tarted treatment? 🔲 Yes, o	date of next treatment:	
, c		☐ No, a	nticipated date of first tre	atment:
Dispense:vial(s) of SYFOVRE® (	pegcetacoplan injec	tion) NDC: <b>73606-0020-0</b>	1 Refills #:	
SIG: ☐ Inject 15 mg (0.1 mL) intravitr		days Ancillary suppl 0 60)		on Kit (29G thin-wall needle and 5M filter needle)
*Section 9. Physician Dec	laration and Au	ıthorization		
This form allows Apellis Pharmaceutic patient support, resources and educe necessary written authorization from or other patient information included include, without limitation: (1) financial for alternate funding; and (3) Patient knowledge; (ii) the patient on this for through Apellis to my patient is not more prescribe, or use an Apellis medication and no claim for reimbursement will be a supported to the patient of the patient of the patient of the patient is not more supported to the patient of the patient	ation ("Patient Resou the patient reference therein for allowing p al assistance progran Resources. I certify the mhas a diagnosis for hade in exchange for on or Patient Resource	urces") to eligible patients wed above, or the patient's les participation in programs and ins; (2) verifying insurance conat: (i) the information in this an FDA-approved indication any express or implied agreese. I prescribed SYFOVRE so	who have been prescribed Segal guardian, to release to describe and services offered through overage and/or evaluating the form is complete and accument for SYFOVRE; (iii) any Patiement or understanding the blely on my clinical determination.	SYFOVRE. I have the Apellis the medical and/Apellis Assist, which may he patient's eligibility arate to the best of my ent Resource provided at I would recommend, nation and medical necessity

The SYFOVRE Co-Pay Program is for eligible patients enrolled in the ApellisAssist® program, are commercially insured, and are not covered under government insurance programs such as Medicare, Medicaid, VA/DoD, or TRICARE. The program assists only with the cost of SYFOVRE and its administration (injection) up to the program maximum. It does not assist with the cost of other administrations, medicines, procedures or office visits. Eligible patients residing in Massachusetts or Rhode Island can only receive assistance with the cost of SYFOVRE but not the cost of its administration. Patients receiving assistance through another program or foundation, free trial, or other similar offer or program, are not eligible for the program. Apellis reserves the right to modify or terminate the program at any time without notice.

for related medical procedures and services; nor will the free product be sold, traded, or distributed for sale. I will notify Apellis immediately if SYFOVRE is no longer medically necessary for this patient or if my patient's insurance status changes; (iv) I authorize Apellis to forward the

r	/	
<b>Physician Signature</b> (Dispense As Written) <b>Physician Signature</b> (If Drug Substitution Allowed)	Date (MM/DD/YYYY)	
This form cannot be processed without the physician's signature (no stamps).		

above prescription to the applicable pharmacy as allowed under applicable law.

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SYFOVRE (pegcetacoplan injection)

*Patient First Name:	Middle Initial:	_*Last Name:	

Please read the following authorizations carefully, and if you agree, provide your signature on page 1 of this form. You may keep a copy of this form for your records.

\*Section 10. Patient Authorizations

### Section 10.1 Authorization to Share Personal Health Information

l authorize my healthcare team and staff, my pharmacies, and my insurance ("Health Care Providers and Insurers") to use and to share my personal health information, including information relating to my medical condition, treatment, care management, health insurance, and all information provided on any prescription form for SYFOVRE ("My Information") to Apellis Pharmaceuticals, Inc. and its affiliates, vendors, and other agents (collectively, "Apellis") for the purposes of receiving services from ApellisAssist ("Patient Support Program"), which include but are not limited to:

- receiving product support and resources from Apellis, including insurance verification, product coverage, and financial assistance;
- disease and medication-related educational resources and communications, including disease state education and information about the medication by an Apellis Care Educator;
- and communications with me and my Health Care Providers and Insurers about my medical condition, treatment, care management, and health insurance

I also authorize Apellis Assist to share my information with my caregiver, if I have selected that option in this form.

I further authorize Apellis and its agents to de-identify my health information and use it in performing research, education, business analytics, and marketing studies, or for other commercial purposes, including linkage with other de-identified information Apellis may receive from other sources.

Once My Information has been shared with Apellis, I understand that it is outside of the control of my Health Care Providers and Insurers, and that the recipient may share this information with others and may not be required to comply with federal privacy laws or otherwise protect the information. However, I also understand that Apellis will protect My Information by sharing it only for the purposes for which I have provided permission. I understand and agree that if my SYFOVRE is received through a specialty pharmacy, that specialty pharmacy may receive payment from Apellis in exchange for giving My Information to Apellis. I understand that I do not have to sign this Authorization. A decision by me not to sign this Authorization will not affect my ability to receive health insurance benefits or my ability to get my medications or medical advice and treatment from my physician.

However, if I do not sign this Authorization, I understand I will not be able to participate and receive services from the Patient Support Program. I understand that this Authorization expires the earlier of (1) 10 years from the date signed below, (2) 1 year after the date of my last prescription, or (3) as may be required by applicable state law.

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SYFOVRE. (pegcetacoplan injection)

*Patient First Name:	Middle Initial:	_*Last Name:

## \*Section 10. Patient Authorizations (continued)

Imay change my mind and cancel this Authorization at any time by calling 888-APELLIS (888-273-5547), by notifying Apellis in writing at Attn: Privacy Office, Apellis Pharmaceuticals, Inc., 100 5th Avenue, Waltham, MA, 02451, or by emailing **privacy@apellis.com**. Cancellation of this Authorization will end further uses and sharing of My Information with Apellis and my participation in the Patient Support Program, but will not affect any uses or sharing of My Information based on this Authorization before cancellation. I understand I may request a signed copy of this Authorization.

## Section 10.2 Authorization to Enroll in Apellis Assist Patient Support Program

I authorize Apellis to collect My Information from me, my caregivers, and my Health Care Providers and Insurers, and to use and disclose My Information to provide product support and resources, including enrollment in the Patient Support Program. The Patient Support Program resources include, but are not limited to, providing:

- i) reimbursement and financial assistance information and
- ii) disease and medication-related educational resources and communications, including education provided by an Apellis Care Educator including but not limited to Geographic Atrophy ("Patient Resources"), if approved by prescribing physician.

I also authorize Apellis to communicate with me and/or my caregivers by mail, phone, email and/or text message for the Patient Support Program to receive education. I authorize Apellis to provide me and/or my caregivers with appropriate education on my disease state and medication by an Apellis Care Educator, and to provide me and/or my caregivers with helpful information and resources about SYFOVRE and Geographic Atrophy.

I understand that this education does not include medical advice and it does not replace or substitute the medical treatment and care I receive by my doctor. I further certify that I have discussed this with my doctor, and my doctor informed me of the potential risks and side effects associated with SYFOVRE and how to manage them if they occur. By signing below, I certify that the information contained in this form is complete and accurate to the best of my knowledge.

I authorize Apellis to send text messages to the phone number(s) I provide. I understand this consent is not a condition of participating in Apellis Assist or purchasing anything from Apellis. I may revoke this authorization and choose not to receive automated calls and text messages by replying STOP to any such text from Apellis or by contacting Apellis in writing at the address in section 10.1.

For support via the SYFOVRE Co-pay Program (if applicable), I certify that I am not a beneficiary of a federal or state healthcare program, including but not limited to Medicaid, Medicare, VA, DoD, TRICARE, or any state pharmaceutical assistance programs. I understand that once enrolled, Apellis will pay my eligible co-pay

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\*Patient First Name: \_\_\_\_\_\*Last Name: \_\_\_\_\_\*

## \*Section 10. Patient Authorizations (continued)

and/or co-insurance costs up to the program maximum, but that any costs over the program maximum or those that are not eligible for payment under the SYFOVRE Co-pay Program are my responsibility.

For support via the Patient Assistance Program (if applicable), I authorize Apellis to use my demographic information to access reports on my individual credit history from consumer reporting agencies. I understand that upon request, Apellis will tell me whether an individual consumer report was requested and the name and address of the agency that furnished it. I further understand and authorize Apellis to use any consumer reports about me and information collected from me, along with other information they obtain from public and other sources to estimate my income in conjunction with the patient assistance program eligibility determination process, if applicable. I certify that I will not submit a claim for reimbursement for any free product I receive from Apellis to any payer, including Medicare and Medicaid; and that no free product may be sold, traded, or distributed for sale. By signing, I verify that the information on this application and other supporting documentation is complete and accurate. I also verify that unless I have identified otherwise in this application, I have no other coverage for prescription medications, including Medicaid, Medicare or any public or private assistance programs, or any other form of insurance. If my insurance coverage should change, I will notify Apellis Assist immediately.

## Section 10.3 Authorization to Receive Marketing Communications (optional)

I authorize Apellis to communicate with me (by mail, phone, text and/or email) for marketing purposes or to otherwise provide me with information about Apellis products, services, and programs or other topics of interest, and to conduct market research or otherwise ask me about my experience with or thoughts about such topics. I understand and agree that any information I provide may be used by Apellis to help develop new products, services, and programs. I understand that I do not need to provide this authorization to receive marketing communications to participate in the Patient Support Program through ApellisAssist. I understand that this authorization will be in effect until such time as I opt-out of communications from Apellis.

I understand that I may revoke the Authorizations and choose not to receive information from Apellis by clicking the "unsubscribe" link provided in emails I receive from Apellis, calling Apellis at 888-APELLIS (888-273-5547), mailing a letter to Attn: Privacy Office, Apellis Pharmaceuticals, Inc., 100 5th Avenue, Waltham, MA, 02451, or emailing privacy@apellis.com.

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## **Indication and Important Safety Information**

### INDICATION

SYFOVRE® (pegcetacoplan injection) is indicated for the treatment of geographic atrophy (GA) secondary to age-related macular degeneration (AMD).

### IMPORTANT SAFETY INFORMATION

#### **CONTRAINDICATIONS**

• SYFOVRE is contraindicated in patients with ocular or periocular infections, and in patients with active intraocular inflammation

#### WARNINGS AND PRECAUTIONS

### • Endophthalmitis and Retinal Detachments

o Intravitreal injections, including those with SYFOVRE, may be associated with endophthalmitis and retinal detachments. Proper aseptic injection technique must always be used when administering SYFOVRE to minimize the risk of endophthalmitis. Patients should be instructed to report any symptoms suggestive of endophthalmitis or retinal detachment without delay and should be managed appropriately.

### Neovascular AMD

o In clinical trials, use of SYFOVRE was associated with increased rates of neovascular (wet) AMD or choroidal neovascularization (12% when administered monthly, 7% when administered every other month and 3% in the control group) by Month 24. Patients receiving SYFOVRE should be monitored for signs of neovascular AMD. In case anti-Vascular Endothelial Growth Factor (anti-VEGF) is required, it should be given separately from SYFOVRE administration.

#### Intraocular Inflammation

o In clinical trials, use of SYFOVRE was associated with episodes of intraocular inflammation including: vitritis, vitreal cells, iridocyclitis, uveitis, anterior chamber cells, iritis, and anterior chamber flare. After inflammation resolves, patients may resume treatment with SYFOVRE.

### • Increased Intraocular Pressure

o Acute increase in IOP may occur within minutes of any intravitreal injection, including with SYFOVRE. Perfusion of the optic nerve head should be monitored following the injection and managed as needed.

### **ADVERSE REACTIONS**

 Most common adverse reactions (incidence ≥5%) are ocular discomfort, neovascular age-related macular degeneration, vitreous floaters, conjunctival hemorrhage.

Please see full Prescribing Information for more information.

