RETEVMO™ (selpercatinib) Enrollment Form

PP-RC-US-1342 12/2020 ©Lilly USA, LLC 2020. All rights reserved.

Please complete and fax this form to 1-877-427-4030

If you have any questions, please call the Lilly Oncology Support Center™ at 1-866-472-8663, Monday-Friday 8am-10pm ET

By enrolling in the RETEVMO™ Savings and Support Program, Patients may receive various forms of support and information to help access RETEVMO™, which may include the following:

- Benefits Investigation Support
- Copay Savings and Other Financial Support
- Field Reimbursement Support
- Interim Access Support
- Ongoing Support

In order to process the requested services, we require 2 Patient signatures and 1 Prescriber signature. If Interim Access Support is being requested, we will require another Patient signature and another Prescriber signature accepting the Interim

Access Program Terms and Conditions. Not signing this form will result in an incomplete submission and a delay in requested services. **Patient Enrollment Checklist: Prescriber Enrollment Checklist:** Page 2 Page 4 ☐ Complete all sections in the Prescriber ☐ Complete all sections in the Patient **Enrollment section Enrollment section** ☐ Complete the prescription section, including: primary ☐ Document insurance information or provide copies diagnosis, dosing, and number of refills of your insurance and prescription card(s) ☐ If Interim Access Support is requested, please fill out ☐ Select optional Lilly Oncology Support Center[™] services that you would like to receive the Interim Access Support Prescription Section on Page 5 and sign Be sure to sign and date where "Signature of Patient or Authorized Representative" is located ☐ Select appropriate Benefits Investigation Support Option If selecting Specialty Pharmacy Conducted Benefits Page 3 Investigation, indicate which Specialty Pharmacy the prescription should be sent to Read and sign Patient HIPAA Authorization Confirm the Patient tested positive for a RET Alteration Page 5 (only if requesting RETEVMO™ Interim Access Support) Be sure to sign and date where "Signature of Manually sign and date the form Patient or Authorized Representative" is located Page 5 (only if requesting RETEVMO™ Interim Access Support) Pages 6-8 ☐ Complete the prescription section, including: primary ☐ Read and acknowledge the Consent, Terms and diagnosis and dosing Conditions, and Privacy Notice on remaining pages Confirm the Patient tested positive for a RET Alteration Manually sign and date the form



Complete and fax this form to 1-877-427-4030

PATIENT ENROLLMENT SECTION RETEVMO™ (selpercatinib)

OFFICE: Please fax to 1-877-427-4030

PP-RC-US-1342 12/2020 ©Lilly USA, LLC 2020. All rights reserved.

Patient: Fill out only the Patient section and sign where indicated

participation in the program, please contact us at 1-866-472-8663.

Authorized Representative: Fill out both the Patient section and the Authorized Representative section and sign where indicated

Adduses		DOB (MM/DD/YYYY)
Address	City	State ZIP Code
US or Puerto Rico Resident ☐ Yes ☐ No	Gender □ M □ F Preferred I	_anguage ☐ English ☐ Spanish ☐ Other
*By providing my telephone number and signir Company. I understand that I am not required	ng this form, I agree to receive automa to provide my number as a condition of	ted marketing calls and texts from and on behalf of Eli Lilly a of purchase. Message and data rates may apply.
Authorized Representative should fill out the section b	elow and sign on behalf of the Pedia	tric Patient 12 years of age or older
	·	DOB (MM/DD/YYYY)
Relationship to Patient		
Address Ves No	City	State Zip
		anguage □ English □ Spanish □ Other
		ed marketing calls and texts from and on behalf of Eli Lilly a
Company. I understand that I am not required to	to provide my number as a condition o	f purchase. Message and data rates may apply.
select one of the following: No Insurance Coverage hary Prescription Insurance Company	•	ard (Front and Back) Is Attached
rance Company Phone #		
	Cardholder Name _	
cy/ID	Cardholder Name _ Group #	
cy/ID	Cardholder Name _ Group # PCN	
sy/ID	Cardholder Name _ Group # PCN nmercial coverage plan and is active	
sy/ID	Cardholder Name Group # PCN nmercial coverage plan and is active ur prescriptions? Examples include N	ledicaid, Medicare, Medicare Part D, and others.
BIN	Cardholder Name Group # PCN nmercial coverage plan and is active ur prescriptions? Examples include Name hecking the corresponding checkbox	ledicaid, Medicare, Medicare Part D, and others. es below.
sy/ID	Cardholder Name Group # PCN nmercial coverage plan and is active ir prescriptions? Examples include Note the Card Terms and Conditions	ledicaid, Medicare, Medicare Part D, and others. es below.
ey/ID	Cardholder Name Group # PCN nmercial coverage plan and is active ar prescriptions? Examples include Note the Saving Card Terms and Conditions a statements in order to be eligible)	es below. on page 6
SIN	Cardholder Name Group # PCN	es below. on page 6
SIN	Cardholder Name Group # PCN	ledicaid, Medicare, Medicare Part D, and others. es below. on page 6 e or older les include Medicaid, Medicare, Medicare Part D, and others

Lilly

boxes above, I consent to my enrollment in the additional Lilly Oncology Support Center™ services as described in the Consent on page 7. To cancel your

PATIENT HIPAA AUTHORIZATION

OFFICE: Please fax to 1-877-427-4030

PP-RC-US-1342 12/2020 ©Lilly USA, LLC 2020. All rights reserved.

Before the Lilly Oncology Support Center[™] can start helping you, Lilly may ask for some information about you and your health from your Health Care Entities (as defined below). This is known as your *Protected Health Information*, or *PHI*. By signing this form, you understand and agree that your PHI may be shared with or used by Lilly as explained below.

PHI includes information like:

- Your health insurance or benefits, including how much coverage you have
- All records about your treatment
- Whether you're staying on your medicine or treatment

If you agree, your PHI may be shared by these entities (together "Health Care Entities"):

- Your doctors and other healthcare providers
- Your healthcare plan or health insurance company
- Clearinghouses or other agents
- Your pharmacv
- Others who might have your PHI on behalf of your healthcare providers, pharmacies and healthcare plans

Your PHI is used in ways like these:

- To learn how much of your Lilly treatment is covered by your insurance
- To help you find other ways to afford your treatment
- To track your use of your Lilly treatment
- To share information with your healthcare provider
- To make sure that you receive high-quality services from the program
- To measure program performance and make program improvements
- Internal Lilly use of data to drive business decisions and metrics on hub performance
- Reports to our sales force regarding HCP use of hub services
- Conversations/messages to your HCP regarding trends and hub performance

Other things you should know about sharing and using your PHI:

- We only ask for and share the PHI that we need to provide the benefits you want. We do not ask for any PHI that we do not need, but we may receive some in the health records sent to us. Your PHI will be released to Eli Lilly and Company and Lilly USA, LLC and its affiliates, agents, representatives, and service providers (together "Lilly")
- You don't have to give permission to share your PHI with Lilly to receive treatment from your healthcare providers, your prescription from your pharmacy, or benefits from your healthcare plan, but the Lilly Oncology Support Center™ may not be able to help you without it
- After your PHI has been shared, it may no longer be covered by federal and state privacy laws (such as HIPAA), and it may be shared again with others by Lilly
- Your signed permission to share and use your PHI lasts for 3 years from the date of your signature unless you are a resident of Maryland, Maine, or Montana, in which case the permission will last for 1 year from the date of your signature. In either case, you may revoke your permission before then by writing to PO Box 12307, La Jolla, CA 92039, which will preclude reliance on the authorization after the date your written revocation is received
- Your healthcare providers (such as pharmacies) may be paid by us in exchange for sharing your PHI. They may also be paid by us to use your PHI to provide services, such as contacting you about Lilly products
- You can stop sharing your PHI with us or change what you share by calling us at 1-866-472-8663, or by writing us at PO Box 12307, La Jolla, CA 92039
- Your cancellation or revocation of this Authorization will be effective when your Health Care Entities receive notice
 of your cancellation or revocation, and will not apply to any information shared with Lilly by your Health Care
 Entities prior to the time those Health Care Entities receive notice

I have read and agree to the Patient HIPAA Authorization. By signing this Authorization, I represent that I am the Authorized Representative for the Pediatric Patient 12 years of age or older. I understand I am entitled to a copy of this signed Authorization.



Signature of Patient or Authorized Representative	_ Date Signed (MM/DD/YYYY)
Printed Name of Patient/Authorized Representative	_ Date of Birth (MM/DD/YYYY)
Not signing this form will result in an incomplete submission and a delay in requested services	



PRESCRIBER ENROLLMENT SECTION RETEVMO™ (selpercatinib)

OFFICE: Please fax to 1-877-427-4030

PP-RC-US-1342 12/2020 ©Lilly USA, LLC 2020. All rights reserved.

	Address	City	State Zip	
	Group Tax ID Office Contact N	Name Office Cont	act Phone	
	Office Contact Email	NPI #		
ă.	Patient Name (First, MI, Last)	DOB	(MM/DD/YYYY)	
3,	Address	City State	e Zip _	
	RETEVMO™ Prescription - Fill out corresponding p	rescription below and sign at the bottom of	f page	
	Diagnosis:	RETEVMO™ Prescribing Information (PI) Dosing:	Days Supply	Refill
	Primary Diagnosis: Adult patients with metastatic RET fusion-positive nonsmall cell lung cancer (NSCLC)	☐ 160mg (2 x 80mg capsules) orally twice daily Recommended for patients with body weight 50kg or greater	Please Do Not Change As This Is A Set Value	
must ct a nosis	Primary Diagnosis: Adult and pediatric patients 12 years of age and older with advanced or metastatic RET-mutant medullary thyroid cancer (MTC) who require temic therapy	☐ 120mg (3 x 40mg capsules) orally twice daily Recommended for patients with body weight less than 50kg ☐ 80mg (80mg capsule) orally twice daily	Days Supply	Refil
	Primary Diagnosis: Adult and pediatric patients 12 years of age and older with advanced or metastatic <i>RET</i> fusion-positive thyroid cancer who require systemic therapy and who are radioactive iodine-refractory (if radioactive iodine is appropriate)	☐ 40mg (40mg capsule) orally twice daily ☐ 40mg (40mg capsule) orally once daily	30	
Ronofi	ts Investigation Support (select are choice)			
optical selection of the control of	ts Investigation Support (select one choice) illy Conducted Benefits Investigation—The Lilly Oncology Supports to help identify the lowest out-of-pocket cost available for RETE'sts. A Lilly Oncology Support Center™ representative will help triage PRESCRIPTION SECTION ABOVE. Specialty Pharmacy Conducted Benefits Investigation—Special Properties of the Properties of th	VMO™ and will forward the prescription to the Specialts and troubleshoot access issues on the Patient's behavity Pharmacy where prescription was sent	y Pharmacy that the lf. IF CHECKED, MU /MO™ at no charge coverage. Not availa erage delay, the Pat uct for the RETEVM bligation accompani	Patient JST FIL for able to tient, O™ Inte
optic select OUT Selec	illy Conducted Benefits Investigation—The Lilly Oncology Supports to help identify the lowest out-of-pocket cost available for RETE'ts. A Lilly Oncology Support Center™ representative will help triage PRESCRIPTION SECTION ABOVE. Specialty Pharmacy Conducted Benefits Investigation—Special Presentative Pharmacy Conducted Benefits Investigation—Special Presentative Pharmacy Conducted Benefits Investigation—Special Presentative Pharmacy Conducted Benefits Investigation—Special Presentation and experients who are new to RETEVMO™ and experients whose insurers have made a final determination to deny contents whose insurers have made a final determination to deny contents whose insurers have made a final determination to deny contents whose insurers have made a final determination to deny contents are propried to the propried of the proprie	MO™ and will forward the prescription to the Specialt e and troubleshoot access issues on the Patient's behavior that the prescription was sent	y Pharmacy that the If. IF CHECKED, MI IMO™ at no charge coverage. Not availaterage delay, the Patuct for the RETEVM Diligation accompanichange or end the pi Lilly and Company, Lilly Usted to, and has directed made through the duration of orm, the prescription complunderstand that by signir recognized Compendia. P	Patient JST FIL for able to tient, O™ Inte es program SA, LLC, t hy disclose the Patien blies with ng this for



RETEVMO™ INTERIM ACCESS PROGRAM REQUIREMENTS

OFFICE: Please fax to 1-877-427-4030

PP-RC-US-1342 12/2020 ©Lilly USA, LLC 2020. All rights reserved.

RETEVMO™ Interim Access Program Requirements

To be eligible for the RETEVMO™ Interim Access Program, a Patient must: 1) be a new RETEVMO™ Patient; 2) be prescribed RETEVMO™ for an FDA-approved indication or an indication medically supported by CMS recognized Compendia; 3) be 18 years of age or older or an authorized representative of a Patient under age 18; and 4) be a resident of the United States or Puerto Rico

Please note: This program is provided by Sonexus rather than your in-office dispensary or any other specialty pharmacy your Patient may later use. We will contact your office as soon as the Patient has received his/her dose so you can begin the process of starting his/her next month's script.

Terms and Conditions:

The RETEVMO™ Interim Access Program (or "Program") provides a 15-day supply of RETEVMO™ at no charge for eligible, insured patients who are:

- 1) prescribed RETEVMO™ for the first time after testing positive for a RET alteration,
- 2) experiencing a minimum 5-business-day delay in insurance coverage determination,

I confirm the Patient tested positive for a RET Alteration

- 3) prescribed RETEVMO™ for an FDA-approved indication or an indication medically supported by CMS-recognized compendia, and
- 4) enrolled in the Lilly Oncology Support Center™

5) residents of the United States or Puerto Rico. May not be combined with any other offer. Not available to patients whose insurers have made a final determination to deny the patient coverage for RETEVMOTM. If a denial is received after the initial 5 business days have passed and appeal rights are being pursued, or if there is a persistent coverage delay, the patient, under appropriate circumstances, may be eligible for up to 3 additional 15-day supplies of RETEVMOTM. Product provided through the Program is only available through the Program non-commercial specialty pharmacy. Product is provided free of charge and may not be sold, bartered, or returned for credit. Reimbursement cannot be sought from any third party for product provided under the program. Patients enrolled in Medicare Part D are prohibited from counting any portion of the cost of the product provided under the Program towards true out-of-pocket ("TrOOP") costs for prescription drug calculations. No purchase contingency or other obligation accompanies program participation. This Program is not health insurance and does not guarantee coverage. Lilly reserves the right to change or end the program at any time. Benefits under the program are not transferable.

l certif Repres	I the program at any time. Benefits under the program are not traffy and agree to the above RETEVMO™ Interim Access Terms sentative for the Pediatric Patient 12 years of age or older. gnature of Patient or Authorized Representative ot signing this form will result in an incomplete submission	and Conditions. By signing this Authorization, I re	epresent that I am t		
4	,	· ·	DOB (MM/DD/YYYY)		
ent	Address	City Sta	te Zip _		
	RETEVMO™ Prescription - Fill out corresponding	-			
	Diagnosis:	RETEVMO™ Prescribing Information (PI) Dosing:	Days Supply	Refill	
	Primary Diagnosis: Adult patients with metastatic RET fusion-positive non-small cell lung cancer (NSCLC)	☐ 160mg (2 x 80mg capsules) orally twice daily Recommended for patients with body weight 50kg or greater	Please Do Not Change As This Is A Set Value		
nust et a nosis	Primary Diagnosis: Adult and pediatric patients 12 years of age and older with advanced or metastatic RET-mutant medullary thyroid cancer (MTC) who require temic therapy	☐ 120mg (3 x 40mg capsules) orally twice daily Recommended for patients with body weight less than 50kg ☐ 80mg (80mg capsule) orally twice daily		Refills	
	Primary Diagnosis: Adult and pediatric patients 12 years of age and older with advanced or metastatic <i>RET</i> fusion-positive thyroid cancer who require systemic therapy and who are radioactive iodine-refractory (if radioactive iodine is appropriate)	☐ 40mg (40mg capsule) orally twice daily ☐ 40mg (40mg capsule) orally once daily	15		

Prescriber: I certify that I understand and agree: 1) To the terms and conditions of the RETEVMO™ Interim Access Program; 2) I am licensed to prescribe the medication identified in this form, and that the prescription complies with my state-specific prescribing requirements; 3) In my medical judgment, RETEVMO™ is clinically appropriate for the Patient named above and its use is consistent with the FDA-approved indication or an indication medically supported by CMS recognized Compendia; and 4) This supply of RETEVMO™ is specifically for the Patient named above.

FRESCRIBER SIGNATURE: FRESCRIBER MUST	I WANDALLI SIGIN AND DATE. NUDD	er starrips, signature by other orn	ice personnerior the Frescriber, a	and computer-generated signature:	s will flot be accepted.



Dispense as written

Not signing this form will result in an incomplete submission and a delay in requested services

Date Signed (MM/DD/YYYY)



SAVINGS CARD TERMS AND CONDITIONS

PP-RC-US-1342 12/2020 ©Lilly USA, LLC 2020. All rights reserved.

Terms and Conditions:

By using the Retevmo Savings ("Card"), you attest that you meet the eligibility criteria, agree to and will comply with the terms and conditions described below:

Offer good for up to 12 months until 12/31/2021. Patients must have coverage for Retevmo through their commercial drug insurance coverage to pay as little as \$0 for a 30-day supply of Retevmo. Offer subject to a monthly cap of wholesale acquisition cost plus usual and customary pharmacy charges and a separate annual cap of \$25,000. Participation in the program requires a valid patient HIPAA authorization. Offer void where prohibited by law. Patient is responsible for any applicable taxes, fees, or amounts exceeding monthly or annual caps. This offer is invalid for patients without commercial drug insurance or whose prescription claims for Retevmo are eligible to be reimbursed, in whole or in part, by any governmental program, including, without limitation, Medicaid, Medicare, Medicare Part D, Medigap, DoD, VA, TRICARE®/CHAMPUS, or any state patient or pharmaceutical assistance program. This offer is not valid for: Massachusetts residents if an AB-rated generic equivalent is available; California residents if an FDA-approved therapeutic equivalent is available. Available only in the US and Puerto Rico for residents of the US and Puerto Rico. By accepting this offer, you agree that if you are required to do so under the terms of your insurance coverage for this prescription or are otherwise required to do so by law, you should notify your insurance carrier of your redemption of this Card. This offer cannot be combined or utilized with any other program, discount, discount card, cash discount card, coupon, incentive, or similar offer involving Retevmo. It is prohibited for any person to sell, purchase or trade; or to offer to sell, purchase or trade, or to counterfeit this Card. This offer may be terminated, rescinded, revoked or amended by Lilly USA, LLC at any time without notice. Card activation required. This Card is not health insurance. This Card expires on 12/31/2021.



PP-RC-US-1342 12/2020 ©Lilly USA, LLC 2020. All rights reserved.

What to Know About the RETEVMO™ Ongoing Support Program:

Your healthcare provider has talked with you about using RETEVMOTM, an Eli Lilly and Company medicine. The Lilly Oncology Support CenterTM was created to help you have a positive experience as you get started with and use this medicine. The Lilly Oncology Support CenterTM offers personalized support to Patients at no charge.

OPTIONAL RETEVMO™ ONGOING SUPPORT ENROLLMENT CONSENT

Ongoing Support Enrollment Consent:

The Ongoing Support Services included in the Lilly Oncology Support Center™ provide support after you've received your medication, like check-in calls to answer any questions you might have about RETEVMO™. As part of your participation in the Ongoing Support Services, Eli Lilly and Company and Lilly USA, LLC and its affiliates, agents, representatives, and service providers (together "Lilly") may use, disclose, and/or transfer the personal information you supply to provide services related to your condition and treatment to administer the program.

Services include:

Contacting you by email, mail or telephone to provide personalized services, delivered by your Lilly Oncology Support Center[™] team, such as information and marketing materials; responding to customer service requests and/or questions about your treatment; requesting feedback on your experience with the related products, services, and programs, including market research and medical research; disclosing your enrollment and use of these services to your doctors and insurers; analyzing and/or measuring program performance and program effectiveness for future enhancements; and other activities related to your condition and therapy that are not part of the Lilly Oncology Support Center[™]. These activities include opportunities to share your story and participate in studies about products and services. To cancel your participation in the program, please contact us at **1-866-472-8663** Mon-Fri, 8am–10pm ET.



PP-RC-US-1342 12/2020 ©Lilly USA, LLC 2020. All rights reserved.

Privacy Notice:

We may use and save your personal information to meet legal or regulatory obligations that are in the legitimate interest of Lilly, to fulfill legitimate and lawful business purposes in accordance with Lilly's record retention policies and applicable laws and regulations, and to respond to lawful requests by public authorities, including to comply with national security or law enforcement requests.

Your information may be combined with other information that you have previously provided or that Lilly has received. We do not sell personal information.

We may transmit personal information about you to other Lilly affiliates worldwide. These affiliates may in turn transmit personal information about you to other Lilly affiliates. Some of Lilly's affiliates may be located in countries that do not ensure the same level of data protection. Nevertheless, all of Lilly's affiliates are required to treat personal information in a manner consistent with this notice. To obtain additional information about Lilly's privacy practices, including the basis for transfers and safeguards that Lilly has in place for cross-border transfers of personal information, please contact us at privacy@lilly.com or visit https://www.lilly.com/privacy.

We provide reasonable physical, electronic and procedural safeguards to protect information we work with and maintain. We limit access to your information to authorized employees, agents, contractors, vendors, subsidiaries, and business partners, or others who need such access to information to carry out their assigned roles and responsibilities on behalf of Lilly. Please be aware, although we try to protect the information we work with and maintain, no security system can prevent all potential security breaches.

Upon verification, you have the right to request information from us regarding how your personal information is being used and with whom that information is being shared. You also have the right to request to see and get a copy of the personal information that we have about you, request its correction or request its erasure/deletion.

There may be exceptions that apply to your request.

In limited circumstances, you may have the right to have your information transmitted to another entity or person in a machine-readable format.

You will not be discriminated against for exercising any of your rights.

To exercise your rights, you or your authorized representative may submit a request by contacting us using one of the methods listed below.

You may make any of the above requests by contacting us at: The Lilly Answers Center, Lilly USA, LLC, Lilly Corporate Center, Indianapolis, IN 46285 or by calling 1-800-545-5979.

If you wish to raise a complaint on how we have handled your personal information, you can contact the Global Privacy Office and Data Protection Officer at privacy@lilly.com who will investigate the matter.

If you are not satisfied with our response or have any concerns about how your data is being processed, you can register a complaint with a relevant regulatory authority (e.g. a Data Protection Authority (DPA) or Attorney General).

