

SUBMIT ONLY REQUESTED DOCUMENTS

Required field (\*) M-US-00006694(v3.0)

Services Requested (Check all that apply)
[ ] Benefits Investigation/Prior Authorization [ ] Refer Patient to Co-pay Assistance [ ] Appeals Support

Step 1 Patient Information

\*First name: \*Last name:
\*Date of birth (MM/DD/YYYY): Gender: [ ] Male [ ] Female
Street: Apt:
City: \*State: ZIP:
Home phone: Cell phone: [ ] Do not contact patient
Preferred language: [ ] English [ ] Spanish [ ] Other: Has patient started therapy? [ ] Yes [ ] No
Alternate contact name: Relationship: Alt. phone:

Step 2 Insurance Information

Is the patient insured? [ ] Yes [ ] No
If patient is uninsured, please complete the Genentech Patient Foundation Enrollment Form or call (888) 941-3331 for assistance.
If insured, please fill out the information below or attach a copy of the patient's insurance cards.

Table with 2 columns: Primary Insurance, Secondary Insurance. Rows include Insurance name, Subscriber name, Subscriber/Policy ID #, Group #, Insurance phone.

Step 3 Patient's Therapy (Check all that apply)

[ ] LUCENTIS (ranibizumab injection) [ ] SUSVIMO (ranibizumab injection) [ ] VABYSMO (faricimab-svoa)
[ ] Initial Fill and Implant Procedure [ ] Refill Exchange Procedure

Step 4 Diagnosis and Clinical Information

Please provide the appropriate diagnosis code(s) to the highest level of specificity. For coding information, please visit Genentech-Access.com/Ophthalmology.
Anticipated date of treatment: \*Diagnosis code(s):

Step 5 Prescriber Information

\*First name: \*Last name:
\*Practice name:
\*Street: Suite: \*City:
\*State: \*ZIP: Prescriber tax ID #:
Prescriber NPI #: Group NPI #:
Office contact: Contact phone: Contact fax:

Step 6 Administration Information (Complete for SUSVIMO only)

[ ] Ambulatory Surgical Center [ ] Hospital Outpatient Department
Place of administration name: Tax ID #:
Street: Suite: City:
State: ZIP: NPI #:

Step 7 Health Care Provider Certification

By submitting this form, I certify: (a) The above therapy is medically necessary for this patient and the treatment decision has been made by the prescribing physician. (b) If the indication for which this Genentech product is being prescribed to treat is not listed in the FDA-approved label, the prescriber is prescribing the medication for an "unapproved" use... (f) No action on these services will be taken until the patient consent document has been received.