

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: _____
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.

Primary Insurance

Secondary Insurance

Insurance name		
Subscriber name (if not patient)		
Subscriber/Policy ID #		
Group #		
Insurance phone		

Step 3

Diagnosis and Clinical Information

*To the highest level of specificity, provide diagnosis code:

☐ E84.0 Cystic fibrosis with pulmonary manifestations ☐ E84.8 Cystic fibrosis with other manifestations
☐ E84.9 Cystic fibrosis, unspecified ☐ Other code: _____

*Prescription type: ☐ New start ☐ Continuing therapy ☐ Restart therapy Has patient started therapy? ☐ Yes ☐ No

Anticipated date of treatment: ____ / ____ / ____

Step 4

Prescription Information

Pulmozyme regimen 2.5 mL (dornase alfa) inhalation solution

Dispense: ☐ 30-day supply ☐ 60-day supply ☐ 90-day supply Refill _____ times SIG: ☐ QD ☐ BID

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

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, meaning that the FDA has not approved the efficacy, dosage amount or safety of this medication for such a use. (c) The provider's office received the authorization to release the information above and other protected health information (as defined by the Health Insurance Portability and Accountability Act of 1996 [HIPAA]) to Genentech, Inc., Genentech Access Solutions, the contracted dispensing pharmacy, or other contractors for the purpose of requesting reimbursement support, assisting in initiating or continuing therapy, as a break in treatment would negatively impact the patient's therapeutic outcome. (d) The provider's office will not attempt to seek reimbursement for free product provided to the patient. (e) The services requested on behalf of the patient may include benefits investigation (BI), prior authorization (PA) and appeals support, co-pay card and co-pay assistance foundation referral. (f) No action on these services will be taken until the patient consent document has been received.

[†]National Provider Identifier.

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