

Tezspire

Patient Inf	ormation:			
Name:				
Member ID	:			
Address:				
City, State,	Zip:			
Date of Bir				
	,			
Prescriber	Information:			
Name:				
NPI:				
Phone Nur	nber:			
Fax Number	er			
Address:				
City, State,	Zip:			
_				
•	l Medication			
Rx Name:	1			
Rx Strengt				
Rx Quantit	•			
Rx Freque	-			
Rx Route o				
Administration: Diagnosis and ICD Code:				
Diagnosis	and ICD Code:			
prescribed a quantities can Upon receipt SECTION requests. medication	medication for your h be provided. Plea t of the completed A: Please no Pharmacy pri ns that are no	efit requires that we review certain requests for coverage with the proposition patient that requires Prior Authorization before benefit coverage or consecomplete the following questions then fax this form to the toll-free not form, prescription benefit coverage will be determined based or the that supporting clinical documentation is required or authorization reviews can be subject to trial with a listed within the criteria. The policies are subject to the trial with a listed within the criteria. The policies are subject to the trial with a listed within the criteria. The policies are subject to the trial with a listed within the criteria.	verage of a umber liste of the plan for AL addition ochange	additional ed below. n's rules. LPA al
1 I	s the request an II	NITIAL or CONTINUATION of therapy?		
	·	• •		
L	Initial (If checked	1, go to 7)		
	Continuation (If c	checked, go to 2)		
		ently receiving the request medication and taking it with one oid OR one inhaled corticosteroid-containing combination	Yes	No

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	[If no, skip to question 7.]		
3	I I a the contract to the cont		
	Has the patient been receiving medication samples of the requested medication?	Yes	No
	[If yes, skip to question 7.]		
	Does the patient have a previously approved prior authorization (PA) on file with the current plan?	Yes	No
	[Note: If the patient does NOT have a previously approved PA on file for the requested medication with the current plan, the renewal request will be considered under initial therapy.]		
	[If no, skip to questions 7.]		
	Will the patient be concurrently receiving the requested medication in combination with any anti-IgE, anti-IL4, OR anti-IL5 monoclonal antibody agents (benralizumab, omalizumab, mepolizumab, reslizumab, dupilumab, et cetra)?	Yes	No
	[If yes, no further questions.]		
	Has documentation been submitted to confirm that the patient has responded to therapy as determined by the prescriber? ACTION REQUIRED: Submit supporting documentation.	Yes	No
	[Note: Examples of a response to Tezspire therapy are improvement in FEV1 from baseline, decreased asthma exacerbations; decreased asthma symptoms; decreased hospitalizations, emergency department/urgent care, or medical clinic visits due to asthma; improved lung function parameters; and/or a decreased requirement for oral corticosteroid therapy.]		
	[No further questions.]		
7	What is the indication?		
	[] Asthma (If checked, go to 8)		
	[] Other (If checked, no further questions)		
8	Is the patient 12 years of age or older?	Yes	No
	[If no, no further questions.]		
9	Is the requested medication being prescribed by or in consultation with an allergist,	Yes	No

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	immunologist, or pulmonologist?		
	[If no, no further questions.]		
10	Does the patient have asthma that is uncontrolled or was uncontrolled at baseline as defined by the following: patient experienced two or more asthma exacerbations requiring treatment with systemic corticosteroids in the previous year?	Yes	No
	[Note: "Baseline" is defined as prior to receiving any Tezspire, anti-interleukin-5 therapies (that is, Cinqair, Fasenra, or Nucala), Dupixent, or Xolair.]		
	[If yes, skip to question 14.]		
11	Does the patient have asthma that is uncontrolled or was uncontrolled at baseline as defined by the following: patient experienced one or more asthma exacerbation(s) requiring hospitalization, an Emergency Department visit, or an urgent care visit in the previous year?	Yes	No
	[Note: "Baseline" is defined as prior to receiving any Tezspire, anti-interleukin-5 therapies (that is, Cinqair, Fasenra, or Nucala), Dupixent, or Xolair.]		
	[If yes, skip to question 14.]		
12	Does the patient have asthma that is uncontrolled or was uncontrolled at baseline as defined by the following: patient has a forced expiratory volume in 1 second (FEV1) LESS THAN 80% predicted?	Yes	No
	[Note: "Baseline" is defined as prior to receiving any Tezspire, anti-interleukin-5 therapies (that is, Cinqair, Fasenra, or Nucala), Dupixent, or Xolair.]		
	[If yes, skip to question 14.]		
13	Does the patient have asthma that is uncontrolled or was uncontrolled at baseline as defined by the following: patient has an FEV1/forced vital capacity (FVC) LESS THAN 0.80?	Yes	No
	[Note: "Baseline" is defined as prior to receiving any Tezspire, anti-interleukin-5 therapies (that is, Cinqair, Fasenra, or Nucala), Dupixent, or Xolair.]		
	[If no, no further questions.]		
14	Does the patient have asthma that is uncontrolled or was uncontrolled at baseline as defined by the following: the patient has asthma that worsens upon tapering of oral corticosteroid therapy?	Yes	No
	[Note: "Baseline" is defined as prior to receiving any Tezspire, anti- interleukin-5		
	therapies (that is, Cinqair, Fasenra, or Nucala), Dupixent, or Xolair.]		

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15	Will the requested medication be used in combination with anti-IgE, anti- IL4, or anti-IL5 monoclonal antibody agents (benralizumab, omalizumab, mepolizumab, reslizumab, dupilumab, et cetera)?	Yes	No
	[If yes, no further questions.]		
16	Will the requested medication be administered concurrently with live vaccines?	Yes	No
	[If yes, no further questions.]		
17	Will the requested medication be used for the relief of acute bronchospasm or status asthmaticus?	Yes	No
	[If yes, no further questions.]		
18	Do the patient and provider agree that the requested medication WILL NOT be used as monotherapy AND WILL be used as an add on maintenance treatment with an inhaled corticosteroid?	Yes	No
	[If no, no further questions.]		
19	Does the patient have an active or untreated helminth infection?	Yes	No
	[If yes, no further questions.]		
20	Does the patient have a documented intolerance, contraindication to, or failed treatment for AT LEAST 4 months of therapy with Xolair?	Yes	No
	[If no, no further questions.]		
21	Has documentation been submitted to confirm that the patient has had an intolerance, contraindication to, or failed treatment for at least 3 months with preferred IL-4 and IL13R inhibitor, Dupixent? ACTION REQUIRED: Submit supporting documentation.	Yes	No
	[If no, no further questions.]		
22	Has documentation been submitted to confirm that the patient has had an intolerance, contraindication to, or failed treatment for at least 3 months with IL-5 inhibitor, Nucala? ACTION REQUIRED: Submit supporting documentation.	Yes	No
	[If no, no further questions.]		
23	Does the requested dose exceed Food and Drug Administration (FDA) approved	Yes	No

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	label dosing for this indication?		
	[Note: Dosing is 210 mg once every 4 weeks]		
Dia		4 4 - 4hi	
Plea	ase document the diagnoses, symptoms, and/or any other information important	t to this r	eview:
SEC	TION B: <u>Physician Signature</u>		

FAX COMPLETED FORM TO: 1-833-896-0656

Disclaimer: An authorization is not a guarantee of payment. Member must be eligible at the time services are rendered. Services must be a covered Health Plan Benefit and medically necessary with prior authorization as per Plan policy and procedures.

PHYSICIAN SIGNATURE

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DATE