AETNA BETTER HEALTH® OF VIRGINIA REQUEST FORM

Tezspire® (tezepelumab-ekko)

Fax back to: 1-855-799-2553

If the following information is not complete, correct, or legible, the PA process can be delayed.

Please use one form per member.

MEMBER INFORMATION													
Last Name:	First Name:												
Medicaid ID Number:	Date of Birth:												
Weight in Kilograms:	_												
PRESCRIBER INFORMATION													
Last Name:	First Name:												
NPI Number:													
Phone Number:	Fax Number:												
DRUG INFORMATION													
Drug Name/Form:													
Strength:													
Dosing Frequency:													
Length of Therapy:													
Quantity per Day:													
The Virginia Department of Medical Assistance Service Cinqair®, Dupixent®, Fasenra®, Nucala®, Tezspire™ an and efficacy of theses combinations have NOT been e	d Xolair® to be experimental and investigational. Safety												

(Form continued on next page.)

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AETNA BETTER HEALTH® OF VIRGINIA REQUEST FORM: Tezspire® (tezepulumab-ekko)

Member's Last Name:											Member's First Name:													
DIAGNOSIS AND MEDICAL INFORMATION																								
Fo	r sev	ere* a	asth	ma i	nitia	ıl app	rova	al, co	mpl	ete 1	the f	ollo	owin	g qu	estio	ns to	rec	eive	a 6-n	nont	h ap	prov	al:	
1.		ne me Yes	mbe	er 12	yea] No		age (or ol	der?	ANI	D													
2.	Does the member have a diagnosis of severe* asthma? AND Yes No																							
3.	Will coadministration with another monoclonal antibody be avoided (e.g., omalizumab, mepolizumab, reslizumab, benralizumab, dupilumab)? AND Yes No																							
4.	Will this be used for add-on maintenance treatment in members regularly receiving both (unless otherwise contraindicated) of the following: • Medium- to high-dose inhaled corticosteroids; AND												wise											
	•	An a Yes	addi	tion	al co] No	ntrol	ler m	nedio	catio	n (e.	g., lc	ng	-acti	ng be	eta a	gonis	st, le	ukot	riene	e mod	difier	s)?		
5.	Has the member had two or more exacerbations in the previous year requiring oral or injectable corticosteroid treatment (in addition to the regular maintenance therapy defined above) or one exacerbation resulting in a hospitalization? AND Yes No																							
6.	•	Use Nur	of sof index	syste nhal r of l	mic ed co	corticortico ortico italiz ry vol	coste oster ation	eroid oids ns, El	ls R visi	its, o	or uns	sch	edul	ed vi							due t	o cor	nditic	n
	(For	m co	ntinu	ued (on ne	ext p	age.))																

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Member's Last Name:	Member's First Name:													
7. Has the member tried and failed an adequate trial of the 2 different preferred products (Fasenra® and Xolair®)?														
☐ Yes ☐ No ☐ N/A														
If N/A was selected for question 7 please answer the following:														
a. Does the member lack an eosinophilic phenotype with blood eosinophils ≥150 cells/µL? AND														
☐ Yes ☐ No														
b. Does the member lack a serum IgE level < 30 IU/mL? OR														
Yes No														
c. Does the member have another predicted intolerance the preferred agents? (Answer below)														
☐ Yes ☐ No														
 8. Has the member been assessed for toxicity? AND Yes No 9. Does the member have improvement in asthma symptoms or asthma exacerbations as evidenced by decrease in one or more of the following: Use of systemic corticosteroids Hospitalizations ER visits Unscheduled visits to healthcare provider Improvement from baseline in forced expiratory volume in 1 second (FEV₁)? Yes No 														
*Components of severity for classifying asthma as so	evere may include any of the following (not all-inclusive):													
Symptoms throughout the day														
 Nighttime awakenings, often 7 times/week SABA use for symptom control occurs several times per 	r dov													
 SABA use for symptom control occurs several times per Extremely limited normal activities 	luay													
■ Lung function (percent predicted FEV ₁) < 60%														
	re generally more frequent and intense relative to moderate													
asthma														

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Member's Last Name:											N	Member's First Name:											
Pre	Prescriber Signature (Required)															D	ate						
•	_	iture ifiabl	•						ıe ab	ove i	nfor	mati	on is	accur	ate								
Plea	ase ii	nclud	de A	LL re	eque	sted	info	rma	tion;	Inco	mpl	ete f	orms	will c	delay	the I	PA pr	roces	s.				

Submission of documentation does NOT guarantee coverage.

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