

Updated: 04/2024 DMMA Approved: 04/2024

Request for Prior Authorization for Qbrexza (glycopyrronium) Website Form – www.highmarkhealthoptions.com Submit request via: Fax - 1-855-476-4158

All requests for Qbrexza (glycopyrronium) require a Prior Authorization and will be screened for medical necessity and appropriateness using the criteria listed below.

Qbrexza (glycopyrronium) Prior Authorization Criteria:

Coverage may be provided with a <u>diagnosis</u> of **primary axillary hyperhidrosis** and the following criteria is met:

- The member must be 9 years of age or older
- There is documentation that the axillary hyperhidrosis is severe, intractable and disabling in nature as documented by at least one of the following:
 - Significant disruption of professional and/or social life as a result of excessive sweating
 - The condition is causing persistent or chronic cutaneous conditions (e.g., skin maceration, dermatitis, fungal infections, secondary microbial infections)
- Potential causes of secondary hyperhidrosis have been ruled out (e.g., hyperthyroidism)
- Must provide documentation showing the member has tried and failed or had an intolerance or contraindication to at least 2 months of topical aluminum chloride 20%
- The requested dose and frequency is in accordance with FDA-approved labeling, nationally recognized compendia, and/or evidence-based practice guidelines
- **Initial Duration of Approval:** 6 months
- Reauthorization criteria
 - o Documentation of improvement from baseline
- **Reauthorization Duration of Approval:** 12 months

Coverage may be provided for any non-FDA labeled indication if it is determined that the use is a medically accepted indication supported by nationally recognized pharmacy compendia or peer-reviewed medical literature for treatment of the diagnosis(es) for which it is prescribed. These requests will be reviewed on a case by case basis to determine medical necessity.

Drugs are authorized in generic form unless the branded product is on the preferred drug list or the prescriber has indicated in writing that the branded product is medically necessary. If only the branded product is on the preferred drug list, the generic form will be considered non-preferred and shall not require the prescriber to indicate in writing that the branded product is medically necessary.



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QBREXZA (GLYCOPYRRONIUM) PRIOR AUTHORIZATION FORM

Please complete and fax all requested information below including any progress notes, laboratory test results, or chart documentation as applicable to Highmark Health Options Pharmacy Services. FAX: (855) 476-4158

If needed, you may call to speak	to a Pharmacy Services Rep	presentative. PHONE:	(844) 325-6251 Mon –	Fri 8 am to 7 pm	
	PROVIDER I	NFORMATION			
Requesting Provider:		NPI:	NPI:		
Provider Specialty:	Office Cor	Office Contact:			
Office Address:		Office Pho	Office Phone:		
	Office Fax	Office Fax:			
MEMBER INFORMATION					
Member Name:					
Member ID:		Member weight:	er weight: Height:		
REQUESTED DRUG INFORMATION					
Medication:	-	Strength:			
Directions:		Quantity:	Refills:		
Is the member currently receiving re-	quested medication? Yes	No Date I	Medication Initiated:		
Is this medication being used for a chronic or long-term condition for which the medication may be necessary for the life of the					
patient? Yes No					
Billing Information					
This medication will be billed: at a pharmacy OR medically, JCODE:					
Place of Service: Hospital	Provider's office Memb	er's home Other			
Place of Service Information					
Name:	NPI:				
Address:		Phone:			
MEDICAL HISTORY (Complete for ALL requests)					
Diagnosis: ICD Code:					
Is there documentation the axillary hyperhidrosis is severe, intractable and disabling? Yes No					
Is there significant disruption of professional and/or social life as a result of excessive sweating? Yes No					
Does the condition cause persistent of				fections, secondary	
	No	` ` ` `	, , ,	•	
Has secondary hyperhidrosis been ru	led out? Yes No				
CURRENT or PREVIOUS THERAPY					
Medication Name	Strength/ Frequency	Dates of Therapy	Status (Discontinue	ed & Why/Current)	
1/20urouron 1 (unic	serengen, rrequency	Dutes of Therapy	Status (Discontinue	ou co ((iij) cultone)	
	REAUTHO	ORIZATION			
Has the member experienced improvement with treatment? Yes No					
SUPPORTING INFORMATION or CLINICAL RATIONALE					
D					
Prescribing Provide	er Signature		Date		