

PRIOR AUTHORIZATION REQUEST FORM **Zurzuvae™**

For authorization, please answer each question and fax this form PLUS chart notes back to Real Rx at 385-425-4052.

Failure to submit clinical documentation to support this request will result in a dismissal of the request.							
If yo	ou have prior authorization question	ns, please call for assistance	385-42!	5-5094.			
Discl	aimer: Prior authorization request for	ms are subject to change in acc	ordance	with Fede	ral and State notice requirements.		
		AA A		15.4			
Date:		Member Name:		ID#:	ID#:		
DOB	OB: Gender:		Physician:				
Office Phone: Office Fax		Office Fax:	:		Office Contact:		
Height/Weight:							
preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria. Product being requested: □ Zurzuvae™ (zuranolone) Dosing/Frequency:							
		for reauthorization, procee					
1 1	Questions	tation with an OR/CVN or	Yes	No	Comments/Notes Please provide documentation		
 Is the request made by, or in consultation with, an OB/GYN or Psychiatric provider? 				Please provide documentation			
	Does clinical documentation show t	he member does NOT					
r	have a history of pre-partum major depressive disorder						
	previously treated within the last 12 months prior to the						
	oregnancy?	and although the construction			Bloom of the day of the state of		
	Does the member have a document depressive episode that began no e				Please provide documentation		
	rimester and no later than the first						
	delivery, as diagnosed by a structure	· ·					
	DSM-5?						
4. [Does clinical documentation include	one of the following?			Please provide documentation		
•		le (HAM-D) score is ≥ 24					
	(severe depression) OR;						
•		•					
_	(MADRS) score is ≥ 35 (severe d	•					
•	Patient Health Questionnaire-9						
	(severe depression) plus Edinbu Scale (EPDS) ≥18 Score OR;	igii Postiiatai Depression					
•	Beck Depression Inventory (BDI	1>29 (severe denression)					
	OR;	, (301010 00010331011)					
•	 Quick Inventory of Depressive S 	ymptomatology (self-					
	reported) (QIDS-SR) ≥16 OR;						

		Please provide documentation
n the pa	ast for this	condition? Please document

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Policy: PHARM-158

Origination Date: 12/20/2023 Reviewed/Revised Date: 01/17/2024 Next Review Date: 01/17/2025 Current Effective Date: 02/01/2024

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