

## ADHD Products Prior Authorization Request Form

Please complete this <u>entire</u> form and fax it to: 866-940-7328. If you have questions, please call 800-310-6826. This form may contain multiple pages. Please complete all pages to avoid a delay in our decision.

Allow at least 24 hours for review.

Section A – Member Inform	mation							
First Name:	Last Name	Last Name:				Member ID:		
Address:								
City:	City: State:				ZIP Code:			
Phone:		DOB:			Allergies:			
Primary Insurance Information	(if any):	1			•			
Is the requested medicati	on: □ New or □	Continuat	ion of Thera	apy? If continuation,	list sta	rt date: _		
Is this patient currently h	ospitalized?	Yes □ No	If recently	discharged, list disc	harge o	date:		
Section B - Provider Infor	mation							
First Name:			Last Name:				M.D./D.O.	
Address:			City:		State:		ZIP code:	
Phone:	Fax:		NPI#:		Specia	ılty:		
Office Contact Name / Fax atte	ention to:							
Section C - Medical Inform	nation							
Medication:						Strength:		
Directions for use:						Quantity:		
Diagnosis (Please be specific	& provide as muc	ch information	as possible)			ICD-10 CC	DDE:	
Is this member pregnant?	Yes □ No	If yes,	what is this	member's due date?				
Section D - Previous Med	ication Trials							
Medication Name	Strength	Dire	ctions	Dates of Therapy		Reason for failure / discontinuation		
Section E – Additional info								
Please refer	to the patient's	PDL at ww	w.uhcprovi	der.com for a list of	preferre	ed alterna	tives	



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Member Firs	t name:	Member Last name:	Member DOB:			
	Clinical and Drug Specific Information					
ALL REQUESTS						
□ Yes □ No	Does the patient have one of the following diagnoses? (If yes, check which applies)  □ Attention deficit hyperactivity disorder/attention deficit disorder (ADHD/ADD)  □ Autism Spectrum Disorder  □ Binge eating disorder (BED)  □ Depression					
□ Yes □ No	and is new to the plan?		n-preferred behavioral health medication			
□ Yes □ No		ceiving treatment with the requested i Il and must continue upon discharge?				
□ Yes □ No		ure to THREE preferred alternatives? above/MEDICAL RECORDS MUST BE	SUBMITTED)			
□ Yes □ No		istory of <u>intolerance or contraindication</u> above, including the intolerance or contraindication				
	LESS THAN THE FDA APPROVED MINIMUM AGE					
□ Yes □ No	Is the patient unresponsi	ve to, or has had an inadequate respo	onse to behavioral therapy?			
□ Yes □ No	Is the patient experiencin therapy?	g moderate-severe continuing disturl	pance in function despite behavioral			
		TI-SOURCE BRAND MEDICATION REC				
□ Yes □ No	<ul> <li>□ The multi-source brand is equivalent. Specify the assertion of the multi-source brand is RECORDS MUST BE State of the multi-source brand is destabilization of the past of</li></ul>	adverse reaction, allergy, or sensitivity: _ s being requested due to a therapeutic fa SUBMITTED s being requested because transition to a	e reaction, allergy, or sensitivity to a generic allure with the generic equivalent. <i>MEDICAL</i> a generic equivalent could result in heric version of the multi-source brand			
KAPVAY / CLONIDINE ER						
□ Yes □ No		•	AL RECORDS MUST BE SUBMITTED)			
□ Yes □ No	· -	•				



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Member Firs	t name:	Member Last name:	Member DOB:			
QELBREE						
□ Yes □ No	<ul> <li>(If yes, check which applied</li> <li>□ Amphetamine/dextroam</li> <li>□ Atomoxetine (generic S</li> <li>□ Clonidine ER (generic K</li> <li>□ Dexmethylphenidate EF</li> <li>□ Guanfacine ER (generic Methylphenidate extend</li> <li>□ Methylphenidate extend</li> <li>□ Methylphenidate extend</li> </ul>	(apvay) R capsule (generic Focalin XR)	eneric Adderall XR)			
□ Yes □ No	<ul> <li>⟨If yes, check which applied</li> <li>□ Amphetamine/dextroam</li> <li>□ Atomoxetine (generic S</li> <li>□ Clonidine ER (generic K</li> <li>□ Dexmethylphenidate EF</li> <li>□ Guanfacine ER (generic Methylphenidate extend</li> <li>□ Methylphenidate extend</li> <li>□ Methylphenidate extend</li> </ul>	(apvay) R capsule (generic Focalin XR)	g the intolerance or contraindication) eneric Adderall XR)  D)			
□ Yes □ No	Does the physician attest that use of a stimulant medication is not appropriate for the patient?					
		VYVANSE				
□ Yes □ No	( <i>If yes, check which applie.</i> □ Amphetamine/dextroam □ Dexmethylphenidate EF	Iure to any of the following? Is and complete Section D above/MEDICA Inphetamine salts extended-release (gene R capsule (generic Focalin XR) Ided-release tablet or capsule (generic Co	ric Adderall XR)			
□ Yes □ No	Does the patient have a history of intolerance or contraindication to any of the following?  (If yes, check which applies and complete Section D above, including the intolerance or contraindication)  □ Amphetamine/dextroamphetamine salts extended-release (generic Adderall XR)  □ Dexmethylphenidate ER capsule (generic Focalin XR)  □ Methylphenidate extended-release tablet or capsule (generic Concerta or Metadate CD)					
□ Yes □ No	Does the patient have a l	history of, or potential for, a substanc	e abuse disorder?			
□ Yes □ No		lure to atomoxetine (generic Strattera) above/MEDICAL RECORDS MUST BE				
□ Yes □ No		nistory of <u>intolerance or contraindication</u> above, including the intolerance or contr				

Provider Signature:	Date:	
	•	

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