SENTARA HEALTH PLANS

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions</u>: The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; fax to 1-800-750-9692. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. If information provided is not complete, correct, or legible, authorization may be delayed.

Non-Preferred Central Nervous System (CNS) Stimulants (For all ages)

 A review of written documentation to substantiate a complete, appropriate, and covered diagnosis for both new starts and members currently receiving any CNS stimulant listed below will be required before Prior Authorization approval. <u>Prescribing history alone WILL NOT meet criteria for approval.</u>

MEMBER & PRESCRIBER INFO	PRMATION: Authorization may be delayed if incomplete.
Member Name:	
Member Sentara #:	
Prescriber Name:	
Prescriber Signature:	
Office Contact Name:	
	Fax Number:
NPI #:	
DRUG INFORMATION: Authoriza	
Drug Form/Strength:	
	Length of Therapy:
Diagnosis:	ICD Code:
Weight (if applicable):	Date weight obtained:
Will the member be discontinuing a prev medication if approved for requested med	iously prescribed central nervous system (CNS) stimulant dication?
	□ Yes OR □ No
 If yes, please list the medication that will approval along with the corresponding ef 	be discontinued and the medication that will be initiated upon fective date.
Medication to be discontinued:	Effective date:
Medication to be initiated:	Effective date:

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DRUG(S) REQUESTED:	Check applicable drug(s) belo	ow. Box(es) must be checked to q	ualify, or
authorization process will be dela	ayed.		

Adhansia XR®	Adzenys XR-ODT® Adzenys ER® Suspension	amphetamine sulfate (Evekeo®)	Azstarys [®]
Cotempla XR- ODT®	Dyanavel® XR Suspension Dyanavel® XR Chewable Tablets	Evekeo ODT®	Jornay PM®
methylphenidate ER (Aptensio XR®)	methylphenidate TD Patch (Daytrana®)	Mydayis®	Quillichew® ER
Quillivant XR®	Xelstrym [™] (dextroamphetamine)		

CLINICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

	Member must	have tried	and failed	30 davs o	of therapy	with t	wo (2)	of the	following:
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- amphetamine-dextroamphetamine IR/ER (generic Adderall/Adderall XR®)
- □ dexmethylphenidate IR/ER (generic Focalin®/Focalin XR®)
- □ dextroamphetamine IR/SR (generic Dextrostat®/Procentra®/Zenzedi®/Dexedrine® IR/ER)
- □ methylphenidate IR/ER (generic Ritalin®/Methylin®/Ritalin SR®/Ritalin LA®/Concerta®/ Metadate CD®/Metadate ER®
- ☐ Member must have tried and failed <u>30 days of therapy</u> with Vyvanse[®] (<u>NOT</u> required for amphetamine sulfate (Evekeo[®]) or Evekeo ODT[®] requests)
- ☐ If the member is <u>over the age of 18</u>, member <u>must</u> also meet diagnostic criteria. The prior authorization form "CNS Stimulants for Adults Age 19 and Above" can be downloaded from:

 http://www.sentarahealthplans.com/providers/

Not all drugs may be covered under every Plan.

If a drug is non-formulary on a Plan, documentation of medical necessity will be required.

**Use of samples to initiate therapy does not meet step edit/preauthorization criteria. **

*Previous therapies will be verified through pharmacy paid claims or submitted chart notes. *