

Prescriber Signature:

## Office of Medical Assistance Programs – Fee-for-Service, Pharmacy Division Phone 1-800-537-8862 Fax 1-866-327-0191

STIMULANTS AND RELATED AGENTS PRIOR AUTHORIZATION FORM (Form effective 2/15/19) Prior authorization guidelines are accessible on the DHS Pharmacy Services website at <a href="http://www.dhs.pa.gov/provider/pharmacyservices/index.htm">http://www.dhs.pa.gov/provider/pharmacyservices/index.htm</a>. PRIOR AUTHORIZATION REQUEST INFORMATION PRESCRIBER INFORMATION total # of pgs: New request Renewal request Prescriber name/specialty: NPI: Name/phone of office contact: State license #: LTC facility contact/phone: Street address: Suite #: City/state/zip: Beneficiary name: DOB: Phone: Fax: Beneficiary ID#: **CLINICAL INFORMATION** Medication Requested (Names in parentheses are the brand name equivalents for reference purposes. IR = immediate-release; ER/XR = extended-release) Preferred Agents Non-Preferred Agents ☐amphetamine mixed salts IR Focalin XR capsule ☐ Adderall tablet ☐ Dyanavel XR suspension methylphenidate solution tablet (Adderall) ☐guanfacine ER tablet Adderall XR capsule Evekeo tablet (Methylin) Aptensio XR capsule methylphenidate IR tablet ☐ Adzenys ER suspension Intuniv tablet ■ Mydayis ER capsule ☐ProCentra solution ☐atomoxetine capsule ☐Adzenys XR-ODT ☐ Kapvay tablet (Ritalin) methylphenidate ER/SR ☐clonidine ER tablet (Kapvay) methamphetamine tablet Relexxii ER tablet □ Daytrana patch dextroamphetamine ER cap tablet (Ritalin-SR) ☐Concerta tablet Methylin solution ☐Ritalin tablet methylphenidate ER 24HR dextroamphetamine IR tablet ☐Cotempla XR-ODT methylphenidate chew (Methylin) ☐Ritalin LA capsule ☐ Desoxyn tablet
☐ Dexedrine Spansule ER Strattera capsule
Zenzedi tablet (Dexedrine IR) tab (Concerta) methylphenidate CD capsule dextroamphet/amphetamine Quillichew ER tablet (Metadate CD) mixed salts combo XR capsule ☐ Quillivant XR suspension dexmethylphenidate IR tablet (*Focalin*) methylphenidate ER capsule dexmethylphenidate XR cap (Focalin XR)
dextroamphetamine sol'n (ProCentra) (Ritalin LA) (Adderall XR) □ Vyvanse capsule Focalin tablet ■Vyvanse chewable tablet methylphenidate ER 72 mg tablet Quantity: Strength: Directions: # months requested: Weight (if <4 years old): Diagnosis: Diagnosis code (required): Did the prescriber or prescriber's delegate search the PDMP to review the beneficiary's controlled □Yes Submit documentation. substance prescription history before issuing this prescription for the requested agent?  $\square$ No If request is for a NON-PREFERRED agent, does the beneficiary have a history of trial and ☐Yes – Submit documentation. failure, contraindication, or intolerance to the preferred agents (listed above)? □No ---OR--- not applicable If request for a NON-PREFERRED agent, has the beneficiary been taking the requested non-□ Yes Submit documentation of drug regimen and  $\square$ No preferred medication within the past 90 days? clinical response Request for a Beneficiary LESS than 4 Years of Age Yes – Submit documentation of diagnosis. 1. Does the beneficiary have one of the following diagnoses? Check all that apply. □No – Submit medical literature supporting the use of the □ ADHD □autism ☐brain injury requested medication for the beneficiary's age and diagnosis. Yes
No (prescriber's specialty: 2. Is the requested medication prescribed by, or in consultation with, one of the following specialists? pediatric neurologist ☐ child/adolescent psychiatrist ☐ child development pediatrician 3. Has the beneficiary had a comprehensive evaluation by, or in conjunction with, the above specialist? Yes – Submit documentation of evaluation.  $\square$ No Request for a Beneficiary 18 Years of Age and Older □ADD/ADHD Initial request – Submit documentation of an initial evaluation that shows a history of symptoms that meet the current DSM criteria (note: a rating scale alone is not sufficient documentation). Renewal request – Submit documentation supporting the continued need for the medication to manage symptoms. Narcolepsy – Submit documentation of beneficiary's symptom history and results of an overnight sleep study (a PSG) AND a multiple 1. What is the sleep latency test (MSLT). beneficiary's diagnosis? Moderate to severe binge eating disorder (Vyvanse request) Initial request – Submit documentation of ALL of the following: an initial evaluation that shows a history of symptoms that meet the current DSM criteria; if the beneficiary does NOT have ADD/ADHD, the beneficiary has tried, or cannot try, SSRIs or topiramate, AND an offer of referral for cognitive behavioral therapy or other psychotherapy. Renewal request – Submit documentation that the beneficiary experienced a reduction in binge eating. 2. *Stimulant requests*: Does the beneficiary have a history of or currently have substance use disorder [SUD] Yes Submit documentation of a recent eval. (drugs OR alcohol)? No for current or past substance use. Yes – Submit documentation of treatment. 3. For a beneficiary with a history of or current SUD, does the beneficiary have documentation of active participation in, or successful completion of, a substance use disorder treatment program? No ---OR--- not applicable Yes – Submit documentation of test results. 4. For a beneficiary with a history of or current SUD, does the beneficiary have documentation of a recent □No urine drug screen (UDS) testing for licit (including fentanyl, oxycodone, tramadol, carisoprodol) and illicit drugs? ---OR--- not applicable PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO DHS - PHARMACY DIVISION

Date: