

NYRx the Medicaid Pharmacy Program Atypical Antipsychotics (AAP) Prior Authorization Worksheet

Fax this form to 1-800-268-2990

Processing may be delayed if information submitted is illegible or incomplete. Preferred drugs will not require PA if the required coverage parameters, outlined in the PDL, are found in the member's Medicaid claim history at the time of pharmacy claim submission. Non-preferred drugs will require PA. https://newyork.fhsc.com/

Enrollee Information			
Enrollee Last Name:			
Enrollee First Name:			
Enrollee Medicaid ID (2 letters, 5 numbers,	1 letter):		
Date of Birth:			
PRESCRIBER INFORMATION			
Prescriber Last Name:			
Prescriber First Name:			
National Provider Identifier (NPI) Number: _			
Board Certified Specialty:			
Prescriber Street Address:			
City:	State:	Zip Code:	
Prescriber Phone:	Prescribe	er Fax:	
REQUESTED DRUG INFORMATION			
Drug Name:			
Drug Strength:			
Quantity:	Number of Refills:		
Directions:			
Is this a New Prescription? ☐ Yes ☐ No			
If No , date therapy was initiated:			

Enr	ollee's Name (Last, First):
CRI	TERIA
1.	What is the patient's diagnosis?:
	ICD-10 diagnosis code:
2.	Is the diagnosis an off-label use for this medication?
	☐ Yes ☐ No
	If Yes, what is the clinical reason for off-label use?
For	major depressive disorder (MDD)
3.	Are you requesting Symbyax?
	☐ Yes ☐ No
	If Yes , has the patient experienced a treatment failure or adverse reaction with at least 1 antidepressant other than fluoxetine?
	☐ Yes ☐ No
	If No , has the patient experienced a treatment failure or adverse reaction with at least 2 different antidepressants?
	☐ Yes ☐ No
4.	If either selection is No , what is the clinical reason for use without meeting the required "trial of an antidepressant"?
For	PDL questions (non-preferred medications)
5.	Are you willing to prescribe a preferred agent?
	☐ Yes ☐ No
	If Yes , please see Antipsychotics – Second Generation PDL class for a list of preferred agents (https://newyork.fhsc.com/)
6.	Has the patient experienced treatment failure with a preferred atypical antipsychotic?
	☐ Yes ☐ No
7.	Has the patient experienced an adverse drug reaction with a preferred atypical antipsychotic?
	☐ Yes ☐ No

Enr	ollee's Name (Last, First):
CR1	ITERIA (CONTINUED)
8.	Is there a documented history of successful therapeutic control with a non-preferred atypical antipsychotic and transition to a preferred atypical antipsychotic is medically contraindicated?
	☐ Yes ☐ No
9.	What is the reason the patient is unable to use a preferred agent in the same drug class? (if necessary, fax additional pages)
For	Abilify MyCite
10.	Are you attesting that the patient has a smart phone and has been taught how to use the app?
	☐ Yes ☐ No
11.	Has the patient experienced a treatment failure or adverse reaction to oral aripiprazole?
	☐ Yes ☐ No
12.	Has the patient experienced a treatment failure of adverse reaction to injectable aripiprazole?
	☐ Yes ☐ No
	If No , what is the clinical reason the patient cannot use another formulation of aripiprazole?
	drugs with dose optimization limits (see posted PDL for dose optimization limits tps://newyork.fhsc.com/)
13.	Does this request exceed the Dose Optimization limit?
	☐ Yes ☐ No
14.	Are you willing to use a higher strength (If Yes , skip question #16)?
	☐ Yes ☐ No
15.	What is the clinical reason for not using a higher available strength or exceeding the daily limit?

Enro	ollee's Name (Last, First):
	TERIA (CONTINUED)
Fred	quency/quantity/duration (F/Q/D) limits (see posted PDL for limits)
Max	kimum dose criteria
16.	Does the dose requested exceed the FDA maximum recommended dose?
	☐ Yes ☐ No
	If Yes , what is the clinical reason for exceeding the FDA-approved max daily dose (MDD)?
Min	imum dose criteria (quetiapine/quetiapine ER, Seroquel®/Seroquel XR®)
	What is the clinical reason for prescribing a low dose of Seroquel that is under the minimum amount of 50 mg/day?
Dail	ly Quantity Limit (quetiapine/Seroquel®)
18.	Does the dose requested exceed the established daily quantity limit?
	☐ Yes ☐ No
	If Yes , what is the clinical reason for exceeding the FDA-approved quantity limit?
DUF	REDITS
DUF of a	R - Concurrent use of CNS stimulants and atypical antipsychotic for patients < 18 years age
	What diagnosis requires the concurrent use of a CNS stimulant and an atypical antipsychotic?
	R - Concurrent use of 2 or more atypical antipsychotics for > 90 days for patients < 21 rs of age
20.	What are the names of the atypical antipsychotics?
	What is the clinical reason for the concurrent use of 2 or more oral atypical antipsychotics for > 90 days?
- 22.	Do you plan to discontinue any of the antipsychotics? Yes No

Enrollee's Name (Last, First):					
DUR EDITS (CONTINUED)					
If Yes, what is the drug name and approximate date of discontinuation?					
Drug Name:					
Date of Discontinuation:					
DUR – Concurrent use of 3 or more atypical antipsychotics for > 180 days for patients \geq 21 years of age					
23. What are the names of the atypical antipsychotics?					
24. What is the clinical reason for the concurrent use of 3 or more oral atypical antipsychotics for > 180 days?					
25. Do you plan to discontinue any of the antipsychotics?					
☐ Yes ☐ No					
If Yes, which drug and what is the approximate date of discontinuation?					
Drug(s):					
Date of Discontinuation:					
Submission of this form confirms the information is accurate and true, and that the supporting					

Submission of this form confirms the information is accurate and true, and that the supporting documentation is available for review upon request of the NYSDOH or CMS. The submitter understands that any person who knowingly makes or causes to be made a false record to statement that is material to a Medicaid claim may be subject to civil penalties and treble damages under both federal and NYS False Claims Acts.

Fax Number: 800-268-2990

Billing Questions: 800-343-9000

For clinical concerns or Preferred Drug Program questions, please visit http://newyork.fhsc.com or call 877-309-9493.