

**(All requests must be approved in advance to insure authorization)**

Member Name: \_\_\_\_\_ Today's Date: \_\_\_\_\_  
Provider: \_\_\_\_\_ Medicaid ID #: \_\_\_\_\_  
Contact Phone Number: \_\_\_\_\_ Contact: \_\_\_\_\_  
Service Location: \_\_\_\_\_ Provider MIS #: \_\_\_\_\_

Current Diagnosis:  Schizophrenia  Other (please explain): \_\_\_\_\_

Is the member over age 18?  Yes  No    Is the member over age 65?  Yes  No  
Dementia related psychosis?  Yes  No

What are the member's specific symptoms that are being targeted with this treatment?

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The client's ability to tolerate extended exposure to Risperdal or Invega has been established by the use of oral Risperdal (including M-Tabs), oral Invega, or Risperdal Consta prior to the patient receiving Invega Sustenna. Please list the names of medications that establish this exposure, doses and dates of treatment:

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There is clear documentation that the client cannot take oral Risperdal (including M-tabs), oral Invega, or Risperdal Consta. Include patient-specific reasons why Invega Sustenna is expected to be effective even when these other medications were not:

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There is clear documentation that the client cannot be treated with Haldol Decanoate or Prolixin Decanoate:

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There is clear documentation that the client has been prescribed several oral antipsychotic medications, but could not be safely and effectively treated with any of those medications.  Yes  No (explain below)

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The client has agreed to receive the injections on a regular basis, at the interval prescribed, and a person or agency that is geographically accessible and capable of dispensing the injections at the required frequency has been identified.  Yes  No

There is not more than one provider prescribing antipsychotic medications to this client.  Yes  No

Recent laboratory tests (CBC, lipid panel, FBS) have been completed and reviewed:

Yes Date reviewed: \_\_\_\_\_  No

Results: \_\_\_\_\_

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The maximum FDA approved dosage is 234 mg each month. Amounts in excess of this dose and frequency have not been shown to have additional efficacy, so will not be authorized.

**Please list all current medications and doses:**

Medication:	Dosage:

Initial Prior Authorization for Invega Sustenna will be for 6 months. Subsequent prior authorization frequency may be determined, and will be contingent upon evidence of clinical efficacy and appropriate clinical monitoring.

**Dosage Information for Authorization:**

Please authorize for \_\_\_\_\_ months or \_\_\_\_\_ injections.

Dosage given on each appointment date \_\_\_\_\_ (mg)

Dates of injections: 


J2426 (Invega) x \_\_\_\_\_ Units (1 mg = 1 unit)  
96372 (injection) x \_\_\_\_\_ (number of injections)