

**Pharmacy Prior Approval Request for  
Aduhelm**

**Beneficiary Information**

1. Beneficiary Last Name: \_\_\_\_\_ 2. First Name: \_\_\_\_\_  
3. Beneficiary ID #: \_\_\_\_\_ 4. Beneficiary Date of Birth: \_\_\_\_\_ 5. Beneficiary Gender: \_\_\_\_\_

**Prescriber Information**

6. Prescribing Provider NPI #: \_\_\_\_\_  
7. Requester Contact Information - Name: \_\_\_\_\_ Phone #: \_\_\_\_\_ Ext. \_\_\_\_\_

**Drug Information**

8. Drug Name: \_\_\_\_\_ 9. Strength: \_\_\_\_\_ 10. Quantity Per 30 Days: \_\_\_\_\_  
11. Length of Therapy (in days):  up to 30 Days  60 Days  90 Days  120 Days  180 Days  365 Days

**Clinical Information**

1. Does the beneficiary have mild cognitive impairment due to Alzheimer's Disease or mild Alzheimer's Dementia?  Yes  No
2. Has the beneficiary received all of the tests listed below?
  - a. Clinical Dementia Rating (CDR) -Global Score of 0.5  Yes  No
  - b. Objective evidence of cognitive impairment at screening  Yes  No
  - c. Mini-Mental Status Exam (MMSE) score between 24 and 30 (inclusive) OR equivalent tool indicating MCI or mild dementia (NOTE: range of scores may be adjusted based on educational status of patient)  Yes  No
  - d. Positron Emission Tomography (PET) scan is positive for amyloid beta plaque or Cerebrospinal Fluid Test (collected via lumbar puncture) is positive for amyloid  Yes  No
3. Is the beneficiary age 50 or older?  Yes  No
4. Has the beneficiary undergone testing to rule out reversible causes of dementia  Yes  No
5. Has the beneficiary had an assessment including a review of current medications as a cause of intellectual decline?  Yes  No
6. Has the beneficiary had a recent (within one year) brain MRI prior to beginning treatment?  Yes  No
7. Has the Prescriber has assessed and documented baseline disease severity utilizing an objective measure/tool?  Yes  No
8. Does the Beneficiary does NOT have history or increased risk of amyloid related imaging abnormalities-edema (ARIA-E), which includes brain edema or sulcal effusions and amyloid related imaging abnormalities hemosiderin deposition (ARIA-H), which includes microhemorrhage and superficial siderosis?  Yes  No
9. Has the beneficiary had a failure of or inability to tolerate at least one other preferred cholinesterase inhibitor Alzheimer therapy for at least four months?  Yes  No Please List \_\_\_\_\_
10. Does the provider attests to obtain MRIs prior to the 7th infusion (first dose of 10 mg/kg) and 12th infusion (sixth dose of 10 mg/kg)?  Yes  No
11. Does the beneficiary have hypersensitivity to any components of Aduhelm™?  Yes  No
12. Is Aduhelm™ being prescribed by or in consultation with a neurologist or geriatrician or geriatric psychiatrist?  Yes  No

Signature of Prescriber: \_\_\_\_\_ Date: \_\_\_\_\_

**(Prescriber Signature Mandatory)**

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.

Fax this form to: (833) 404-2393

Pharmacy PA Call Center: (833) 585-4309

<https://www.covermy meds.com/main/prior-authorization-forms/>