

Standard Aduhelm® (aducanumab-avwa) Plan of Treatment

NOTE: We require MD office notes to support clinical treatment and may require a Letter of Medical Necessity (depending on diagnosis), to be able to verify eligibility and payment for this treatment through patient Medicare and/or other insurance plan.

1. Patient Name: _____ Height (inches): _____ Weight (lbs): _____

2. Allergies: _____

3. Diagnosis: Please select the appropriate G and F codes:

- G30.0 Alzheimer’s disease with early onset
- G31.84 Mild Cognitive Impairment, so stated
- G30.1 Alzheimer’s disease with late onset
- G30.8 Other Alzheimer’s disease
- F02.8 Dementia without behavior disturbance
- F02.81 Dementia with behavior Disturbance
- Other ICD-10 Code (Diagnosis/Description):** _____

<p>Acetaminophen:</p> <ul style="list-style-type: none"> <input type="checkbox"/> 1000 mgs PO <input type="checkbox"/> 650mgs PO <input type="checkbox"/> 500mgs PO <input type="checkbox"/> 325mgs PO 	<p>Diphenhydramine: <input type="checkbox"/> 25 mgs PO, <input type="checkbox"/> 50mgs PO, <input type="checkbox"/> 25 mgs IVP, <input type="checkbox"/> 50mgs IVP or</p> <p>Alternate oral antihistamine to diphenhydramine: <input type="checkbox"/> Cetirizine 10 mg,</p> <p><input type="checkbox"/> Loratadine 10 mg, Fexofenadine <input type="checkbox"/> 60mgs or <input type="checkbox"/> 180mgs</p> <p>Methylprednisolone <input type="checkbox"/> 40mgs IVP <input type="checkbox"/> 125mgs IVP or other _____ mgs IVP</p> <p>Famotidine: <input type="checkbox"/> 20mgs PO, <input type="checkbox"/> 40mgs PO, <input type="checkbox"/> 20mgs IVP, <input type="checkbox"/> 40mgs IVP</p>
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Pre-medicate with other: _____

Dose: Aduhelm® (aducanumab-avwa) over approximately one hour every 4 weeks and at least 21 days apart as directed to infuse per protocol via pump with 0.2 or 0.22-micron filter.

- Initial titration dosing:** Aduhelm® (aducanumab-avwa) according to dosing table below followed by maintenance dose of Aduhelm® (aducanumab-avwa) 10mg/kg every 4 weeks (at least 21 days apart):

IV Infusion (every 4 weeks)	Aduhelm® Dosage (administer over approximately one hour)
Infusion 1 and 2	1mg/kg
Infusion 3 and 4	3mg/kg
Infusion 5 and 6	6mg/kg
Infusion 7 and beyond	10mg/kg

- Maintenance Dosing:** Aduhelm® (aducanumab-avwa) is 10mg/kg in 100ml of 0.9% Sodium Chloride every 4 weeks (at least 21 days apart)

Monitoring: Referring provider to obtain MRIs prior to the 7th infusion (first dose of 10mg/kg) and 12th infusion (sixth dose of 10mg/kg).

Special orders: _____

If adverse drug reaction occurs, utilize the ADVERSE DRUG REACTION GUIDELINES

6. Physician’s Signature: _____ / _____ Date: _____

No Stamp Signatures

(Dispense as written)

(Substitution permitted)

Printed Physician’s Name with Credentials: _____ NPI: _____

7. Fax updated supporting clinical MD notes with each order renewal or change in orders

Infusion order forms and Adverse Drug Reaction Guidelines are available at www.palmettoinfusion.com



Phone: 1-800-809-1265 Fax: 1-866-872-8920

MRN: _____

DOB: _____

Patient Name: _____ Referral Date: _____

- Include signed and completed **Plan of Treatment**. *(MD must complete sections 1-7)*
(Infusion order forms & Standard Adverse Reactions orders are available at www.palmettoinfusion.com under Agency/MD tab)
- Include patient demographic information and insurance information. *(Copy of insurance cards if*
- Supporting clinical MD notes to include any past tried and/or failed therapies, intolerance, outcomes or contraindications to conventional therapy. Include any lab results and/or tests to support diagnosis.**
- Include recent (within one year) brain magnetic resonance imaging (MRI).
- Include primary caregiver contact information, authorization to release protected health information, and/or POA if applicable.

Name: _____ Phone: _____ Relationship: _____

Diagnostic test confirming the presence of amyloid plaque build up

Other as requested: _____

WARNINGS AND PRECAUTIONS. Amyloid Related Imaging Abnormalities. ADUHELM can cause amyloid related imaging abnormalities-edema (ARIA-E), which can be observed on MRI as brain edema or sulcal effusions, and amyloid related imaging abnormalities hemosiderin deposition (ARIA-H), which includes microhemorrhage and superficial siderosis. Obtain recent (within one year) brain magnetic resonance imaging (MRI) prior to initiating treatment. The safety of ADUHELM® in patients with any pre-treatment localized superficial siderosis, 10 or more brain microhemorrhages, and/or with a brain hemorrhage greater than 1 cm within one year of treatment initiation has not been established.

Hypersensitivity Reactions. Promptly discontinue the infusion upon the first observation of any signs or symptoms consistent with a hypersensitivity