

State of Louisiana

Louisiana Department of Health Bureau of Health Services Financing

PRIOR AUTHORIZATION REQUEST COVERSHEET

Please check the member's appropriate health plan listed below:

Retail Pharmacy Requests
Magellan Medicaid Administration, LLC For Aetna Better Health of Louisiana, AmeriHealth Caritas Louisiana, Healthy Blue, Humana, LA Healthcare Connections, United Healthcare Phone: 1-800-424-1664 / Fax: 1-800-424-7402
Fee-for-Service (FFS) Louisiana Legacy Medicaid Phone: 1-866-730-4357 / Fax: 1-866-797-2329 / www.lamedicaid.com
Requests for Medications Through Medical Benefit
Aetna Better Health of Louisiana – Medical Benefit – Physician Administered Drugs Phone: 855-242-0802 / Fax: 844-227-9205 / TTY: 855-242-0802, 711
AmeriHealth Caritas Louisiana Phone: 1-800-684-5502 / Fax: 1-855-452-9131 / www.amerihealthcaritasla.com/pharmacy/priorauth.aspx
Healthy Blue – Medical Injectables 1-844-521-6942 (M–F 7 a.m.–7 p.m., Sat. 9 a.m.–1 p.m. CT) / Fax: 844-487-9291 CenterX®: Submit through EPIC EMR
Humana – Professionally Administered Drugs <u>Availity.com</u> (registration required) Phone: 1-866-461-7273 (M–F 7 a.m.–10 p.m. CT) / Fax: 1-888-447-3430 / (request form at <u>Humana.com/medPA</u>
LA Healthcare Connections – Physician Administered Medication (Buy and Bill) Phone: 1-866-595-8133 / Fax: 1-866-925-3006
United Healthcare – Medical Benefit Phone: 1-888-397-8129 / Fax: 877-271-6290 / www.UHCprovider.com
DDH/ACV AND CONFIDENTIALITY WADNING

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PLEASE CALL IF YOU HAVE ANY PROBLEMS RECEIVING THIS FAX OR IF PAGES ARE MISSING

Magellan Medicaid Administration

Louisiana Medicaid

Aducanumab-avwa (Aduhelm®) Clinical Authorization Form

Fax this form to 1-800-424-7402

Please fill out all applicable sections on all pages completely and legibly. Attach any additional documentation that is important for the review (e.g., chart notes or lab data, to support the prior authorization). Incomplete forms will not be approved. Information contained in this form is Protected Health Information under HIPAA.

SECTION 1: SUBMISSION		
Submitted to:		
Receiver Phone:	Receiver Fax:	Date:
SECTION 2: PRESCRIBER INFORM	ATION	
Prescriber Last Name:		
Prescriber First Name:		Middle Initial:
Prescriber NPI: Pl	an Provider #:	Specialty:
Prescriber Street Address:		
City:	State:	Zip:
Prescriber Phone:	Prescriber F	ax:
Office Contact Name:	Cor	ntact Phone:
SECTION 3: PATIENT INFORMATION	ON	
Patient Last Name:		
Patient First Name:		Middle Initial:
Date of Birth: Pa	atient Phone:	
Sex: Male Female] Other	
Patient Street Address:		
City:	State:	Zip:
Plan Name (if different from Section	1):	
Member #: Medica	nid #:	Plan Provider ID:
CCN #:		
EPSDT Support Coordinator contact	information (if applicable)	:
EPSDT Support Coordinator First Nam	e:	
EPSDT Support Coordinator Last Nam	e:	
EPSDT Support Coordinator Phone: _		-
SECTION 4: PRESCRIPTION DRUG	INFORMATION	
Drug Name: aducanumab-avwa (Ad	duhelm®) Quantity:	Day Supply:
Drug Strength: 170 mg/ 1.7 mL	☐ 300 mg/ 3 mL	

Patient's Name:			
SECTION 4: PRESCRIPTION DRUG INFORMATION (CONTINUED)			
Titration Dosing (Select one): 1 mg/kg/dose IV q4 weeks x 2 doses 6 mg/kg/dose IV q4 weeks x 2 doses Maintenance Dosing: 10 mg/kg/dose IV q4 weeks Other:			
1. This request is for:			
☐ Initiation of Treatment ☐ Continuation of Treatment			
SECTION 5: PATIENT CLINICAL INFORMATION			
2. Does the patient have a diagnosis of Alzheimer's disease?Yes NoIf Yes, date diagnosed:			
 3. Specify severity of cognitive impairment / dementia: Mild Cognitive Impairment Moderate Dementia Severe Dementia 			
 4. Was the presence of beta-amyloid plaques confirmed by one of the following? Positron emission tomography (PET) scan: Yes No 			
If Yes, date of test: Prescriber Initials: Cerebrospinal fluid (CSF) testing: Yes No			
If Yes, date of test: Prescriber Initials:			
SECTION 6: FOR INITIATION OF THERAPY REQUESTS ONLY			
Document objective evidence of mild cognitive impairment or mild dementia due to Alzheimer's disease below. (Both are required.) Clinical Dementia Rating-Global Score (CDR-GS) Score: Date: Mini-Mental State Exam (MMSE)			
Score: Date:			
Specify tool used to document baseline disease severity. (Note: Same tool must be used for baseline assessment and for ongoing assessments.) Alzheimer's Disease Assessment Scale – Cognitive Subscale (ADAS-Cog-13)			
Score: Date:			
Clinical Dementia Rating – Sum of Boxes (CDR-SB) Score: Date:			
Montreal Cognitive Assessment (MoCA)			
Score: Date:			
Repeatable Battery for Assessment of Neuropsychological Status (RBANS) Score: Date:			
Other:			
Score: Date:			
(Name of tool and defining parameters for disease severity for this tool must be included.)			

Pat	ient's Name:		
SE	CTION 6: FOR INITIATION OF THERAPY REQUESTS ONLY (CONTINUED)		
5.	Does the patient have any contraindication to magnetic resonance imaging (MRI)? Yes No If Yes, explain:		
6.	Most recent MRI Date:		
7.	Please initial below to confirm the results of the MRI: Were there any findings of localized superficial siderosis? Yes No Prescriber Initials: Were there findings of less than 5 brain microhemorrhages? Yes No Prescriber Initials: Were there findings of any brain hemorrhages > 1 cm within the past year? Yes No Prescriber Initials:		
8.	Is the patient currently taking blood thinners (except ≤ 81 mg aspirin)? ☐ Yes ☐ No		
9.	Is the patient ambulatory? ☐ Yes ☐ No		
10.	Has the patient had a bleeding disorder or cerebrovascular abnormalities (including, but not limited to, stroke or transient ischemic attack [TIA]) in the last 12 months? \square Yes \square No		
11.	Have other causes of cognitive impairment been ruled out (including, but not limited to, alcohol/substance abuse, frontotemporal dementia [FTD], Lewy body dementia [LBD], Parkinson's disease dementia, unstable psychiatric illness, and vascular dementia)? Yes No		
12.	Does the patient have a history of unstable angina, myocardial infarction, advanced chronic heart failure, clinically significant conduction abnormalities or unexplained loss of consciousness within 1 year of treatment initiation? Yes No		
13.	Has the patient had a seizure in the past 3 years? $\hfill \Box$ Yes $\hfill \Box$ No		
SE	CTION 7 - FOR CONTINUATION OF THERAPY REQUESTS ONLY		
Dat	te of treatment initiation: Number of doses since initiation:		
Pro	vide date of most recent MRI: (See criteria for MRI recommendations.)		
Not	te: It is recommended that practitioners use the same MRI device with the same imaging protocol for a given patient whenever possible to assist in comparing the images.		
	continuation of therapy requests, current clinical symptom severity and MRI findings must be sed below (see next page).		

Patient's Name:
SECTION 7 - FOR CONTINUATION OF THERAPY REQUESTS ONLY (CONTINUED)
ARIA-E clinical symptom severity:
☐ None ☐ Mild ☐ Moderate ☐ Severe
ARIA-E radiographic severity:
□ None □ Mild □ Moderate □ Severe
ARIA-H clinical symptoms:
☐ Yes ☐ No
ARIA-H radiographic severity:
□ None □ Mild □ Moderate □ Severe
14. Has the patient progressed to the moderate or severe stage of Alzheimer's disease?☐ Yes☐ No
15. Since baseline assessment, has the patient had a positive clinical response to treatment
demonstrated by assessment with the same validated tool that was used to establish baseline
disease severity?
☐ Yes ☐ No
16. Name of tool used to assess baseline disease severity and ongoing assessments:
Date of baseline assessment: Score:
Date of most recent follow-up assessment: Score:
SECTION 8 - ADDITIONAL CLINICAL INFORMATION
SECTION 9: PHARMACY INFORMATION (OPTIONAL)
Name of Dispensing Pharmacy:
Pharmacy NPI: Pharmacy Phone:
Pharmacy Street Address:
City: State: Zip:
City State Zip
☐ Attachments
By signing this request, the prescriber attests that the information provided herein is true and accurate to the
best of his/her knowledge. Also, by signing and submitting this request form, the prescriber attests to statements in the 'Attestation' section of the criteria specific to this request, if applicable.
Prescriber Signature: Date:
(Proxy signatures are not accepted.)
Mail requests to:
Magellan Medicaid Administration, LLC
Attn: GV - 4201
P.O. Box 64811
St. Paul, MN 55164-0811 Phone: 1-800-424-1664 Fay this form to 1-800-424-7402