

New Hampshire Medicaid Fee-for-Service Program

Prior Authorization Drug Approval Form

Monoclonal Antibodies Directed Against Amyloid for the Treatment of Alzheimer's Disease

DATE OF MEDICATION REQUEST: /	/												
SECTION I: PATIENT INFORMATION AND MEDICATION I	REQUESTED												
LAST NAME:	FIRST NAME:												
MEDICAID ID NUMBER:	DATE OF BIRTH:												
	_	_											
GENDER: Male Female													
Drug Name:		Strength:											
Dosing Directions:		Length of Therapy:											
SECTION II: PRESCRIBER INFORMATION													
LAST NAME:	FIRST NAME:												
SPECIALTY:	NPI NUMBER:												
PHONE NUMBER:	FAX NUMBER:												
	-												
SECTION III: CLINICAL HISTORY													
 Does the patient have a confirmed diagnosis of mild of Alzheimer's Disease or mild Alzheimer's Disease? 	ognitive impairmen	t (MCI) due to	Yes No										
Clinical Dementia Rating (CDR) – Global Score	of 0.5 to 1 🗌 Yes [No											
Objective evidence of cognitive impairment at	screening 🗌 Yes [No											
Mini-Mental Status exam (MMSE) score betwee	en 22 and 30 (inclu	sive) 🗌 Yes 🗌 No											
 Positron Emission Tomography (PET) is positiv cerebrospinal fluid assessment of amyloid bet to confirm diagnosis Yes No 		•											

Fax to Prime Therapeutics Management if medications will be dispensed by a pharmacy and will be administered by the patient or caregiver at home. Phone: 1-866-675-7755 Fax: 1-888-603-7696

Fax to DHHS if medication is dispensed/administered by the office or outpatient setting: Phone: 1-603-271-9384 Fax: 1-603-314-8101





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PA	TIENT	LAST	ΓN	AME	:									ΡΑΤΙ	ENT	FIRST	NAN	VE:							
 Have the following conditions been ruled out: vascular dementia, dementia with Lewy bodies, frontotemporal dementia, normal pressure hydrocephalus?] Ye	s 🗌] No										
3.	Has th the pa	-					roke	ort	ransie	ent is	chen	nic at	tacl	k or ı	unexp	olaine	ed los	s of (conso	ciousi	ness	in [] Ye	s 🗌] No
4.	4. Has the patient had a brain hemorrhage, bleeding disorder, cerebrovascular abnormality, or Yes Yes No cardiovascular conditions (e.g., unstable angina, myocardial infarction, advanced congestive heart failure, clinically significant conduction abnormalities) in the past 12 months?] No												
5.	Is the	pati	ent	: on a	an a	anti	i-pla	telet	, antio	coagi	ulant	, or a	nti-	thro	mbin	med	icatio	on?					Ye	s] No
6.	6. Is the prescriber a neurologist or gerontologist or has a neurologist or gerontologist been consulted?] Ye	s 🗌] No											
7.	Has tl mont		atie	nt re	ecei	ive	d a b	asel	ine m	agne	tic re	esona	nce	e ima	ging	(MRI)) with	nin th	e pas	st 12] Ye	s 🗌] No
8.	(Aduł	nelm	® 0	nly):	Wi	ill tł	ne pa	atier	it rece	eive a	h brai	in MF	RI p	rior t	o the	5th,	7th,	9th, a	and 1	L2th c	doses	s? [] Ye	s 🗌] No
9.	(Leqe	mbi®	® Or	۱y): ۱	Wil	ll th	ne pa	atien	t rece	ive a	brai	n MR	l pr	ior to	o the	5th,	7th, a	and 1	.4th c	doses	?	[Ye	s	No
10	(Kisur	าla™	on	ly): V	Vill	the	e pat	ient	receiv	ve a l	orain	MRI	prie	or to	the 2	2nd, 3	3rd <i>,</i> 4	lth, a	nd 7t	th do	ses?	[Ye	s	No
11	Has t	he pa	atie	nt ex	kpe	erie	nced	l any	ofth	e foll	owir	ıg?											Ye	s 🗌] No
	• Pi	re-tre	eati	ment	t lo	cali	ized	supe	erficia	l side	rosis	5													
				n mic				•	S																
				norrh	-																	_	_	_	-
12.	Has a			e ass	sess	sme	ent b	been	comp	leteo	d wit	h at l	eas	t one	e of th	ne fo	llowiı	ng?				L	_ Ye	s 🗋	No
	AAIn	lzhei npair	me me me	r's Di ent ve	isea ersi	ase ion	Coo [AD	pera CS-A	ient S itive S DL-M n of B	tudy CI]	-Acti	vities			-		•	-	ild Co	ogniti	ive				
13	Has ti clinica	•							•					•		l risk	s anc	l min	imal	estak	lishe	ed [Ye	s] No
	CIINIC	ai pe	ner	it ba	sec	ı or		ncal	unais	10 02	ne w	itri tr	eat	men	Lſ										

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PATIENT LAST NAME:								I	PATIENT FIRST NAME:															
14. F	14. For renewals (every 6 months): Has the patient demonstrated stability, improvement, or slowed Yes No																							

rate of progression in one of the following assessments?

- ADAS-Cog 13
- ADCS-ADL-MCI
- MMSE
- CDR-SB

Renewal assessment results:

15. Has the patient progressed to moderate or severe Alzheimer's Disease?	Yes No
16. Has the patient continued dosing at 10 mg/kg every 4 weeks (Aduhelm®) or every 2 weeks (Leqembi™) or 1,400 mg every 4 weeks (Kisunla™)?	Yes No
17. Has the patient received ongoing MRI monitoring as directed in the package insert (questions 8–10 above)?	Yes No
18. Did the MRI show > 10 new incident microhemorrhages or ≥ 2 focal areas of superficial siderosis?	Yes No
19. Will a follow-up MRI be performed to assess stability?	🗌 Yes 🗌 No
20. Do the benefits outweigh the risks based on the MRI results?	Yes No
Please provide any additional information that would help in the decision-making process. If addit	ional space is

needed, please use a separate sheet.

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PATIENT LA	ST NAME:					ΡΑΤΙ		ST NA	ME:						
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PRESCRIBEI Facility whe				d.						DATE:					
racinty with			provide	u.											

Medicaid provider number of facility:

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