

Checklist for Kisunla (donanemab) Referral

Patien	nt Name:	: Dob: Dob: Date:
Referr	ring Phys	sician:NPI:NPI:
Office	Contact,	:/Title/Email:
Office	Address	s:
Office	Phone:	Office Fax:
Best c	ontact n	number for physician in case of reaction:
Please	return c	completed checklist and checklist items to initiate referral. Use this form as fax cover sheet.
	Patien	t demographic information
	Insura	nce information and copy of insurance card/s (front and back). *Include primary and secondary insuranc
	Suppo	orting clinical notes and office visits. Two notes preferred.
	0	Note should include any therapies tried/failed, and must include discussion about Kisunla Medication list and allergies
	o Suppo	Cognitive assessment and functional assessment with score and interpretation orting lab reports/imaging for Kisunla treatment
	0	MRI within 1 year of treatment start
	0	Confirmation of amyloid beta pathology (LP or PET Scan)
	0	ApoE testing to determine ARIA risk
	0	CMS Registration (must be completed every 6 months)
	Durabl	le Power of Attorney for Health Care (DPAHC), if applicable
	Kisunla	a Prescribing Order and Indication Checklist (see attached)

Fax all information to our Infusion Coordinator: 508-698-8671

Call with any questions: 781-551-5812 option 4



Prescribing Order: Kisunla (donanemab)

Date of	Order	·	☐ New Start ☐ Maintenance Date of last infusion:			
Patient	Name	:	DOB:		M/F:	
Diagno	sis (inc	lude ICD-10 code/s):				
□ NKI	DA	Allergies:				
Patient	Weigh	nt:				
Premedication: ☐ Acetaminophen 1000mg PO ☐ Loratadine 10mg or Cetirizine 10mg PO ☐ Diphenhydramine 25mg PO			Diphenhydramine 25m Solu-medrol 125mg IV Other:	-		
Lab Ord						
	New s	lication Order: start: Kisunla 700mg every 4 weels for enance: Kisunla 1400mg every 4 wee	ks over 30min			
Admini	stratio ✓ Ho ✓ Ho	on: old infusion if no MRI Brain prior to the old infusion and notify provider if pation Headache, Confusion, Dizziness case of infusion reaction, STOP infus	ie 2 nd , 3 rd , 4 th ar ient experiencin s, Nausea, Visio	nd 7 th infusion ng any of the following s n Changes, Gait Change	igns of ARIA: s, Seizures	
Orderir	ng Prov	vider Name		NPI		
Signatu	ire			Date		



Patient Name:	DC	DB:	

Legembi/Kisunla Indication Checklist

	INFORMATION REQUIRED F	OR TREATMENT INITIATION				
1)	Patient ICD-10 (select all that apply)	Clinical Diagnosis (select one)				
	G30.0 Alzheimer's disease, early onset	☐ Mild cognitive impairment due to AD				
	☐ G30.1 Alzheimer's disease, late onset	☐ Mild AD Dementia				
	G30.9 Alzheimer's disease, unspecified					
	G31.84 Mild cognitive impairment					
2)	Cognitive Screening: within 6 months Date:					
	Name of Test Used: Score:					
	Interpretation: Functional Screening: within 6 months Date:					
	Name of Test Used:	Score:				
3)	Confirmation of Amyloid-Beta Pathology: MUST provide supporting documentation					
	☐ Amyloid PET Scan Date:	Result:				
	☐ CSE Amyloid Confirmation Date:	Result:				
	,					
4)) Monitoring for Amyloid Related Imaging Abnormalities (ARIA)					
	LEQEMBI/KISUNLA requires brain MRI within 1 year of treatment start date					
	Initial MDI Prain Data					
	☐ Initial MRI Brain Date:					
	Evidence of ARIA-E					
	Evidence of Aria-n - ivegative	- FUSITIVE				
5)	Schedule for MRI Monitoring: the following is	required by Infusion Center of New England				
-,	☐ Legembi: Obtain MRI prior to the 5 th , 7					
	☐ Kisunla: Obtain MRI prior to the 2 nd , 3 rd	•				
6)		□ No				
7)	_	□ No				
8)		□ No				
٥,		Result:				
9)	CMS REGISTRATION NUMBER					