

Phone: 732-390-7750 Fax: 844-683-2244 AsteraCancerCare.org

GFNFRAI

PATIENT REFERRAL FORM

Patient Name:				Pt. DOB: / /			
	Last	First	Middle				
Patient Address: _							
Patient City:			Pt. State:	Pt. Zip:			
Patient Phone: (_)			Pt. Height:	in.		
DX:				Pt. Weight:	lbs.		
Patient Allergies: _							
Insurance:				ID#:			
Referred by:			·	NPI#:			
Office Contact (Re			Offic	ce Ph:(
			Offic	ce Fax: ()			
Office Administrat	or (Required)	·	Administrat	or Ph: ()			
□Edison □Jersey C Required Items/Inf	ity	Robbinsville	□ Brick □ Bridgewater □ Rutherford □ Some	erset T Toms River			
(prescription on		_	·	act dosage, and direction	ns		
☐ Copy of current in	nsurance card						
☐ Recent MD consu	ıltation notes:	relevant diseas	e being treated must	be mentioned in report			
☐ Allergies and curi	ent medicatio	n list					
☐ Current labs requ	iired for specif	c medication, a	s noted on the follow	ing page(s) of this form			
Has the patient initi	ated treatmen	t at your office?	P \(\subseteq \text{Ye}	es 🗆 No			
$\ \square$ If any future lab t	ests are neede	d, please provi	de patient with a pres	cription, and have patien	t bring on		
day of treatment. R	Results will be s	ent to referring	g physician.				
Please note:							
	-	-	-	eir initial infusion at Aste	-		
include diagnosis, p	revious treatm	ents/response	to treatments and be	on letterhead with physi	cian signatur		

- 2. Benefit investigations, copay assistance and prior authorizations will be handled by the Astera precert staff if required by the payer. Right to auto-substitute biosimilars based on payer's preference. Detailed clinical notes providing supportive documentation are required for authorization requests which may take 3-5 business days depending on the payer and receipt of complete documentation from the referring office. The precert staff will update the referring doctor's office during this process and contact the patient to discuss cost and financial assistance options. For certain medications, patients will be required to register/enroll with the pharmaceutical company prior to rendered services and will receive a call from an Astera Financial Counselor to assist with this
- 3. A pretreatment education session will be provided by an Advanced Practice Provider.
- 4. Once the infusion is complete, a follow-up notice will be faxed to the to the referring provider.

Patient Name:				DOB:	/_	/					
	Last First	٨	Middle								
	box for medication requested, at Once all documentation is received Required Current Lab Re	d, we will contac		=							
Note: Progress	notes and labs must be comple	eted within the	e previous 6 month	s for all new and	renew	ed prescriptio	าร				
□ Actemra	CBC, Lipid Panel, Liver Functio	n, PPD (prior t	o initiation)								
☐ Benlysta (IV)	None										
□ Briumvi	CBC, Quantitative Serum Immunoglobulin, Prior to initiation – Hep B Serology (Hep B surface antiged Hep B surface antibody and Hep B core antibody) Confirm No Vaccinations within 4 Weeks of Therapy										
☐ Cimzia	CBC, Prior to initiation – PPD and Hep B Serology (Hep B surface antigen, Hep B surface antibody and Hep B core antibody)										
□ Cytoxan	CBC, CMP, UA	CBC, CMP, UA									
☐ Entyvio	Liver Function, PPD (prior to i	nitiation)									
□ Evenity	CMP, Dexa Scan within 2 year	rs 🗆 Confi	rm pt. has not had	an MI or stroke	within	previous year					
☐ Fasenra	Peak Flow and Other Pulmonary Function Tests										
□ Ilumya	CBC, CMP, Prior to initiation – PPD and Hep B Serology (Hep B surface antigen, Hep B surface antibody and Hep B core antibody) ☐ Confirm up to date with vaccines and no live vaccinations within 4 weeks prior to starting therapy or have an active infection. Evaluated for malignancy.										
□ IVIG	Hematocrit, Hemoglobin, IgG Concentrations, Platelets, Renal Function Tests, Urine Output Provide dose basis in mg/kg. Doses will be rounded to the nearest vial size available.					utput					
☐ Kisunla	Prior to initiation – confirm presence of amyloid beta pathology and brain MRI (within 1 year), completed Benefits investigation and Care Coordination Forms										
☐ Krystexxa	G6PD Deficiency, Serum Uric Acid Levels, Confirm Oral Urate Lowering Agent Discontinued										
□ Leqembi	Prior to initiation – confirm pr	esence of amy	vloid beta patholog	gy and brain MRI	(within	1 year)					
☐ Leqvio	Lipid Panel										
□ Nucala	FEV1, Peak Flow and Other Pulmonary Function Tests										
□ Ocrevus	CBC, prior to initiation - Hep B Serology (Hep B surface antigen, Hep B surface antibody and Hep B core antibody) □ Confirm No Vaccinations within 4 Weeks of Therapy										
□ Orencia (IV)	Prior to initiation – PPD and I Hep B core antibody)	Hep B Serology	y (Hep B surface ar	ntigen, Hep B sur	face an	tibody and					

Radicava	None
Remicade/Inf	lectra (Biosimilar might be replaced if appropriate) CBC, Liver Function, Prior to initiation – PPD and Hep B Serology (Hep B surface antigen, Hep B surface antibody and Hep B core antibody)
Rituxan/Riabı	ni/Truxima/Ruxience (CMS approved indications only - Biosimilar might be replaced if appropriate) CBC, prior to initiation - Hep B Serology (Hep B surface antigen, Hep B surface antibody and Hep B core antibody) Confirm No Vaccinations within 4 Weeks of Therapy
Saphnelo	Up to date with all immunizations before treatment initiation and confirm no live or live attenuated vaccines are given concurrently.
Simponi Aria	(IV) CBC, Liver Function, Prior to initiation – PPD and Hep B Serology (Hep B surface antigen, Hep B surface antibody and Hep B core antibody)
, , ,	Crohn's Disease Indication only - CBC, CMP (with LFTs), Prior to initiation – PPD and Hep B Serology (Hep B surface antigen, Hep B surface antibody and Hep B core antibody) Confirm No Vaccinations within 4 Weeks of Therapy or have an active infection
Soliris	Meningococcal Vaccination
Tezspire	FEV1, Peak Flow and Other Pulmonary Function Tests
Tysabri	MRI (MS patients), TOUCH Program Registration
Vpriv	Gene Testing (GBA – Velaglucerase Alfa)
Vyepti	None
Vyvgart	Anti-AChR Antibody Positive, No Live Vaccines During Therapy
	ulo (SQ – CIDP and Myasthenia Gravis) Anti-AChR Antibody Positive (Myasthenia Gravis only). No live vaccines during therapy. Confirm no active infection.
Xolair	Asthma - Baseline Serum IgE, FEV1, Peak Flow, Other Pulmonary Function Test Chronic Idiopathic Urticaria – None