

Phone: 732-390-7750 Fax: 844-683-2244 AsteraCancerCare.org **PATIENT REFERRAL FORM GENERAL**

Patient Name:				/		
	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#:		
				ice Ph: ()		
				ice Fax: ()		
Office Administrator (Required): Adm						
Astera Infusion Thera	apy scheduling I	ocation request:				
■Bridgewater ■Ea	ast Brunswick	□Edison □Jers	ey City	Robbinsville □Ruthe	erford D S	omerset
Required Items/In	fusion Process	:				
□ Valid/signed wr	<u>itten</u> prescript	ion including na	me of medication, ex	kact dosage, and dire	ctions	
(prescription or	nly valid for 6 i	months, includin	g refills)			
☐ Copy of current	insurance card					
☐ Recent MD cons	ultation notes	: relevant diseas	se being treated must	be mentioned in rep	ort	
 Allergies and cur 	rent medication	on list				
 Current labs req 	uired for speci	fic medication, a	s noted on the follow	ving page(s) of this fo	rm	
Has the patient init	′es 🗆	No				
□ If any future lab	tests are need	ed, please provi	de patient with a pres	scription, and have p	atient brin	ig on
day of treatment.	Results will be	sent to referring	g physician.			
Please note:						

- 1. A Letter of Medical Necessity is required for all patients receiving their initial infusion at Astera (letter must include diagnosis, previous treatments/response to treatments and be on letterhead with physician signature).
- 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. 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 process.
- 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	Middle				
	ne box for medication requested, attach Once all documentation is received, we Required Current Lab Result:	e will contact your patient to s				
Note: Progres	ss notes and labs must be completed	within the previous 6 mont	hs for all new and renewed prescriptions.			
☐ Actemra	CBC, Lipid Panel, Liver Function, P	PD (prior to initiation)				
☐ Benlysta (IV	/) None					
□ Briumvi	CBC, Quantitative Serum Immunoglobulin, 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					
□ Cimzia	CBC, Prior to initiation – PPD and Hep B Serology (Hep B surface antigen, Hep B surface antibody and Hep B core antibody)					
☐ Cinqair	Peak Flow and Other Pulmonary Function Tests					
☐ Cytoxan	CBC, CMP, UA					
☐ Entyvio	Liver Function, PPD (prior to initiation)					
☐ Evenity	CMP, Dexa Scan within 2 years	☐ Confirm pt. has not ha	d an MI or stroke within previous year			
☐ Fasenra	Peak Flow and Other Pulmonary I	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.					
☐ Krystexxa	G6PD Deficiency, Serum Uric Acid	Levels, Confirm Oral Urate	Lowering Agent Discontinued			
□ Leqvio	Lipid Panel					
□ Nucala	FEV1, Peak Flow and Other Pulmo	onary Function Tests				
□ Nulojix	CBC, EBV Serology, Magnesium, C	Operative Report, Potassiun	n, PPD (prior to initiation)			
□ Ocrevus	CBC, prior to initiation - Hep B Ser Hep B core antibody) Confirm No Vaccinations within	-	en, Hep B surface antibody and			

☐ Orencia (IV)	Prior to initiation – PPD and Hep B Serology (Hep B surface antigen, Hep B surface antibody and Hep B core antibody)				
□ Panhematin	CMP, Iron Studies, Prior to initiation - Urinary levels of porphobilinogen (PBG), delta aminolevulinic acid (ALA), and total porphyrin				
☐ Prolastin	Alpha 1 Proteinase Inhibitor Serum Levels and Lung Function ☐ IgA antibodies negative for patient with IgA deficiency				
☐ Radicava	None				
☐ Remicade/Ir	nflectra (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/Rial	bni/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	a (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)				
□ Skyrizi (IV)	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				
☐ Stelara (IV)	CBC, PPD				
☐ 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	CBC, Anti-AChR Antibody Positive, No Live Vaccines During Therapy				
□ Xolair	Asthma - Baseline Serum IgE, FEV1, Peak Flow, Other Pulmonary Function Test Chronic Idiopathic Urticaria – None				