RHEUMATOLOGY INFUSION REFERRAL FORM (PAGE 1 OF 2)

PHONE 855.896.9254 | **FAX** 855.370.0086



Remove above portion before faxing. Please complete the prescription form in its entirety and fax with secure cover sheet to the number above.

PATIENT INFORMATION Last Name Address Email Primary Contact Method (chec	First Name	DO	В		Gender □ M						
Address Email	First Name	DO	7B	(-onder M						
Email					Gerider 🗆 M	⊔F L	ast 4 SSN		Primary L	anguage	
				City				State		ZIP	
Primary Contact Method (chec		e Phone			Work Pho				Cell P		
	ck one)	☐ Home Phone	☐ Work Phone	e 🗆 Tex	t 🗆 Email	□ Prima	ry Caregive	r DO NO	T CONTACT		
Primary Caregiver/Alt Contact	Name (If applicable)		Alt Conta	ct Email					Alt Conta	act Phone	
PRESCRIBER INFORMA	TION										
Name of Contact Sending Refe	erral		Title			Preferre	d Contact N	1ethod (check	one) 🗆 E	mail 🗆 F	Phone 🗆 Fax
Referral Contact Email					Office Phone	•		Off	fice Fax		
Practice / Facility Name					Prescriber Na	ame / Spe	cialty				
Address				Cit	ty				State		ZIP
	* Pleas	e include a	copy of the	e front	t and bac	k of ins	surance	card *			
CLINICAL INFORMATIO	N - Please include	applicable clii	nical chart r	notes							
Patient New to Therapy 🗆 Naïv	ve/New Start	/ Restart □ Exis	ting Treatment				-	Therapy Start D	Date		
Sample/Starter Provided? ☐ N	lo ☐ Yes, Provide Qty:	Date Provi	ded:	Pa	tient Height (d	cm/in):	Weig	ht (kg/lbs):	Da	ate Obtain	ed:
Therapies Tried and Failed (ple	ease list medications)										
Other/Concomitant Medication	ns (please list)										
<u> </u>	Allergies (please list)										
	☐ Prescriber's Office ☐ □	Other (please list)									
ICD-10 Code	er forms of systemic lupus	s erythematosus s erythematosus, ı	unspecified		□ M06.8	89 Other		arthritis with rh eumatoid arthr s			nultiple sites
☐ M32.9 Othe		us, organ or system	m involvement :	unspecific				-			
☐ M32.9 Othe ☐ M32.10 Sys ☐ M32.19 Oth ☐ M05.79 Rh	stemic lupus erythematos ner organ or system involv neumatoid arthritis with rh	ement in systemic	c lupis erythema	atosus	☐ Other	r					
☐ M32.9 Othe ☐ M32.10 Sys ☐ M32.19 Oth ☐ M05.79 Rho or systems	stemic lupus erythematos ner organ or system involv neumatoid arthritis with rh s involvement	rement in systemic eumatoid factor o	c lupis erythema of multiple sites	atosus w/o orga	☐ Other	r					
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□ M32.9 Othe □ M32.10 Sys □ M32.10 Sys □ M32.19 Oth □ M05.79 Rh or systems PRESCRIPTION INFORM In order for a brand name p or your state-specific require	stemic lupus erythematosi ner organ or system involv leumatoid arthritis with rh is involvement MATION - Please Es product to be dispense	cribe if required, the prescribe is substitutions.	red by state or must handw This form is	atosus w/o orga law vrite "Bri	□ Other	ary" or "E					REFILLS
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Note: The information contained in this document will become a legal prescription. Prescriber is to comply with his/her state specific Pharmacy and Medical Board guidelines such as e-prescribing, state specific prescription form, fax language, number of prescriptions allowed on a single prescription form, etc. If more than one page is required, make additional copies. Non-compliance with state specific requirements could result in outreach to the prescriber.

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PATIENT INFORMATI	ON	TZIJM)	BE FILLED ON	T TO PROCESS PAGES TOGETHER)		
Patient Last Name		(MUS1			DOB	
	e product	to be dispensed, the	prescriber must h	state law nandwrite "Brand Necessary" or "Brand Medically Necessary," rm is not a valid prescription form for writing controlled med		
MEDICATION	ROUTE	DOSE/STRENGTH		DIRECTIONS	QTY	REFILLS
□ Saphnelo (anifrolumab)	□IV	□ 300 mg/2 mL Vial		□ To be infused over 30 minutes every 4 weeks vi pump with 0.2 or 0.22 micron filter. Upon completion of the infusion, flush infusion set with 25 mL of 0.9 Sodium Chloride Injection, USP. Prior to initiating therapy, is patient positive for autoantibodies relevant to SLE (e.g., ANA, anti-ds DNA, anti-Sm)? □ Yes □ No Positive ANA or anti-dsDNA test? □ Yes (Date of Test:) □ No	1 month 3 months	□1 year □
□ Simponi Aria (golimumab)	□IV	Starting dose 2 mg/kgmg IV at week 0, 4 and every 8 weeks OtherMaintenance Dose 2 mg/kgmg IV every 8 weeks Other		□ Infuse diluted solution over a period of 30 minutes	1 month 3 months	□1 year □
☐ Vascular Access Method	□ peri	pheral 🗆 central	□ other:			
☐ Normal Saline ☐ D5W	□IV	□ 3 mL □ 5 mL		☐ Before and after infusion ☐	□ 1 month □ 3 months	□ 1 year
☐ Heparin 10 units/mL ☐ Heparin 100 units/mL	□IV	3 mL		☐ After infusion ☐	□ 1 month □ 3 months	□ 1 year □
☐ Diphenhydramine	□ PO □ IV □ IM	☐ 25 mg ☐ 50 mg		☐ After infusion ☐ PRN Allergic Reaction:	☐ With each infusion	□ 1 year □
□ Famotidine	□IV	□ 20 mg IVP □ 40 mg IVP		□ Pre-Med:		□1 year □
☐ Methylprednisolone	□IV	□ 40 mg IVP □ 125 mg IVP □		□ Pre-Med:		□1 year
☐ Acetaminophen	□РО		□ 500 mg □ 1 gm	□ Pre-Med:	☐ With each infusion	□ 1 year
☐ Epinephrine	□ IM □ SQ	☐ Adult 1:1000, 0.3 m ☐ Peds 1:2000, 0.3 ml (15-30kg/33-66lbs)		□ PRN Anaphylaxis □ Repeating Dose:	□ Once	□1 year
☐ Other:	 					
Prescriber Signature			Date	Supervising Physician Signature (where required by state law)	NPI #	Date
DAW (Dispense as Written)			Date	Brand Necessary (must handwrite)		

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