RHEUMATOLOGY INFUSION Referral Form (Page 1 of 2)

PHONE 515.225.2930 | **FAX** 515.559.2495



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

First National First	Home Phone	DOB	City	Gender □I	м 🗆 ғ	Last 4 S	SSN		Primary	Language		
check one) □ Cell I act Name (If applica	Home Phone			Gender □ I	M 🗆 F	Last 4 S	SSN		Primary			
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			P	Prescriber N	ame/S _l	pecialty		l				
Practice/Facility Name Address										z	IP	
	* Please include	a copy of the	front a	nd back o	f insui	rance ca	rd. *					
– Please include	applicable clinica	I chart notes.										
Naïve/New Start 🗆	Therapy Restart 🗆 Ex	cisting Treatment					The	erapy Start	Date			
□ No □ Yes, Provide	Qty: Date Prov	/ided:	Patie	Patient Height (cm/in): Weight (kg/lbs):					Dat	ate Obtained:		
(please list medicati	ons)											
tions (please list)												
g Allergies (please li	st)											
☐ Prescriber's Offi	ce 🗆 Other (please lis	t)										
ther forms of system	nic lupus erythematosu	s		ПМО	5.89 O	ther rheur	natoid aı	thritis wit	h rheumate	oid factor of n	ultiple sites	
ther forms of system systemic lupus eryth Other organ or syste Rheumatoid arthritis	nic lupus erythematosu ematosus, organ or sys m involvement in syster	s, unspecified tem involvement mic lupis erythem	atosus	☐ M0 fied ☐ M0 ☐ Otl	6.89 O	ther speci	fied rheu				·	
e product to be di	spensed, the prescrib	oer must handw								ons.		
ROUTE	DOSE/STRENGTH		DIREC	CTIONS						QTY	REFILLS	
□IV	□10 mg/kg		□ 10 r Maint	□ 10 mg/kg IV at weeks 0, 2, 4 and then every weeks Maintenance Dose			eeks	□1 month □3 months	□1 year			
□IV		-	□ 400 week Main	Starting Dose 400 mg (given as two 200 mg subcutaneous injections) at weeks 0, 2, 4 Maintenance Dose 200 mg subcutaneous injection every other week 0 other				□1 month □3 months	□1 year □			
□IV	□ 500 mg Orencia □ 750 mg Orencia □ 1000 mg Orencia		□ Info	use over 30	minute	es				□1 month □3 months	□1 year	
□IV	☐ 3 mg/kgmg IV at ☐ Other Maintenance Dose	t weeks 0, 2, 6	□То	be infused (over a p	period NO	T less tha	n 2 hours		□1 month □3 months	□1 year	
	Date		ıpervisin	g Physician	Signa	ture (wher	e require	_ ed by state	e law) N	 PI#	 Date	
	Naïve/New Start No Yes, Provide (please list medications (please list) g Allergies (please li Prescriber's Offither forms of system ther forms of system ther forms of system is lupus erythother organ or system in the sinvolvement tion – Please Estem product to be divired language to ROUTE NOUTE	Please include applicable clinica Naïve/New Start □ Therapy Restart □ Exit	— Please include applicable clinical chart notes. Naïve/New Start □ Therapy Restart □ Existing Treatment □ No □ Yes, Provide Qty: □ Date Provided: (please list medications) Itions (please list) □ □ Prescriber's Office □ Other (please list) Ither forms of systemic lupus erythematosus, unspecified systemic lupus erythematosus, unspecified byther organ or system involvement in systemic lupis erythematory and in systemic lupis	Please include applicable clinical chart notes. Naïve/New Start	* Please include a copy of the front and back o - Please include applicable clinical chart notes. Naïve/New Start	Please include a copy of the front and back of insular places include applicable clinical chart notes. Naive/New Start □ Therapy Restart □ Existing Treatment □ No □ Yes, Provide Qty: □ Date Provided: □ Patient Height (cm/in) (please list medications) titions (please list) □ Prescriber's Office □ Other (please list) □ M05.89 O M06.89 O M06.89 O M06.89 O M06.89 O M06.89 O M06.90 Rheumatoid arthritis with rheumatoid factor of multiple sites w/o organ The product to system involvement in systemic lupis erythematosus Prescriber must handwrite "Brand Necessary" wired language to prohibit substitutions. This form is not a valid prescription for the prescriber must handwrite "Brand Necessary" wired language to prohibit substitutions. This form is not a valid prescription for the prescriber must handwrite "Brand Necessary" wired language to prohibit substitutions. This form is not a valid prescription for the prescriber must handwrite "Brand Necessary" wired language to prohibit substitutions. This form is not a valid prescription for the prescriber must handwrite "Brand Necessary" wired language to prohibit substitutions. This form is not a valid prescription for the prescriber must handwrite "Brand Necessary" wired language to prohibit substitutions. This form is not a valid prescription for Maintenance Dose □ 10 mg/kg IV every □ Starting Dose □ 10 mg/kg IV every □	* Please include a copy of the front and back of insurance call - Please include applicable clinical chart notes. Naïve/New Start	Please include a poplicable clinical chart notes. Naive/New Start □ Therapy Restart □ Existing Treatment □ The □ No □ Yes, Provide Qty: □ Date Provided: □ Patient Height (cm/in): Weight (liplease list medications) tions (please list) If patient Height (cm/in): Weight (liplease list) If prescriber's Office □ Other (please list) If patient Height (cm/in): Weight (liplease list) If patient Height (cm/in): W	Please include applicable clinical chart notes. Naive/New Start	Please include a copy of the front and back of insurance card. * Please include applicable clinical chart notes. Naive/New Start	Please include a poplicable clinical chart notes. Naive/New Start □ Prescriber □ Patient Height (cm/in): Weight (kg/lbs): Date Obtained: Patient Height (cm/in): Weight (kg/lbs): Patient Height (cm/in): Weight (kg/lbs): Patient Height (cm/in): Weight (kg/lbs):	

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. Confidentiality Statement: This message is intended only for the individual or entity to which it is addressed. It may contain information which may be proprietary and confidential. It may also contain privileged, confidential information, which is exempt from disclosure under applicable laws, including the Health Insurance Portability and Accountability Act (HIPAA). If you are not the intended recipient, please note that you are strictly prohibited from disseminating or distributing this information (other than to the intended recipient) or copying this information. If you received this communication in error, please notify the sender immediately at the address and telephone number set forth herein and obtain instructions as to proper destruction of the transmitted material.

Brand Necessary (must handwrite)

Date

DAW (Dispense as Written)

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Patient Information		Must Be Filled C						
Patient Last Name			Patient First N	ame	DOB			
	e product to be di	spensed, the presci	riber must handv	rite "Brand Necessary" or "Brand Medically Nece ot a valid prescription form for writing controlled		ns.		
MEDICATION	ROUTE	DOSE/STRENGTH		DIRECTIONS		QTY	REFILLS	
□ Rituxan (rituximab)	□IV	□ 1000 mg IV on day 0, day 14 and then repeat the course everyweeks □ 375 mg/m2 IV every 4 weeks □ Other		☐ Infuse as directed	□1 month □3 months	□1 year		
□ Saphnelo (anifrolumab)	□IV	□ 300 mg/2 mL Vial		□ To be infused over 30 minutes every 4 weeks IV p 0.2 or 0.22 micron filter. Upon completion of the infi infusion set with 25 mL of 0.9 Sodium Chloride Inject Prior to initiating therapy, is patient positive for autoantibodies relevant to SLE (e.g., ANA, anti-ds Di anti-Sm)? □ Yes □ No Positive ANA or anti-ds DNA test? □ Yes (Date of test:) □ No	□1 month □3 months	□1 year □		
□ Simponi Aria (golimumab)	□IV	Starting dose 2 mg/kgmg IV every 8 weeks Other Maintenance Dose 2 mg/kgmg IV 0	every 8 weeks	☐ Infuse diluted solution over a period of 30 minute	□1 month □3 months	□1 year □		
☐ Vascular Access Method	□ peripheral	□ 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 □ 5 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:	□ Pre-Med:			
☐ Methylprednisolone	□IV	☐ 40 mg IVP ☐ 125 mg IVP ☐		□ Pre-Med:		□ 1 year		
☐ Acetaminophen	□РО	□ 325 mg □ 650 mg	□ 500 mg □1 gm	□ Pre-Med:		□ With each infusion	□ 1 year	
☐ Epinephrine	□IM □SQ	☐ Adult 1:1000, 0.3 m		□ PRN Anaphylaxis) □ Repeating Dose:	_	Once	□1 year	
☐ Other:								
Prescriber Signature				pervising Physician Signature (where required by state	e law) NF		Date	
DAW (Dispense as Written)		Date	Br	and Necessary (must handwrite)				

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