

**1 PATIENT INFORMATION:**

Name: \_\_\_\_\_  
 Address: \_\_\_\_\_  
 City: \_\_\_\_\_ State: \_\_\_\_\_ Zip: \_\_\_\_\_  
 Phone: \_\_\_\_\_ Alt. Phone: \_\_\_\_\_  
 Email: \_\_\_\_\_  
 DOB: \_\_\_\_\_ Gender:  M  F Caregiver: \_\_\_\_\_  
 Height: \_\_\_\_\_ Weight: \_\_\_\_\_ Allergies: \_\_\_\_\_

**2 PRESCRIBER INFORMATION:**

Name: \_\_\_\_\_  
 Address: \_\_\_\_\_  
 City: \_\_\_\_\_ State: \_\_\_\_\_ Zip: \_\_\_\_\_  
 Phone: \_\_\_\_\_ Fax: \_\_\_\_\_  
 NPI: \_\_\_\_\_ DEA: \_\_\_\_\_  
 Tax I.D.: \_\_\_\_\_  
 Office Contact: \_\_\_\_\_ Phone: \_\_\_\_\_

**3 STATEMENT OF MEDICAL NECESSITY:** (Please Attach All Medical Documentation)

Date of Diagnosis: \_\_\_\_\_  
 ICD-10: \_\_\_\_\_ Other: \_\_\_\_\_  
 TB Test:  Positive  Negative Date: \_\_\_\_\_  
 LFT: ALT: \_\_\_\_\_ AST: \_\_\_\_\_ Date: \_\_\_\_\_  
 Assessment:  Moderate  Mod to Severe  Severe  
 \_\_\_\_\_% BSA affected  
 Scalp  Face  Chest  Arms  Hands  Nails  
 Back  Groin  Buttocks  Legs  Other: \_\_\_\_\_

Patient also taking Methotrexate?  Yes  No  
 Serious or active infection present?  Yes  No  
 Hep B ruled out or treatment started?  Yes  No  
 Does patient have latex allergy?  Yes  No  
 Does patient have joint involvement?  Yes  No  
 If yes, please indicate affected joint(s): \_\_\_\_\_  
**If Prior Authorization is denied, recommended  
 formulary alternatives will be provided to the prescriber  
 based upon the patient's insurance coverage.**

**Prior Failed Treatments:**

Topicals \_\_\_\_\_  
 Methotrexate \_\_\_\_\_  
 Oral Meds \_\_\_\_\_  
 Biologics \_\_\_\_\_  
 UVA  UVB \_\_\_\_\_  
 Others \_\_\_\_\_

**4 INJECTION TRAINING:**  To be Administered by a Healthcare Provider  Pharmacist to Provide Training  Patient Trained in MD Office  Manufacturer Nurse Support

**5 PICK UP OR DELIVERY:**  Delivery to Patient's Home  Delivery to Physician's Office  Pharmacy to Coordinate

**6 INSURANCE INFORMATION:** Please Include Front and Back Copies of Pharmacy and Medical Card

**PRESCRIPTION INFORMATION:** (Please be sure to choose both induction and maintenance dose where applicable)

Patient Name: \_\_\_\_\_ Patient's Date of Birth: \_\_\_\_\_

Medication	Dosage & Strength	Direction	QTY	Refills
<input type="checkbox"/> RASUVO®	Single-dose auto-injector prefilled syringe: <input type="checkbox"/> 7.5mg <input type="checkbox"/> 10mg <input type="checkbox"/> 12.5mg <input type="checkbox"/> 15mg <input type="checkbox"/> 17.5mg <input type="checkbox"/> 20mg <input type="checkbox"/> 22.5mg <input type="checkbox"/> 25mg <input type="checkbox"/> 27.5 mg <input type="checkbox"/> 30mg	<input type="checkbox"/> Inject _____ mg SC once weekly  *An initial test dose of 2.5 to 5 mg is recommended in patients with risk factors for hematologic toxicity or renal impairment*		
<input type="checkbox"/> SILIQ™	<input type="checkbox"/> 210mg/1.5ml Prefilled Syringe	<input type="checkbox"/> Induction Dose: Inject 210mg subcutaneously at Weeks 0, 1 and 2 <input type="checkbox"/> Maintenance Dose: Inject 210mg subcutaneously every 2 weeks	<input type="checkbox"/> 1 Months <input type="checkbox"/> 2 Months <input type="checkbox"/> 3 Months	
<input type="checkbox"/> SIMPONI® (for PsA)	<input type="checkbox"/> 50mg/0.5ml Smartject Injector <input type="checkbox"/> 50mg/0.5ml Prefilled Syringe	<input type="checkbox"/> Inject 50mg SC once a month	1	
<input type="checkbox"/> SKYRIZI™	<input type="checkbox"/> 75mg/0.83ml Prefilled Syringe <input type="checkbox"/> Yes or <input type="checkbox"/> No: SKYRIZI SELF-INJECTION: Healthcare provider certifies that patient has been trained and is eligible for self-injection	<input type="checkbox"/> Induction Dose: Inject 150mg (two 75mg injections) SC at weeks 0 and 4 <input type="checkbox"/> Maintenance: Inject 150mg (two 75mg injections) SC every 12 weeks thereafter	4 2	0
<input type="checkbox"/> STELARA®	<input type="checkbox"/> 45mg/0.5 mL Single-Dose Prefilled Syringe <input type="checkbox"/> 45mg/0.5 mL Solution in a Single-Dose Vial	<b>Plaque Psoriasis:</b> <input type="checkbox"/> Adult dosing (≤100 kg): Inject 45 mg SC initially and at 4 weeks, then every 12 weeks thereafter <input type="checkbox"/> Adult Dosing (>100 kg): Inject 90 mg SC initially and at 4 weeks, then every 12 weeks thereafter <b>Psoriatic Arthritis:</b> <input type="checkbox"/> Inject 45 mg SC initiation and at 4 weeks, then every 12 weeks thereafter		
<input type="checkbox"/> TALTZ®	<input type="checkbox"/> 80mg/ml Single-Dose Prefilled Autoinjector <input type="checkbox"/> 80mg/ml Single-Dose Prefilled Syringe	<b>Psoriatic Arthritis (PsA):</b> <input type="checkbox"/> Induction Dose: Inject 160 mg SC (two 80 mg injections) at week 0 <input type="checkbox"/> Maintenance: Inject 80 mg SC every 4 weeks thereafter <b>Plaque Psoriasis or PsA with Coexistent Moderate-to-Severe Plaque Psoriasis:</b> <input type="checkbox"/> Weeks 0-2: Inject 160mg SC (two 80mg injections) at weeks 0, then inject 80mg SC at week 2 <input type="checkbox"/> Weeks 4-10: Inject 80mg SC at week 4 and every 2 weeks thereafter through week 10 <input type="checkbox"/> Week 12 and onwards: Inject 80mg SC at week 12 and every 4 weeks thereafter	3 4	0 0
<input type="checkbox"/> TREMFYA™	<input type="checkbox"/> 100mg/ml Prefilled Syringe <input type="checkbox"/> 100mg/ml One- Press Patient Controlled Injector	<input type="checkbox"/> Induction Dose: Inject 100mg/ml SC at weeks 0 and 4 <input type="checkbox"/> Maintenance: Inject 100mg/ml SC every 8 weeks thereafter	2 1	0
<input type="checkbox"/> XELJANZ®	<input type="checkbox"/> 5mg Tablet	<input type="checkbox"/> Take one tablet by mouth twice daily in combination with a nonbiologic DMARD	60	
<input type="checkbox"/> XELJANZ® XR	<input type="checkbox"/> 11mg Tablet	<input type="checkbox"/> Take one tablet by mouth once daily in combination with a nonbiologic DMARD	30	
<input type="checkbox"/>				

*Cimzia®, Cosentyx®, Enbrel®, Humira®, Orencia™ and Otezla® are listed alphabetically on respective enrollment forms.*

**PRESCRIBER SIGNATURE:** I authorize pharmacy to act as my designee for initiating and coordinating insurance prior authorizations, nursing services and patient assistance programs.

Signature: \_\_\_\_\_ Date: \_\_\_\_\_ Signature: \_\_\_\_\_ Date: \_\_\_\_\_  
 Substitution Permitted Dispense As Written

Prior authorization approval and insurance benefits will be determined by the payor based upon the patient's eligibility, medical necessity, and the terms of the patient's coverage, among other things. Participation in this program is not a guarantee of prior authorization or of payment. If Prescribing SILIQ™: I certify that I made the prescribing decisions indicated above based on my own independent medical judgment regarding what is in the best interest of the patient and that I have reviewed the current SILIQ Prescribing Information. By signing above, I confirm that: (1) I am certified as a healthcare provider under the SILIQ REMS Program, (2) I have counseled the patient on the risks of suicidal ideation and behavior that may occur with SILIQ, (3) the patient has signed the SILIQ REMS Patient-Prescriber Agreement, (4) I have given the patient the SILIQ REMS patient wallet card, and (5) I have enrolled the patient in the SILIQ REMS Program. I authorize SILIQ Solutions to act on my behalf to transmit this prescription to the appropriate qualified pharmacy designated above by the patient or the patient's plan.