

PATIENT INFORMATION

Patient Name: _____ DOB: _____ Phone: _____ Sex: M F Ht: _____ Wt: _____ lbs kg
 Primary Language: _____ Allergies: _____
 Patient Preferred Location: _____

DIAGNOSIS & CLINICAL INFORMATION

<ICD 10 CODE REQUIRED>

ICD 10 Code (PROVIDE COMPLETE CODE)

M32.1 _____ Systemic lupus erythematosus with organ or system involvement

Other: _____

Prescribing information

Limitations of Use: The efficacy of Saphnelo has not been evaluated in patients with severe active lupus nephritis or severe active central nervous system lupus. Saphnelo has not been studied in combination with other biologics therapies. Therefore, the use of Saphnelo is not recommended for use in combination with biologic therapies.

Evaluation of Immunizations: Should be completed prior to, and live vaccines should not be given for 30 days before or concurrently with Saphnelo.

Missed Dose: Administer as soon as possible but maintain at least 14 days between infusions.

REQUIRED: Demographics & Most Recent: H&P, clinical notes, & medication list. Supporting clinical notes to include any past tried and/or failed therapies, intolerance, outcomes, or contraindications to conventional therapy.
LAB RESULTS: Lab testing documenting the presence of autoantibodies (i.e. ANA, Anti-dsDNA, Anti-Sm, Anti-Ro/SSA, Anti-La/SSB)

PRESCRIPTION

Pre-Medications

Acetaminophen: 650 mg PO
 Cetirizine: 10 mg PO
 Diphenhydramine: 25mg PO

Diphenhydramine: 25mg IVP
 Famotidine: 20 mg PO
 Methylprednisolone: 125 mg SIVP
 Other: _____

SAPHNELO (anifrolumab-fnia)

Loading Dose

IV: Infuse 300 mg in 100 mL of 0.9% Sodium Chloride over 30 minutes using a 0.2-micron filter every 4 weeks for one year
 After the infusion, flush with 25 mL of 0.9% Sodium Chloride

Is the patient on any other disease modifying therapy? Yes No
 If yes, please note therapy and last dose: _____

Post Treatment Observations: The patient is observed for 30 minutes following the first administration.

Adverse Events: In the event of an adverse reaction occurring at a Medix Infusion suite, utilize the Medix Infusion adverse reactions protocol.

Comments: _____

PRESCRIBER INFORMATION

Prescriber Name: _____ Signature: _____
 Date: _____ NPI #: _____ Specialty: _____
 Supervising Physician: _____ (If Applicable)
 Address: _____ City: _____ State: _____ Zip: _____
 Contact Name: _____ Phone: _____ Fax: _____ Email: _____