

OPTIMA HEALTH PLAN

PHARMACY/MEDICAL PRIOR AUTHORIZATION REQUEST

Directions: The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; fax to 1-844-202-5034. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. Incomplete form will delay authorization process.

Drug Requested: **Ilaris® (canakinumab) (J0638) (Medical)**

{Tumor Necrosis Factor Receptor Associated Periodic Syndrome (**TRAPS**), Hyperimmunoglobulin D Syndrome (**HIDS**)/Mevalonate Kinase Deficiency (**MKD**), and Familial Mediterranean Fever (**FMF**)}

DRUG INFORMATION: Complete all information below. If incomplete, authorization process will be delayed.

Drug Form/Strength/Quantity: _____

Dosing Schedule: _____ Length of Therapy: _____

Diagnosis: _____ ICD Code: _____

CLINICAL CRITERIA: Progress notes/chart notes **MUST** be submitted to support lab values and diagnosis. Applicable boxes below **must** be checked to qualify. If incomplete, authorization process will be delayed.

1st Approval: 6 months

Age: ≥ 2 years old Weight kg: _____

Please check all that apply: (All boxes **must** be checked) **1st Approval: 6 months**

Age: ≥ 2 years old Weight kg: _____

Diagnosis:

- Tumor Necrosis Factor Receptor Associated Periodic Syndrome (TRAPS)**
 - Chart notes documenting six (6) flares within a 12 month time frame.
 - Labs document CRP >10mg/L
- Hyperimmunoglobulin D Syndrome (HIDS)/Mevalonate Kinase Deficiency (MKD)**
 - Test result submitted genetic MVK/enzymatic (MKD)
 - History \geq three (3) febrile acute flares within a 6 month period and not receiving prophylactic treatment: YES NO
 - \geq CRP 10 mg/L
- Familial Mediterranean Fever (FMF)**
 - Documented a trial and failure colchicine 1.5-2.0mg/day
 - Type I phenotype
 - Currently active disease the following will meet the criteria:
 - One (1) flare per month (chart notes document five months of flare)
 - \geq CRP 10 mg/L

Reauth Approval 1 year: Please submit current progress notes that document CRP and symptoms.

(signature on next page)

Medication being provided by a Specialty Pharmacy: Briova SpecialtyRx

****Use of samples to initiate therapy does not meet step edit/preauthorization criteria.****

Previous therapies will be verified through pharmacy paid claims or submitted chart notes.

Patient Name: _____

Member Optima #: _____ Date of Birth: _____

Prescriber Name: _____

Prescriber Signature: _____ Date: _____

Office Contact Name: _____

Phone Number: _____ Fax Number: _____

DEA OR NPI #: _____

*Approved by Pharmacy and Therapeutics Committee: 11/18/16

REVISED/UPDATED: 3/28/2017; 4/4/2017; 9/14/2017; 10/4/2017.