

July 12th, 2010

SUBJECT: Prior Authorization Effective July 21, 2010

Ilaris® (canakinumab)

Authorization Criteria

- 1. FDA approved indication of Cryopyrin-Associated Periodic Syndromes (CAPS) verified by genetic testing. This includes Familial Cold Auto-inflammatory Syndrome (FCAS) and Muckle-Wells Syndrome (MWS) in adults and children 4 and older.
- 2. The member should not be using a tumor necrosis factor blocking agent (e.g. adalimumab, etanercept, and infliximab) or anakinra
- 3. Should not be initiated in patients with active or chronic infection including hepatitis B, hepatitis C, human immunodeficiency virus, or tuberculosis.
- 4. Dosing should not be more often than once every 8 weeks.
- 5. Approved dosing schedule based on weight:
 - a. Body weight >40 kg: 150mg
 - b. Body weight 15 kg 40 kg: 2mg/kg. If inadequate response, may be increased to 3mg/kg
- 6. Approval period is for one year.

Requip XL™ (ropinirole) and Mirapex ER™ (pramipexole)

Authorization Criteria

- 1. Diagnosis of Parkinson's Disease, and
- 2. Clinically significant reason why the immediate release products cannot be used

Lovaza® (Omega-3-Acid Ethyl Esters)

Authorization Criteria

- 1. Laboratory documentation of severe hypertriglyceridemia (fasting triglycerides ≥500 mg/dL).
- 2. Previous failure with both nicotinic acid and fibric acid medications.

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PA Criteria: www.okhca.org/providers/rx/pa PA forms: www.okhca.org/rx-forms