

Reference number
1800-A

SPECIALTY GUIDELINE MANAGEMENT

ARCALYST (rilonacept)

POLICY

I. INDICATIONS

The indications below including FDA-approved indications and compendial uses are considered a covered benefit provided that all the approval criteria are met and the member has no exclusions to the prescribed therapy.

FDA-Approved Indications

Arcalyst is indicated for:

- A. Treatment of Cryopyrin Associated Periodic Syndromes (CAPS), including Familial Cold Auto-inflammatory Syndrome (FCAS) and Muckle-Wells Syndrome (MWS) in adults and children 12 years of age and older.
- B. Maintenance of remission of Deficiency of Interleukin-1 Receptor Antagonist (DIRA) in adults and pediatric patients weighing at least 10 kg.
- C. Treatment of recurrent pericarditis (RP) and reduction in risk of recurrence in adults and pediatric patients 12 years and older.

All other indications are considered experimental/investigational and not medically necessary.

II. DOCUMENTATION

Submission of the following information is necessary to initiate the prior authorization review:

- A. Deficiency of interleukin-1 receptor antagonist (DIRA) initial requests: *IL1RN* mutation status
- B. Recurrent pericarditis:
 1. Initial requests: Chart notes, medical record documentation, or claims history supporting previous medications tried, including response to therapy.
 2. Continuation requests: Chart notes or medical record documentation supporting positive clinical response.

III. PRESCRIBER SPECIALTIES

This medication must be prescribed by or in consultation with one of the following:

- A. Cryopyrin associated periodic syndromes (CAPS) and deficiency of interleukin-1 receptor antagonist (DIRA): rheumatologist or immunologist
- B. Recurrent pericarditis: cardiologist, rheumatologist, or immunologist

IV. CRITERIA FOR INITIAL APPROVAL

A. Cryopyrin-associated periodic syndromes (CAPS)

Arcalyst 1800-A SGM P2022a

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Authorization of 12 months may be granted for members 12 years and older for the treatment of CAPS when both of the following criteria are met:

1. Member has a diagnosis of familial cold auto-inflammatory syndrome (FCAS) with classic signs and symptoms (i.e., recurrent, intermittent fever and rash that were often exacerbated by exposure to generalized cool ambient temperature) or Muckle-Wells syndrome (MWS) with classic signs and symptoms (i.e., chronic fever and rash of waxing and waning intensity, sometimes exacerbated by exposure to generalized cool ambient temperature).
2. Member has functional impairment limiting the activities of daily living.

B. Deficiency of interleukin-1 receptor antagonist (DIRA)

Authorization of 12 months may be granted for members weighing at least 10 kg for the treatment of DIRA when both of the following criteria are met:

1. Member has loss-of-function *IL1RN* mutations.
2. Arcalyst will be used for maintenance of remission following treatment with Kineret (anakinra).

C. Recurrent pericarditis

Authorization of 12 months may be granted for members 12 years and older for the treatment of recurrent pericarditis when both of the following criteria are met:

1. Member has had at least two episodes of pericarditis.
2. Member has failed at least two agents of standard therapy (e.g., colchicine, non-steroidal anti-inflammatory drugs [NSAIDs], corticosteroids).

V. CONTINUATION OF THERAPY

A. Cryopyrin-associated periodic syndromes (CAPS)

Authorization of 12 months may be granted for all members 12 years and older (including new members) who are using the requested medication for CAPS who achieve or maintain positive clinical response as evidenced by low disease activity or improvement in signs and symptoms of the condition.

B. Deficiency of interleukin-1 receptor antagonist (DIRA)

Authorization of 12 months may be granted for all members weighing at least 10 kg (including new members) who are using the requested medication for DIRA and who achieve or maintain positive clinical response as evidenced by low disease activity or improvement in signs and symptoms of the condition.

C. Recurrent pericarditis

Authorization of 12 months may be granted for all members 12 years and older (including new members) who are using the requested medication for recurrent pericarditis and who achieve or maintain a positive clinical response as evidenced by decreased recurrence of pericarditis or improvement in signs and symptoms of the condition when there is improvement in any of the following:

1. Pericarditic chest pain
2. Pericardial rubs
3. Electrocardiogram (ECG)
4. Pericardial effusion
5. C-reactive protein (CRP)

VI. OTHER

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For all indications: Member has had a documented negative tuberculosis (TB) test (which can include a tuberculosis skin test [PPD], an interferon-release assay [IGRA], or a chest x-ray)* within 6 months of initiating therapy for persons who are naïve to biologic drugs or targeted synthetic drugs associated with an increased risk of TB.

* If the screening testing for TB is positive, there must be further testing to confirm there is no active disease. Do not administer the requested medication to members with active TB infection. If there is latent disease, TB treatment must be started before initiation of the requested medication.

For all indications: Member cannot use the requested medication concomitantly with any other biologic drug or targeted synthetic drug.

VII. REFERENCES

1. Arcalyst [package insert]. London, UK: Kiniksa Pharmaceuticals (UK), Ltd.; May 2021.
2. Hoffman HM, Throne ML, Amar NJ, et al. Efficacy and safety of riloncept (interleukin-1 trap) in patients with cryopyrin-associated periodic syndromes. Results from two sequential placebo-controlled studies. *Arthritis Rheum.* 2008;58(8):2443-52.
3. Testing for TB Infection. Centers for Disease Control and Prevention. Retrieved on June 15, 2022 from: <https://www.cdc.gov/tb/topic/testing/tbtesttypes.htm>.
4. Adler Y, Charron P, Imazio M, et al. 2015 ESC Guidelines for the diagnosis and management of pericardial diseases: The Task Force for the Diagnosis and Management of Pericardial Diseases of the European Society of Cardiology (ESC) Endorsed by: The European Association for Cardio-Thoracic Surgery (EACTS). *Eur Heart J.* 2015 Nov 7;36(42):2921-64.
5. Chiabrando JG, Bonaventura A, Vecchié, et al. Management of acute and recurrent pericarditis: JACC State-of-the-art review. *J Am Coll Cardiol.* 2020;75(1):76-92.
6. Klein AL, Imazio M, Cremer P, et al. Phase 3 trial of interleukin-1 trap riloncept in recurrent pericarditis. *N Engl J Med.* 2021;384(1):31-41.