



Drug Name: Epogen, Procrit (epoetin alfa)

Line of Business: Exchange

Revised Date: 12/2018

Drug Name:	Epogen, Procrit (epoetin alfa)
Exclusion Criteria:	<ul style="list-style-type: none"> • In patients with cancer receiving hormonal agents, biologic products, or radiotherapy, unless also receiving concomitant myelosuppressive chemotherapy. • In patients with cancer receiving myelosuppressive chemotherapy when the anticipated outcome is cure. • In patients scheduled for surgery who are willing to donate autologous blood. • In patients undergoing cardiac or vascular surgery. • As a substitute for RBC transfusions in patients who require immediate correction of anemia
Required Medical Information:	<ul style="list-style-type: none"> • Anemia Due to CKD Authorization of 12 weeks may be granted for members with pretreatment hemoglobin < 10 g/dL. • Anemia Due to Myelosuppressive Chemotherapy Authorization of 12 weeks may be granted for members with nonmyeloid malignancy who meet ALL of the following criteria: <ol style="list-style-type: none"> 1. The intent of chemotherapy is non-curative 2. Pretreatment hemoglobin < 10 g/dL • Anemia in MDS <ul style="list-style-type: none"> ○ Authorization of 12 weeks may be granted for members with pretreatment hemoglobin < 10 g/dL. • Reduction of Allogeneic Red Blood Cell Transfusion in Patients Undergoing Elective, Noncardiac, Nonvascular Surgery <ul style="list-style-type: none"> ○ Authorization of 12 weeks may be granted for members scheduled to have an elective, noncardiac, nonvascular surgery when the pretreatment hemoglobin is > 10 to ≤ 13 g/dL. • Anemia in Congestive Heart Failure (CHF) <ul style="list-style-type: none"> ○ Authorization of 12 weeks may be granted for members with pretreatment hemoglobin < 9 g/dL. • Anemia in Rheumatoid Arthritis (RA) <ul style="list-style-type: none"> ○ Authorization of 12 weeks may be granted for members with

pretreatment hemoglobin < 10 g/dL.

- **Anemia Due to Hepatitis C Treatment**
 - Authorization of 12 weeks may be granted for members with pretreatment hemoglobin < 10 g/dL who are receiving ribavirin in combination with either interferon alfa or peginterferon alfa.
- **Anemia Due to Zidovudine in HIV-infected Patients**
 - Authorization of 12 weeks may be granted for members currently receiving zidovudine with pretreatment hemoglobin < 10 g/dL.
- **Anemia in Members Whose Religious Beliefs Forbid Blood Transfusions**
 - Authorization of 12 weeks may be granted for members with pretreatment hemoglobin < 10 g/dL.
- **Anemia in Primary Myelofibrosis (MF), Post-polycythemia Vera MF, and Post-Essential Thrombocythemia MF**
 - Authorization of 12 weeks may be granted for members who meet ALL of the following criteria:
 - Member has symptomatic anemia
 - Pretreatment hemoglobin < 10 g/dL
 - Pretreatment serum erythropoietin level < 500 mU/mL

Renewal Criteria

For all indications below: all members (including new members) requesting authorization for continuation of therapy after at least 12 weeks of ESA treatment must show a response with a rise in hemoglobin of ≥ 1 g/dL. Members who completed less than 12 weeks of ESA treatment and have not yet responded with a rise in hemoglobin of ≥ 1 g/dL may be granted authorization of up to 12 weeks to allow for sufficient time to demonstrate a response.

- **Anemia Due to CKD**
 - Authorization of 12 weeks may be granted for continuation of therapy when the current hemoglobin is ≤ 12 g/dL.
- **Anemia Due to Myelosuppressive Chemotherapy**
 - Authorization of 12 weeks may be granted for the continuation of therapy in members with nonmyeloid malignancy who meet BOTH of the following criteria:
 - The intent of chemotherapy is non-curative
 - Current hemoglobin is < 11 g/dL

- **Anemia in MDS**
 - Authorization of 12 weeks may be granted for continuation of treatment when the current hemoglobin is ≤ 12 g/dL.
- **Anemia in CHF, RA**
 - Authorization of 12 weeks may be granted for continuation of treatment when the current hemoglobin is ≤ 12 g/dL.
- **Anemia Due to Hepatitis C Treatment**
 - Authorization of 12 weeks may be granted for continuation of treatment when the member meets ALL of the following criteria:
 - The member is receiving ribavirin in combination with either interferon alfa or peginterferon alfa
 - The current hemoglobin is ≤ 12 g/dL.
- **Anemia Due to Zidovudine in HIV-infected Patients**
 - Authorization of 12 weeks may be granted for continuation of therapy in members receiving zidovudine when the current hemoglobin is ≤ 12 g/dL.
- **Anemia in Members Whose Religious Beliefs Forbid Blood Transfusions**
 - Authorization of 12 weeks may be granted for continuation of treatment when the current hemoglobin is ≤ 12 g/dL.
- **Anemia in Primary Myelofibrosis, Post-polycythemia Vera Myelofibrosis, and Post-Essential Thrombocythemia Myelofibrosis**
 - Authorization of 12 weeks may be granted for continuation of treatment when the current hemoglobin is ≤ 12 g/dL.