

OPTIMA HEALTH PLAN

PHARMACY 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 1-800-750-9692. 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.*

This form is to be completed ONLY if the patient is self-administering.

The FDA has placed a Black Box Warning on all Erythropoietin Stimulating Agents (ESA).

Drug Requested (check one below):

- | | | |
|---|--|---|
| <input type="checkbox"/> Aranesp® (darbepoetin alfa) | <input type="checkbox"/> Epogen® (epoetin alfa) | <input type="checkbox"/> Procrit® (epoetin alfa) |
|---|--|---|

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

Drug Form/Strength: _____

Dosing Schedule: _____ Length of Therapy: _____

Diagnosis: _____ ICD Code, if applicable: _____

Procrit® is Preferred for Chronic renal Failure and Anemia in Cancer Patients

CLINICAL CRITERIA: ALL lines below must be completed to qualify. Authorization process will be delayed if incomplete. ALL lab results/tests MUST be submitted for review.

*Patient's most recent hemoglobin level. Hg = _____ *

Diagnosis: Anemia associated with (check **one** of the diagnoses below):

- | | |
|---|--|
| <ul style="list-style-type: none"><input type="checkbox"/> <u>Chronic Renal Failure</u><ul style="list-style-type: none"><input type="checkbox"/> Patient has tried and failed:<ul style="list-style-type: none"><input type="checkbox"/> Procrit®<input type="checkbox"/> <u>HIV/AIDS receiving zidovudine</u><ul style="list-style-type: none"><input type="checkbox"/> Endogenous erythropoietin <500mUnits/mL<input type="checkbox"/> Receiving zidovudine ≤ 4200mg/week<input type="checkbox"/> <u>Myelodysplasia Syndrome (MDS)</u><ul style="list-style-type: none"><input type="checkbox"/> Combination with G-CSF<input type="checkbox"/> Recent erythropoietin level <500mU/ml<input type="checkbox"/> <u>Anemia of prematurity</u><ul style="list-style-type: none"><input type="checkbox"/> Combination with iron supplementation<input type="checkbox"/> Birth weight of <1500grms<p style="text-align: center;">OR</p><input type="checkbox"/> Gestational age <33 weeks <input type="checkbox"/> <u>Surgery undergoing elective therapy:</u><ul style="list-style-type: none"><input type="checkbox"/> Noncardiac Surgery<p style="text-align: center;">OR</p><input type="checkbox"/> Nonvascular Surgery<input type="checkbox"/> Hgb >10 to ≤13 g/dL | <ul style="list-style-type: none"><input type="checkbox"/> <u>Anemia in Cancer patient</u><ul style="list-style-type: none"><input type="checkbox"/> Non-myeloid Malignancies (i.e. Solid tumors, Multiple Myeloma, Lymphoma, Lymphocytic Leukemia)<input type="checkbox"/> Other Malignancies _____<input type="checkbox"/> Name/Date of Chemotherapy _____<input type="checkbox"/> H/H initial _____<input type="checkbox"/> H/H after 8 weeks _____<input type="checkbox"/> <i>Patient has tried and failed:</i><ul style="list-style-type: none"><input type="checkbox"/> Procrit®<input type="checkbox"/> <u>Hepatitis C treated with ribavirin and Interferon</u><ul style="list-style-type: none"><input type="checkbox"/> Hg ≤10g/dL<input type="checkbox"/> Unresponsive to 200mg/day reduction of ribavirin<p style="text-align: center;">OR</p><input type="checkbox"/> Symptomatic: anemia, cirrhosis, liver transplant, or HIV coinfection <input type="checkbox"/> <u>Sickle Cell Anemia</u> |
|---|--|

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****NOTE: ESAs Not a Covered Benefit:** for cancer treatment due to vitamin deficiencies, hemolysis, bleeding or bone marrow fibrosis not related to chemotherapy, anemia associated with chemotherapy for CML or AML, anemia associated with radiotherapy without concomitant chemotherapy, prophylactic use to prevent chemotherapy induced anemia or tumor hypoxia, patients with EPO-type resistance, patients with treatments including angiogenic drugs and anemia of chronic disease **

Iron studies: shows member has adequate iron stores to support erythropoiesis (*Submit lab test results for review*)

- Patient's serum ferritin is at least 100ng/mL. Ferritin _____
- Patient's most recent transferrin saturation is at least 20% _____
- Drug and dosage regimen prescribed: _____
- Anticipated length of therapy: _____
- IRON THERAPY PRESENT:** _____

Medication being provided by (check applicable box(es) below):

- Physician's office**
- OR**
- 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: 7/20/2012**
REVISED/UPDATED: 7/12/12; 3/12/2014; 8/8/2014; 10/30/2014; 5/21/2015; 12/24/2015; 8/13/2016; 9/20/2016; 11/16/2016; 12/12/2016; 9/12/2017; 10/6/2017.