



15 Earhart Drive, Suite 110, Amherst, NY 14221

ERYTHROPOIETIN AUTHORIZATION AND REAUTHORIZATION REQUEST

TEL: (716) 929-1000 | 1-800-809-4763 FAX: (716) 532-7360

Member Name:			Today's Date:			Date Needed:				
Date of birth:		Sex:	Weight:		Prescriber:			Hospital/Clinic:		
Home Phone Number: () ()				Phone Number: () ()			Fax Number: () ()			
Home Address:			City:	State:	Zip:	Address:		City:	State:	Zip:
Payor: <input type="checkbox"/> Independent Health <input type="checkbox"/> Anne Arundel Health System <input type="checkbox"/> Pharmacy Benefit Dimensions		<input type="checkbox"/> Commercial <input type="checkbox"/> Medicare <input type="checkbox"/> Medicaid <input type="checkbox"/> Self-funded		Prescriber specialty:						
Insurance ID:			Group Number:			Allergies:				
Will patient be self-administering medication?				<input type="checkbox"/> Yes <input type="checkbox"/> No						

STATEMENT OF MEDICAL NECESSITY

Primary Diagnosis: _____

ICD10 Code _____

****Please provide clinical office notes to support this request****

IS EPOETIN ALPHA BEING ADMINISTERED FOR THE TREATMENT OF:

1. Symptomatic anemia associated with chronic renal failure including patients on dialysis (End Stage Renal Disease) and not on dialysis? Yes No

OR:

2. Chronic Kidney disease defined as GFR between 30-75mL/min/1.73m²? Yes No

Provide GFR: _____

- Non-Dialysis patient's hemoglobin is less than 10g/dL? Yes No
- Rate of hemoglobin decline indicated the likelihood of requiring a red blood cell transfusion? Yes No
- Reducing the risk of alloimmunization and/or other RBC transfusion-related risks is a goal? Yes No

OR:

• Dialysis patient's hemoglobin is less than 10g/dL? Yes No

OR:

3. Symptomatic anemia associated with zidovudine treated HIV infected patients when endogenous serum erythropoietin level is ≤ 500 mUnits/ml and patient is receiving a dose of zidovudine ≤ 4200mg/wk? Yes No

• Patient's hemoglobin is less than 11g/dL? Yes No

OR:

4. Symptomatic anemia in a patient with a solid tumor, multiple myeloma, lymphoma or lymphocytic leukemia where anemia is due to the effect of concomitantly administered myelosuppressive chemotherapy? Yes No

- Anticipated duration of myelosuppressive chemotherapy: _____
- Provider must enroll in and comply with ESA APRISE Oncology Program.
- Patient's hgb is less than 10g/dL (or HCT less than 30%)? Yes No

DRUG SELECTION

EPOGEN

PROCIT

RETACRIT

New Authorization

Re-authorization*

Dose: _____

Frequency: _____

Expected duration of therapy: _____

OR:

5. Is patient anemic and scheduled to undergo elective, noncardiac, nonvascular surgery to reduce the need for allogenic blood transfusion? Yes No

6. If not being requested for any of the above indications, please provide the diagnosis for use and clinical rationale to support this request.

FOR EPOGEN AND PROCIT ONLY:

Has Patient tried and failed or have a contraindication to Retacrit?

Yes No

HCT: _____ Date: _____

Hgb: _____ Date: _____

Ferritin Level: _____ Date: _____

Transferrin Saturation: _____ Date: _____

Most recent Blood Pressure reading: _____ Date: _____

FOR RE-AUTHORIZATION:

Please submit lab work or office notes providing the following information

HCT: _____ Date: _____

Hgb: _____ Date: _____

Ferritin Level: _____ Date: _____

Transferrin Saturation: _____ Date: _____

Most recent Blood Pressure reading: _____ Date: _____