Betibeglogene Autotemcel

Goal(s):

• Approve Betibeglogene Autotemcel (ZYNTEGLO) for conditions supported by evidence of benefit

Length of Authorization:

• Once in a lifetime dose.

Requires PA:

• Betibeglogene Autotemcel (billed as pharmacy or physician administered claim)

Covered Alternatives:

- Current PMPDP preferred drug list per OAR 410-121-0030 at www.orpdl.org
- Searchable site for Oregon FFS Drug Class listed at www.orpdl.org/drugs/

Approval Criteria			
1. What diagnosis is being treated?	Record ICD10 code.		
2. Is this an FDA approved indication?	Yes: Go to #3	No: Pass to RPh. Deny; medical appropriateness	
3. Is there documentation that the patient has never received another gene therapy for any diagnosis?	Yes: Go to #4	No: Pass to RPh. Deny; medical appropriateness	
Does patient have confirmed Beta- thalassemia?	Yes: Go to #5	No: Pass to RPh. Deny; medical appropriateness	
5. Is the genotype documented?	Yes: Go to #6 Genotype	No: Pass to RPh. Deny; medical appropriateness	
 6. Is the patient transfusion dependent, defined as requiring in each of the past 2 years: 100 mL/kg/year or more of packed red blood cells (any patient age) OR 8 transfusions or more of packed red blood cells per year (patients 12 years and older) 	Yes: Go to #7	No: Pass to RPh. Deny; medical appropriateness	
7. Is the patient 5 years old or older?	Yes: Go to #9	No: Go to #8	

Approval Criteria			
8. Does the patient weigh at least 6 kg?	Yes : Go to #9	No: Pass to RPh. Deny; medical appropriateness	
Does the patient have cirrhosis or advanced liver disease?	Yes: Pass to RPh. Deny; medical appropriateness	No: Go to #10	
10. Is there documentation that the patient does not have active or chronic infections of HIV, hepatitis B, or hepatitis C?	Yes : Go to #11	No: Pass to RPh. Deny; medical appropriateness	
11. Does the prescriber attest that the patient's general health and comorbidities have been assessed and that the patient is expected to safely tolerate myeloablation?	Yes : Go to #12	No: Pass to RPh. Deny; medical appropriateness	
12. Has the patient (and/or guardian, if applicable) been educated on the risk of insertional oncogenesis and need for lifelong monitoring (bloodwork) at least annually?	Yes : Go to #13.	No: Pass to RPh. Deny; medical appropriateness	
13. Is the patient of childbearing potential OR capable of fathering a child?	Yes: Go to #14	No: Approve one lifetime dose.	
14. Is the patient pregnant, actively trying to conceive, or trying to father a child?	Yes: Pass to RPh. Deny; medical appropriateness.	No: Go to #15	
15. Is there documentation that the provider and patient have discussed the teratogenic risks of the drug if the patient were to become pregnant or father a child during treatment and for at least 6 months after administration of the gene therapy?	Yes: Approve for one lifetime dose		

P&T/DUR Review: 10/23 (SF) Implementation: 11/1/23