

Zolgensma® (Onasemnogene Apeparovovec-xioi) Prior Authorization Form

Member Name: _____ **Date of Birth:** _____ **Member ID#:** _____

Drug Information

Physician billing (HCPCS code: _____) Pharmacy billing* (NDC: _____)
*The NDC for this weight-based medication is specific to the dose required. The NDC provided should reflect the member's current weight.
Projected Date of Infusion: _____ **Dose:** _____ **Regimen:** _____

Zolgensma® Billing Provider Information

Provider NPI: _____ **Provider Name:** _____

Provider Phone: _____ **Provider Fax:** _____

Name of outpatient health care facility where Zolgensma® will be delivered to and administered at: _____

Prescriber Information

Prescriber NPI: _____ **Prescriber Name:** _____

Prescriber Phone: _____ **Prescriber Fax:** _____ **Specialty:** _____

Criteria

For Authorization (Only one Zolgensma® infusion will be approved per member per lifetime):

1. If not previously submitted, please provide the **member's recent progress notes discussing respiratory status.**
2. What is the diagnosis for which the medication is being prescribed?
 - Spinal muscular atrophy (SMA)
 - A. Has the diagnosis been confirmed by molecular genetic testing? Yes ___ No ___
 - B. Does member have bi-allelic pathogenic variants in the *survival motor neuron gene 1 (SMN1)*?
Yes ___ No ___
 - Other, please list: _____
3. Will member have reached full-term gestational age prior to the "Projected Date of Infusion" provided in the Drug Information section of this form? Yes ___ No ___
4. Is member currently dependent on permanent invasive ventilation? Yes ___ No ___
If member requires ventilator support, please provide a recent nursing note stating hours on the ventilator per day.
 - A. If member is currently dependent on permanent ventilation, please specify number of hours per day member requires ventilator support: _____
 - B. If member is currently dependent on permanent ventilation, how many continuous days has member required ventilator support: _____
 - C. Has the member required ventilator support in the absence of an acute, reversible illness or a perioperative state? Yes ___ No ___
5. Is Zolgensma® being prescribed by a neurologist, specialist with expertise in treatment of SMA, or an advanced care practitioner with a supervising physician who is a neurologist or specialist with expertise in treatment of SMA? Yes ___ No ___
6. Please provide member's baseline anti-AAV9 antibody titers: _____
7. Does prescriber agree to monitor liver function tests, platelet counts, and troponin-I at baseline and as directed by the Zolgensma® prescribing information? Yes ___ No ___
8. Does prescriber agree to administer systemic corticosteroids starting 1 day prior to the Zolgensma® infusion and continue as recommended in the prescribing information based on member's liver function? Yes ___ No ___
9. Will the facility where Zolgensma® will be delivered to and administered at, and pharmacy if applicable, adhere to the storage and handling requirements in the Zolgensma® prescribing information? Yes ___ No ___
10. Is member currently receiving treatment with Spinraza® (nusinersen)? Yes ___ No ___
11. Is member currently receiving treatment with Evrysdi™ (risdiplam)? Yes ___ No ___
12. Will Spinraza® or Evrysdi™ treatment be used concomitantly with Zolgensma®? Yes ___ No ___
13. Please provide member's current weight: _____ Date taken: _____

Prescriber Signature: _____ **Date:** _____

(By signature, the physician confirms the criteria information above is accurate and verifiable in patient records.) **Specific information/documentation will be requested if necessary. Failure to complete this form in full will result in processing delays.**

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy
Pharmacy Management Consultants
Product Based Prior Authorization Unit

Fax: 1-800-224-4014
Phone: 1-800-522-0114 Option 4

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