



# State of Oklahoma SoonerCare

## Synagis® (Palivizumab) Continuation Form

**Please Note: This form is for continuation of Synagis® therapy only.**

This form should only be submitted following initial approval of Synagis® during the current RSV season in Oklahoma. Each approval will be for a duration of 1 month. Subsequent approval consideration will also require use of this form and will only be granted monthly during the current RSV season in Oklahoma.

For initial Synagis® approval consideration, please submit the Synagis® (Palivizumab) Initiation Prior Authorization Form (PHARM-7A) which is available on the OHCA website at: <https://oklahoma.gov/ohca/providers/forms/rxfoms.html>.

**Member Name:** \_\_\_\_\_ **Date of Birth:** \_\_\_\_\_ **Member ID#:** \_\_\_\_\_

### Drug Information

FDA approved dosing: 15mg/kg intramuscularly. Only those doses that require greater than a vial's dose +10% may use the next vial size or an additional vial (e.g. 1-55mg = 50mg vial, 56-110mg = 100mg vial). Weight must be taken within the last 3 weeks. Each dose is to be given every 30 days.

Physician billing     CPT code 90378 (50mg/unit)  
Pharmacy billing     50mg/0.5ml NDC: \_\_\_\_\_     100mg/ml NDC: \_\_\_\_\_

### Billing Provider Information

Provider: \_\_\_\_\_ Provider NPI: \_\_\_\_\_  
Provider Phone: \_\_\_\_\_ Provider Fax: \_\_\_\_\_

### Prescriber Information

Prescriber: \_\_\_\_\_ Prescriber NPI: \_\_\_\_\_  
Prescriber Phone: \_\_\_\_\_ Prescriber Fax: \_\_\_\_\_

### Synagis® Continuation Information

**For continued authorization of Synagis®, please provide all of the following:**

1. Previous Dose Information:
  - a. Date last dose of Synagis® was received: \_\_\_\_\_  
(each dose is to be given every 30 days)
  - b. Dose of last Synagis® injection: \_\_\_\_\_ (mg)
2. Current Weight Information:
  - a. Member's current weight: \_\_\_\_\_ (kg)
  - b. Date member's weight was recorded: \_\_\_\_\_  
(weight must be taken within the last 3 weeks)
3. Has the member received Beyfortus™ (nirsevimab-alip)? Yes \_\_\_ No \_\_\_
4. Please provide a patient-specific, clinically significant reason why the member cannot receive Beyfortus™ (nirsevimab-alip): \_\_\_\_\_

<p><b>PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:</b></p> <p>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</p>	<p><b>CONFIDENTIALITY NOTICE</b></p> <p><i>This document, including any attachments, contains information which is confidential or privileged. If you are not the intended recipient, be aware that any disclosure, copying, distribution, or use of the contents of this information is prohibited. If you have received this document in error, please notify the sender immediately by telephone to arrange for the return of the transmitted documents or to verify their destruction.</i></p>
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