

Request for Prior Authorization — Antiviral Monoclonal Antibodies

Anthem Blue Cross and Blue Shield | Serving Hoosier Healthwise, Healthy Indiana Plan, Hoosier Care Connect, and Indiana PathWays for Aging

CONTAINS CONFIDENTIAL PATIENT INFORMATION

Complete form in its entirety and fax to Prior Authorization of Benefits Center at 844-864-7860 (retail) or 888-209-7838 (medical injectable)
Provider Help Desk: 866-408-6132 (Hoosier Healthwise)
844-533-1995 (Healthy Indiana Plan)
844-284-1798 (Hoosier Care Connect)
833-569-4739 (Indiana PathWays for Aging)

Today's date:

Month	Day	Year	

Note: This form must be completed by the prescribing provider.

All sections must be completed or the request will be returned.

Patient's Medicaid #:	Date of birth:		
Patient name:	Prescriber's name:		
Prescriber's IN license #:	Specialty:		
Prescriber's NPI #:	Prescriber's signature: **required below within attestation section**		
Return fax #:	Return phone #:		
Mark box if requesting retroactive PA: □	Date(s) of service requested for retroactive eligibility (if applicable):		

Note: Submit PA requests for retroactive claims (dates of service prior to eligibility determination, but within established eligibility timelines) with dates of service prior to 30 calendar days of submission separately from current PA requests (dates of service 30 calendar days or less and going forward).

Requested medication	Strength	Dosage regimen

Providers who are contracted with Anthem Blue Cross and Blue Shield to serve Hoosier Healthwise, Healthy Indiana Plan, Hoosier Care Connect, and Indiana PathWays for Aging through an accountable care organization (ACO), participating medical group (PMG) or Independent Physician Association (IPA) are to follow guidelines and practices of the group. This includes but is not limited to authorization, covered benefits and services, and claims submittal. If you have questions, please contact your group administrator or your Anthem network representative.

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PΑ	requirements for Synagis (palivizumab):				
1.	Patient information:				
	Actual gestational age:weeksdays				
	Current age (Must be < 24 months):months				
	Current weight: □ kg □ lb				
2.	Prescription information: ☐ Inject 15mg/kg IM once per month through March 31st				
	□ Other:				
3.	alivizumab Prior Authorization Criteria Guidelines for a maximum of five doses (approval will branted under any of the following circumstances)^:				
	If member is less than 12 months of age, select one of the following that is applicable:				
	Member was born before 29 weeks, 0 days' gestation				
	Member was born before 32 weeks, 0 days' gestation and has CLD necessitating more than 21% oxygen for at least the first 28 days of life				
	Please provide dates of oxygen supplementation/medication intervention:				
	Member has hemodynamically significant heart disease (for example, acyanotic heart disease receiving medication to control CHF and will require cardiac surgical procedures, or those with moderate to severe pulmonary hypertension)				
	Please provide relevant diagnoses/medical intervention:				
	Member has congenital airway abnormality or neuromuscular disease that impairs the ability to clear secretions				
	Please provide relevant diagnoses/medication intervention:				
	Member has cystic fibrosis with clinical evidence of CLD and/or nutritional compromise				
	If member is less than 24 months of age, select one of the following that is applicable:				
	Member is or will be considered to be profoundly immunocompromised (must provide chart documentation and explicitly state how member is or will be considered to be profoundly immunocompromised during the RSV season), including members undergoing cardiac transplantation during current RSV season.				

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Please explain:
☐ Member was born before 32 weeks, 0 days' gestation and required at least 28 days of supplemental oxygen after birth and who continued to require supplemental oxygen, chronic systemic corticosteroid therapy, diuretic, or bronchodilator therapy within 6 months of the start of the second RSV season
Please provide dates of oxygen supplementation/medication intervention:
 Member has cystic fibrosis with manifestations of severe lung disease (previous hospitalization for pulmonary exacerbation in the first year of life or abnormalities on chest imaging that persist when stable) or weight-for-length <10th percentile
Please provide relevant diagnoses/medical intervention:
Prescriber has submitted valid medical justification for the use of Synagis (palivizumab) over Beyfortus (nirsevimab)** ☐ YES ☐ NO
Medical justification:
Prescriber attests member has not received Beyfortus (nirsevimab) within the same RSV season Prescriber signature:

Note: Prophylaxis will be given only until the infant or child reaches a maximum of five doses or the end of the RSV season, whichever comes first.

The Respiratory Syncytial Virus (RSV) season is defined as November 1st through March 31st. The Office of Medicaid Policy & Planning may extend the season based on statewide virology data. Requests for additional doses beyond the initial five approved doses will require separate prior authorization.

CONFIDENTIAL INFORMATION

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