## MHS PHARMACY BENEFIT ANTIVIRAL MONOCLONAL ANTIBODIES PRIOR AUTHORIZATION REQUEST FORM



## MHS 429 N Pennsylvania St Indianapolis, IN, 46204-1208 Phone: (877) 647-4848 Fax: (866) 399-0929



Today's Date				
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's Name	Prescriber's Name			
Prescriber's IN License #	Specialty			
Prescriber's NPI#	Prescriber's Signature			
Return Fax #	Return Phone #			
Check box if requesting retro-active PA	Date(s) of service requested for retro-active 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).				
PA Requirements for Synagis (palivizumab):  1. Patient Information:				
	daya			
Actual Gestational Age:weeks				
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. Palivizumab Prior Authorization Criteria Guidelines for a maximum of 5 doses (approval will be granted under any of the following circumstances) <sup>^</sup> :				
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:				

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		Member has hemodynamically significant heart disease (e.g., 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	
lf n	nembe	r is less than 24 months of age, select one of the following that is applicable:	
	and	nber is or will be considered to be profoundly immunocompromised (must provide chart documentation explicitly state how member is or will be considered to be profoundly immunocompromised during the 'season', including members undergoing cardiac transplantation during current RSV season	1
	Plea	se explain:	
	after diure	nber was born before 32 weeks, 0 days' gestation and required at least 28 days of supplemental oxygen birth and who continued to require supplemental oxygen, chronic systemic corticosteroid therapy, etic, or bronchodilator therapy within 6 months of the start of the second RSV season are provide dates of oxygen supplementation/medication intervention:	en
	puln	nber has cystic fibrosis with manifestations of severe lung disease (previous hospitalization for nonary exacerbation in the first year of life or abnormalities on chest imaging that persist when stable) with-for-length <10 <sup>th</sup> percentile	or
	Plea	se provide relevant diagnoses/medical intervention:	
4.		riber has submitted valid medical justification for the use of Synagis (palivizumab) over Beyfortus vimab)   Vimab)   Vimab	
	Medic	al justification:	
5.	Presci	riber attests member has NOT received Beyfortus (nirsevimab) within the same RSV season	
	Presci	riber signature:	

**Note:** Prophylaxis will be given only until the infant or child reaches a maximum of 5 doses or the end of the RSV season, whichever comes first

^The Respiratory Syncytial Virus (RSV) season is defined as November 1<sup>st</sup> through March 31<sup>st</sup>. The Office of Medicaid Policy & Planning may extend the season based on statewide virology data. Requests for additional doses beyond the initial 5 approved doses will require separate prior authorization.

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