2021-2022 Synagis® Season – Prior Authorization Form



Phone: 1-800-218-7453 ext. 22080 | Fax: 1-866-683-5631

Section I — Dispensing Pharmacy Information

Name of Pharmacy	National Provider Ide	ntifier (NPI)	Area Code and Telephone	No. Area Code a	Area Code and Fax No.		
Section II — Patient Demographics				<u>'</u>			
Name of Patient	Medicaid ID		Date of Birth (MMDDYY)	Gestational Age			
				weeks and	/ 7th day		
Address of Patient (Street, City, State, ZIP Code	e) Patient Phor	ne Number		County of Reside	ence		
Has patient received a Synagis prophylactic inje	ection during hospitaliz	zation since t	ne start current of the RSV se Date(s):	eason?			
Has the patient been hospitalization due to RS\ No Yes If yes, date of diagnosis:	at any time since the	e start of the o	current RSV season?				
Section III — Patient Diagnosis at the (Diagnosis/conditions must be clearly document							
Patients who are younger than 24 month s chronological age can qualify, for up to five monthly doses of Synagis, based on diagnosis listed to the right.	hematopoietic stem cell transplant, chemotherapy or other condition that leaves the infant profoundly immunocompromised):						
Patients who are between 12 - 24 months chronological age at the start of the RSV season can qualify, for up to five monthly doses of Synagis, based on the diagnosis or conditions listed to the right. Please refer to page 2 for definition.	of the following therapies within the 6 months prior to the current RSV season (che that apply):						
	24-3: Diagnosis of cystic fibrosis with severe lung disease*, or cystic fibrosis with weight or length less than the 10th percentile:						
	ICD-10-CM co	de:					
Patients who are younger than 12 months chronological age at the start of the RSV season can qualify, for up to five monthly doses of Synagis, based on criteria listed to the right.	☐ 12-1: ≤ 28 6/7 weeks gestational age at birth:						
	ICD-10-CM code:						
	12-2: Chronic lung disease (CLD) of prematurity#:						
	ICD-10-CM code:						
	12-3: Severe congenital abnormality of airway OR severe neuromuscular disease that impairs the ability to clear secretions from the upper airway because of ineffective cough:						
	ICD-10-CM co	de:					

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	12-4: Active diagnosis of he	emodynamically significant c	ongenital heart disease (CHD):	
	ICD-10-CM code:			
AN	any of the below			
	Moderate to severe pulmon	nary hypertension.		
	Acyanotic heart disease, or will require cardiac surgery	n medication to control cong y.	jestive heart failure, and	
	Cyanotic heart disease.			
l',	TE: This excludes infants with h r list.)	nemodynamically insignificant he	eart disease - refer to pages 2 and	
	12-5: Diagnosis of cystic fib compromise.	prosis with clinical evidence	of CLD and/or nutritional	
	ICD-10-CM code:			
2 (; N/				
Section IV — Synagis Prescription detail (to be con			n to the specialty pharmacy.	
Rx: Synagis (palivizumab) Injection Quantity:	Dose ((mg): 		
Sig: Inject 15mg/kg one time per month Current V	(kg) 0	1 1 2		
Syringes 1ml 25G 5/8* Syringes 3ml 20G 1	Epinephrine 1:1000 a	amp. Sig: Injected 0.01 mg/k	g as directed.	
Prescriber Name	License	No.	NPI	
Address of Prescriber (Street, City, State and ZIP Co	le) Ar	rea Code and Telephone No	. Area Code and Fax No.	
Physician Signature			Date	
Fax the completed prior a	ithorization form to Sup	erior HealthPlan at 1-86	6-683-5631	
Category	Subcategories			
# Chronic Lung Disease (CLD) of Prematurity	• Infants born < 32 we for at least 28 days	32 weeks, 0 days' gestational age who require >21% oxygen days after birth.		
Hemodynamically significant heart disease	Congestive heart fa	ilure (CHF) requiring med	lication	

Category	Subcategories
# Chronic Lung Disease (CLD) of Prematurity	• Infants born < 32 weeks, 0 days' gestational age who require >21% oxygen for at least 28 days after birth.
Hemodynamically significant heart disease	Congestive heart failure (CHF) requiring medication
	Moderate to severe pulmonary hypertension
	Unrepaired cyanotic congenital heart disease
*Severe lung disease	Previous hospitalization for pulmonary exacerbation in the first year of life or abnormalities on chest radiography or chest computed tomography that persist when stable
The following groups of infants are NOT AT INCREAS	ED risk of RSV and generally should not receive immunoprophylaxis:
1.Hemodynamically insignificant heart disease	Secundum atrial septal defect
	Small ventriculoseptal defect
	Pulmonic stenosis
	Uncomplicated aortic stenosis
	Mild coarctation of the aorta
	Patent ductus arteriosus
Congenital heart disease adequately corrected by failure.	surgery which does not continue to require medication for congestive heart

3. Mild cardiomyopathy that does not require medical therapy for the condition.

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Category Subcategories

4. Children in the second year of life on the basis of a history of prematurity alone.

Note: Tobacco smoke exposure is <u>not</u> an indication for Synagis administration. Tobacco dependent parents should be offered tobacco dependence treatment or referral for tobacco dependence treatment. 877-YES-QUIT (877-937-7848, YesQuit.org) is the Quitline operated in Texas.

Additional Information

- Texas Medicaid has adopted the updated guidance published in 2014 by the American Academy of Pediatrics.
- Infants born at 29 weeks, 0 days' gestation or later are no longer universally recommended to receive prophylaxis with Synagis.
- Infants born at 29 weeks, 0 days' gestation or later, on the basis of chronic lung disease, congenital heart disease, or another condition, may qualify to receive prophylaxis.
- Synagis is not recommended in the second year of life on the basis of prematurity alone.
- Monthly prophylaxis should be discontinued in any child who experiences a breakthrough RSV hospitalization.

References

- "Updated Guidance for Palivizumab Prophylaxis Among Infants and Young Children at Increased Risk of Hospitalization for Respiratory Syncytial Virus Infection." *Pediatrics* 134.2 (2014): 415-420. Web. 11 Aug. 2015.
- Synagis® (palivizumab) [prescribing information]. Gaithersburg, MD: Medimmune, LLC. 2014.
- Epinephrine 1:1000 (1mg/ml) [prescribing information]. Lake Forest, IL: Hospira. 2008.