

Division: Pharmacy Services	Subject: Prior Authorization Criteria
Effective Date:	September 9, 2014
Revision Date	October 30, 2014, November 19,2014, June 30, 2015

SYNAGIS® (Palivizumab)^{i,ii,iii}

LENGTH OF AUTHORIZATION:

- Authorize for a maximum of 5 doses during RSV reason (maximum of 5 monthly doses of 15 mg/kg IM) for all recipients **EXCEPT**:
- Authorize a maximum of 3 doses or up to 90 days of age (whichever occurs first) for infants born between 29 weeks 1 day and 34 weeks, 6 days gestational age AND who are currently less than 3 months of age at the start of RSV season AND who have at least one of the following two risk factors:
 - Recipient has a sibling or other child under age 5 living permanently in their home
 - o Recipient attends child care/day care where multiple children are present
- In infants and children < 24 months, already on prophylaxis and eligible, one post-op dose can be approved after cardiac bypass or after extracorporeal membrane oxygenation (ECMO).

CLINICAL NOTES:

Palivizumab (Synagis) is a respiratory syncytial virus (RSV) F protein inhibitor monoclonal antibody indicated for the prevention of serious lower respiratory tract disease caused by RSV in children at high risk of RSV disease. The American Academy of Pediatrics (AAP) has issued an updated guidance for palivizumab prophylaxis among infants and young children at increased risk of hospitalization for RSV.

- There is variability in the onset and offset of RSV season. Generally it runs from November to April within the continental U.S.
- Florida- Data from the Florida Department of Health can be used to determine the onset and offset of RSV season in different regions of Florida.
- Native American Indian infants- There is limited information about the burden of RSV infection among American Indian populations. Prophylaxis can be considered for Navajo and White Mountain Apache infants in the first year of life.



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APPROVAL CRITERIA

Palivizumab will be approved in the following scenarios:

Infant/Child Age at Start of RSV Season	Criteria
< 12 months (first year of life)	 GA< 29 wks, 0 d (otherwise healthy) CLD of prematurity (GA<32 wks, 0 d and with supplemental O₂ for at least the first 28 d after birth) Anatomic pulmonary abnormalities or neuromuscular disorder, or congenital anomaly that impairs the ability to clear secretions Profoundly immunocompromised with conditions such as SCID, immunocompromised infant with stem cell transplant, severe acquired immunodeficiency syndrome (AIDS) CF with CLD and/or nutritional compromise GA 29 wks 1 d-34 wks, 6 d who are less than 3 months of age at the start of RSV season and have at least one risk factor (siblings or other children < 5 y living permanently in the home OR recipient attends day care/ child care with multiple children)-Maximum of 3 doses
\leq 12 months (first year of life)	 CHD (hemodynamically <i>significant</i>) with acyanotic* heart disease on medications to control CHF and will require cardiac surgery or infants with moderate to severe PH. For <i>cyanotic</i>* heart defects, a pediatric cardiologist should be consulted.
> 12 months to 23 months	 CLD of prematurity (GA < 32 wks, 0 d and supplemental O₂ for at least the first 28 d after birth) and medical support (chronic systemic steroids, diuretic therapy, or supplemental O₂) within 6 months before start of 2nd RSV season CF with severe lung disease** or weight for length < 10th percentile
< 24 months (2 nd year of life)	 Cardiac transplant during RSV season Already on prophylaxis and eligible; give post-op dose after cardiac bypass or after ECMO Profoundly immunocompromised with conditions such as SCID, immunocompromised infant with stem cell transplant, severe acquired immunodeficiency syndrome (AIDS)

GA=gestational age; wks=weeks; d=day; CLD=chronic lung disease; SCID= severe combined immune deficiency; CHD=congenital heart disease; O2=oxygen; HD=heart disease; CHF=congestive heart failure; PH=pulmonary hypertension; CF=cystic fibrosis; ECMO=extracorporeal membrane oxygenation

* Examples of acyanotic heart defects include ventricular septal defects, atrial septal defects, pulmonary valve stenosis and aortic valve stenosis. For *cyanotic* heart defects, a pediatric cardiologist should be consulted

** Examples of severe lung disease: previous hospitalization for pulmonary exacerbation in the 1st year of life, abnormalities on chest radiography [chest X-ray], or chest computed tomography [chest CT] that persist when stable



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DENIAL CRITERIA

Palivizumab will NOT be approved in the following scenarios:

Infant/Child Age at Start of RSV Season	Deny
> 12 months (2 nd year of life)	 Based on prematurity alone
	 CLD without medical support (chronic systemic
	steroids, diuretic therapy or supplemental O_2)
	CHD
	• Otherwise healthy children in 2 nd year of life
Any age	 Breakthrough RSV hospitalization ***
	 Hemodynamically insignificant CHD****
	 CHD lesions corrected by surgery (unless on CHF
	meds)
	 CHD and mild cardiomyopathy not on medical
	therapy
	• CHD in 2 nd year of life
No specific age defined	 Asthma prevention
	 Reduce wheezing episodes
	 Down Syndrome
	 CF (otherwise healthy)
	 Healthcare-associated RSV disease****

***If any infant or child is receiving palivizumab prophylaxis and experiences a breakthrough RSV hospitalization, discontinue palivizumab, because the likelihood of a second RSV hospitalization in the same season is extremely low.

****Examples of hemodynamically *insignificant* CHD: secundum atrial septal defect, small ventricular septal defect, pulmonic stenosis, uncomplicated aortic stenosis, mild coarctation of the aorta, patent ductus arteriosus.

***** No rigorous data exist to support palivizumab use in controlling outbreaks of health care-associated disease; palivizumab use is not recommended for this purpose.

REFERENCES

10.1542/peds.2014-1666. Available at: http://pediatrics.aappublications.org/content/early/2014/07/23/peds.2014-1666 Accessed August 27, 2014

ⁱ American Academy of Pediatrics. Position Statement. Updated guidance for palivizumab prophylaxis among Infants and young children at increased risk of hospitalization for respiratory syncytial virus infection. Pediatrics 2014; 134;415. DOI: 10.1542/peds.2014-1665. Available at:

http://pediatrics.aappublications.org/content/134/2/415.full.pdf+html?sid=c5cf7568-4302-4ccd-9c71-ea785e33e241 Accessed August 27, 2014

ⁱⁱ American Academy of Pediatrics. Technical Report. Updated guidance for palivizumab prophylaxis among Infants and young children at increased risk of hospitalization for respiratory syncytial virus infection. DOI:

ⁱⁱⁱ Synagis [package insert]. Gaithersburg, MD; MedImmune; March 2014.