

**Louisiana Fee-For-Service Medicaid**  
**Palivizumab (Synagis®) for the 2018-2019 Respiratory Syncytial Virus (RSV) Season**

Palivizumab is indicated for the prevention of serious lower respiratory tract infection caused by respiratory syncytial virus (RSV) in selected infants and young children at high risk of RSV disease. Monthly prophylaxis should be discontinued in any infant receiving monthly palivizumab prophylaxis who experiences a breakthrough RSV hospitalization.

**Clinical Pre-Authorization Criteria**

All prescriptions for palivizumab for recipients in Fee-For-Service Medicaid require clinical pre-authorization.

**Prescribing providers**, not the pharmacy, manufacturer or any other third party entity, must complete the *Palivizumab Clinical Pre-Authorization Form* and fax it **directly** to LA Medicaid Rx PA Operations at the University of Louisiana at Monroe College of Pharmacy at **866-797-2329**. Any requests submitted early will not be processed prior to the start of RSV season. Prescribing providers will be notified by fax or mail of the outcomes of clinical pre-authorization requests.

**Clinical pre-authorization will be considered for approval when requests meet the following criteria:**

- Palivizumab clinical pre-authorization requests will be considered in accordance with an RSV season of November 1, 2018 through March 31, 2019; **AND**
- Recipient must meet gestational age **AND** chronological age requirements for the ICD-10-CM diagnosis code(s) and/or other qualifying risk factor(s) submitted with the request. Supporting documentation (i.e. progress notes, hospital discharge notes, pediatric cardiologist consult notes, chart notes, pharmacy profiles, etc.) is required and must be submitted with each request. Requests for palivizumab will be considered for approval when **ONE** of the following ‘high-risk’ criteria are met:
  - 1. Infant born prematurely without chronic lung disease (CLD) OR without hemodynamically significant cyanotic or acyanotic heart disease or without other listed ‘high-risk’ factors:**
    - The infant is younger than 12 months of age on November 1, 2018, **AND** was born before 29 weeks, 0 days’ ( $\leq 28$  weeks, 6 days’) gestation.
  - 2. Infant with chronic lung disease (CLD) (one of the criteria sets below must be met):**
    - **SET 1:** Infant diagnosed with CLD who is 12 months of age or younger, whose first birthday is on or after November 1, 2018, **AND** the infant was born at  $< 32$  weeks, 0 days’ gestation **AND** the infant required  $> 21\%$  oxygen for at least 28 days after birth; **OR**
    - **SET 2:** Infant diagnosed with CLD who is 24 months of age or younger, whose second birthday is on or after November 1, 2018, infant’s second dosing season, **AND** the infant was born at  $< 32$  weeks, 0 days’ gestation **AND** the infant required  $> 21\%$  oxygen for at least 28 days after birth **AND** the infant has required medical therapy (i.e., chronic systemic corticosteroid therapy, diuretic therapy, or supplemental oxygen) during the six (6) months before November 1, 2018, the start of the infant’s second (RSV) season.
  - 3. Infant with congenital heart disease (CHD):**
    - The infant’s first birthday is on or after November 1, 2018; **AND**
    - The infant meets one of the following hemodynamically significant conditions:

- The infant has cyanotic heart defect(s) and decision for use of palivizumab was made with pediatric cardiologist consultation; **OR**
- The infant has acyanotic heart disease **AND** is receiving medication to control congestive heart failure **AND** will require a cardiac surgical procedure; **OR**
- The infant has moderate to severe pulmonary hypertension; **OR**
- The infant has lesions that have been adequately corrected by surgery but continues to require medication for congestive heart failure.

**4. Infant with cardiac transplant**

- The infant is younger than 2 years of age on November 1, 2018; **AND**
- The infant has undergone or will undergo cardiac transplantation from November 1, 2018 through March 31, 2019.

**5. Infant with a congenital anatomic pulmonary abnormality or neuromuscular disease:**

- The infant’s first birthday is on or after November 1, 2018; **AND**
- The infant’s congenital anatomic pulmonary abnormality or neuromuscular disease impairs the ability to clear secretions from the upper airways because of ineffective cough.

**6. Immunocompromised infant:**

- The infant’s second birthday is after November 1, 2018; **AND**
- The child is/will be profoundly immunocompromised (for example, receiving chemotherapy or immunosuppressive therapy) from November 1, 2018 through March 31, 2019.

Medical Reconsideration

Medical Reconsideration of a denied clinical pre-authorization decision may be requested by the prescribing provider. Reconsideration requires completion of the Palivizumab Request for Reconsideration form available at [www.lamedicaid.com](http://www.lamedicaid.com). The form must be completed in full and signed by the prescribing provider. Signature stamps and proxy signatures are not acceptable and will be returned to the requesting provider. The completed form must be faxed from the prescribing provider to the LA Medicaid Rx PA Operations at the University of Louisiana at Monroe College of Pharmacy at 318-812-2940.

**Point-of-Sale (POS) Requirements**

Age Restriction

- Palivizumab claims for recipients who are twenty-four (24) months of age or younger as of November 1, 2018 meet the POS age requirement.

Maximum Number of Doses

- Up to a maximum number of five (5) doses will be reimbursed during the RSV season. Qualifying infants born during the RSV season require fewer doses. For example, infants born in January would receive their last dose in March. A claim submitted for palivizumab outside the maximum number of doses allowed will deny with:

**NCPDP rejection code 88 (DUR Reject Error) mapped to  
EOB code 656 (Exceeds Maximum Duration of Therapy)**

Early Refill

- Palivizumab claims will only process for payment every twenty-eight (28) days.

## **PALIVIZUMAB CRITERIA ICD-10-CM CODE and MEDICATION LIST**

*Note: ANY accepted diagnosis/ICD-10-CM Code listed on the clinical pre-authorization form **MUST** have supporting documentation attached. Supporting documentation is supplemental information submitted to support the patient meeting the criteria and may include copies of progress notes, hospital discharge notes, pediatric cardiologist consult notes, chart notes, pharmacy profiles, etc.*

### **I. Neuromuscular Disorders**

Acceptable ICD-10 codes include:

<b>A80.0-A80.39</b>	Infantile paralysis
<b>G31.9</b>	Cerebral degenerations
<b>G25.3</b>	Myoclonus
<b>G11.1, G11.4</b>	Spinocerebellar disease
<b>G12.0</b>	Werdnig-Hoffman disease (Infantile spinal muscular atrophy)
<b>G12.1, G12.8, G12.9</b>	Spinal muscular atrophy
<b>G12.2*</b>	Motor neuron disease

Exclude (but not limited to) the following (i.e. the following are **NOT** accepted):

<b>G80*</b>	Cerebral palsy
<b>G40.3*</b>	Generalized convulsive epilepsy
<b>G40.4*</b>	Grand mal seizures
<b>G40*</b>	Epilepsy
<b>Q05*</b>	Spina bifida
<b>P90</b>	Newborn seizures
<b>R56*</b>	Infantile seizures

### **II. Congenital Abnormalities of the Airways**

Acceptable ICD-10 codes include:

<b>G47.35</b>	Congenital central alveolar hypoventilation syndrome
<b>Q32.0, Q32.1</b>	Other diseases of the trachea and bronchus, not elsewhere classified (Must specify Tracheomalacia or tracheal stenosis)
<b>Q31.1, Q31.5, Q32.1, Q32.4</b>	Other anomalies of larynx, trachea, and bronchus (Must specify congenital tracheal stenosis, subglottic stenosis, atresia of trachea, laryngomalacia, or absence or agenesis of bronchus, trachea)
<b>Q33.0</b>	Congenital cystic lung
<b>Q33.3, Q33.6</b>	Agenesis, hypoplasia, and dysplasia of the lung
<b>Q33.4</b>	Congenital bronchiectasis
<b>Q38.2</b>	Macroglossia
<b>Q38.5</b>	Uvula anomaly
<b>J98.6</b>	Diaphragmatic paralysis
<b>Q87.3</b>	Beckwith-Wiedemann syndrome

Exclude (but not limited to) the following (i.e. the following are **NOT** accepted):

<b>Q33.9</b>	Anomaly of lung, unspecified
<b>Q33.1, Q33.8</b>	Other anomaly of the lung

**III. Chronic Lung Disease**

Acceptable ICD-10 code:

<b>P27*</b>	Chronic respiratory disease arising in the perinatal period (CLD/BPD/Interstitial pulmonary fibrosis of prematurity/Wilson-Mikity syndrome)
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Exclude (but not limited to) the following (i.e. the following are **NOT** accepted):

<b>J05.0</b>	Croup
<b>J06*</b>	URI
<b>J20*</b>	Bronchitis
<b>J21*</b>	Bronchiolitis
<b>J45*</b>	Asthma
<b>R06.2</b>	Wheezing

**IV. Congenital Heart Diseases (CHD)** Per AAP guidelines, prophylaxis with palivizumab in children with CHD should be made on the degree of cardiovascular compromise. CHD that is deemed hemodynamically insignificant will not meet criteria. Documentation must specifically support CHD being hemodynamically significant (e.g. medications, etc.).

Acceptable ICD-10 codes include:

**A. Acyanotic CHD: Must currently be receiving medication to control CHF (see below)**

<b>Q23.0</b>	Aortic stenosis
<b>I37.0, I37.1, I37.2, Q22.1, Q22.2</b>	Pulmonary valve disorders (incompetence, insufficiency, regurgitation, and stenosis)
<b>I42*, I43</b>	Cardiomyopathy (must be moderate to severe)
<b>Q21.0</b>	Ventricular septal defect
<b>Q21.1</b>	Atrial septal defect
<b>Q21.2</b>	Atrioventricular canal (endocardial cushion defect)
<b>Q22.3</b>	Anomalies of pulmonary valve congenital
<b>Q22.1</b>	Pulmonic stenosis
<b>Q23.0</b>	Congenital stenosis of aortic valve (congenital aortic stenosis) [Excludes: congenital subaortic stenosis; supraaortic stenosis]
<b>Q23.3</b>	Congenital mitral insufficiency
<b>Q25.0</b>	Patent ductus arteriosus
<b>Q25.1</b>	Coarctation of the aorta
<b>Q25.2, Q25.3</b>	Atresia and stenosis of aorta (absence, aplasia, hypoplasia, stricture of the aorta) Supra (valvular)-aortic stenosis [Excludes: congenital aortic (valvular) stenosis or stricture; hypoplasia of aorta in hypoplastic left heart syndrome]

**B. Cyanotic CHD: Does not require use of medication/must not have had or completed surgical correction**

<b>Q20.0</b>	Truncus arteriosus
<b>Q20.3</b>	Transposition of the great vessels
<b>Q21.3</b>	Tetralogy of Fallot
<b>Q22.0</b>	Atresia, congenital
<b>Q22.4</b>	Tricuspid atresia and stenosis, congenital
<b>Q22.5</b>	Ebstein's anomaly
<b>Q23.4</b>	Hypoplastic left heart
<b>Q22.6</b>	Hypoplastic right heart
<b>Q25.5</b>	Pulmonary atresia
<b>Q26.2</b>	Total anomalous pulmonary venous return

**C. Pulmonary Hypertension:**

<b>I26.0*</b>	Acute cor pulmonale
<b>I27.0</b>	Primary pulmonary hypertension
<b>I27.2</b>	Other chronic pulmonary heart disease (pulmonary hypertension, secondary)
<b>P29.3</b>	Persistent fetal circulation (persistent pulmonary hypertension/primary pulmonary hypertension of newborn)

\*any number or letter or combination of **UP TO FOUR** numbers and letters of an assigned ICD-10-CM diagnosis code

**ACCEPTABLE MEDICATIONS USED IN CHD**

Digoxin	ACE Inhibitors	Supplemental oxygen
Beta Blockers	Nitroglycerin	Diuretics
Calcium Channel Blockers	Anti-Coagulants	

**Reference**

STAT!Ref - Red Book®: 2018-2021 Report of the Committee on Infectious Diseases. Online.statref.com. <http://online.statref.com/publictitleinfo/titleinfo.aspx?fxid=76> Published 2018.

Fax this completed form to:  
 La Medicaid Rx PA Operations  
 ULM School of Pharmacy  
 1800 Bienville Drive  
 Monroe, LA 71201-3765  
 FAX 866-797-2329

**State of Louisiana**  
**Department of Health**  
 Bureau of Health Services Financing  
**Palivizumab Clinical Pre-Authorization Form**  
**For 2018- 2019 RSV Season**

**Palivizumab Form: Rx PA01P**  
**Revised: 10/31/2018**  
 VOICE PHONE 866-730-4357

Request must be faxed. Please type or print legibly. Incomplete forms will not be approved.

Date of Request \_\_\_\_\_

<b>Prescribing Provider Information</b>	<b>Recipient Information</b>	
Name (Last, First)	Name (Last, First)	
LA Medicaid Prescribing Provider Number / NPI	LA Medicaid CCN or Recipient Number	
Call-Back Phone Number (include area code)	Date of Birth (mm/dd/yy)	Gestational Age (weeks/days)
FAX Number (include area code)	Recipient Current Weight _____ kg as of _____ (mm/dd/yy)	
Drug and Strength Requested	Diagnosis Code(s) (ICD-10-CM) to Justify Palivizumab Use	
Office Contact Name	EPSDT Support Coordinator (Name / Address) (optional)	

**Does the patient have additional insurance coverage (TPL)?** \_\_\_ Yes \_\_\_ No **If Yes, please contact TPL to determine coverage for this drug.**

**Check the applicable age/condition.** For chronic lung disease (CLD) of prematurity/congenital heart disease (CHD), attach supporting documentation (e.g. hospital birth discharge notes, pediatric cardiologist consult notes and/or chart notes) for any submitted qualifying criteria or ICD-10 diagnosis code(s). Please refer to the Palivizumab Criteria ICD-10-CM Diagnosis Code and Medication List.

- Infant's gestational age is less than 29 weeks, 0 days AND infant's chronological age is less than 12 months old as of November 1, 2018.
- Infant is 12 months old or younger (infant's first birthday is on or after November 1, 2018) with CLD of prematurity, defined as an infant with gestational age of less than 32 weeks, 0 days who required supplemental oxygen greater than 21% for at least the first 28 days after birth.
- Infant is 24 months old or younger (infant's second birthday is on or after November 1, 2018) with CLD of prematurity, defined as an infant with gestational age of less than 32 weeks, 0 days who required supplemental oxygen greater than 21% for at least the first 28 days after birth AND infant continued to require medical support (chronic systemic corticosteroid therapy, diuretic therapy, or supplemental oxygen) during the 6-month period before the start of the infant's second respiratory syncytial virus (RSV) season, which is November 1, 2018.
- Infant is 12 months old or younger (infant's first birthday is on or after November 1, 2018) with hemodynamically significant CHD WITH: (check one) (list applicable diagnosis codes \_\_\_\_\_)  
 \_\_\_\_\_ acyanotic heart disease AND is receiving medication to control congestive heart failure (CHF) such as diuretics, ACE inhibitors, beta-blockers or digoxin AND will require a cardiac surgical procedure.  
 \_\_\_\_\_ moderate to severe pulmonary hypertension.  
 \_\_\_\_\_ lesions that have been adequately corrected by surgery but continues to require medication for CHF such as diuretics, ACE inhibitors, beta-blockers or digoxin.  
 \_\_\_\_\_ cyanotic heart defect(s) AND decision for use of palivizumab was made with pediatric cardiologist consultation.
- Infant is younger than 2 years old on November 1, 2018 AND infant has undergone (or will undergo) cardiac transplantation during the RSV season (November 1, 2018 through March 31, 2019).
- Infant is 12 months old or younger (infant's first birthday is on or after November 1, 2018) AND infant has a congenital anatomic pulmonary abnormality or neuromuscular disease that impairs the ability to clear secretions from the upper airway because of ineffective cough.
- Infant is younger than 24 months old on November 1, 2018 AND infant will be profoundly immunocompromised during RSV season (November 1, 2018 through March 31, 2019) due to \_\_\_\_\_.

**Is the patient currently in the hospital?** \_\_\_\_\_ Yes \_\_\_\_\_ No

**Has the patient been in the hospital since the start of the current RSV season (November 1, 2018)?** \_\_\_\_\_ Yes \_\_\_\_\_ No

**If Yes, was a dose of palivizumab administered while patient was hospitalized?** \_\_\_\_\_ Yes \_\_\_\_\_ No **If Yes, please provide date** \_\_\_\_\_.

Prescribing Physician Signature:\* \_\_\_\_\_ Date: \_\_\_\_\_

\*(Signature stamps and proxy signatures are not acceptable)

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