

UCSF BCH Palivizumab (Synagis®)				
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Patient Information				
Patient Name:	DOB: UC Med Rec #:			
Address:	Sex: □M □ F			
City: State: Zip:	Home Phone: Language:			
Parent/Guardian:	Work/Cell:			
Statement of Medical Necessity/Clinical Information				
Primary Diagnosis: Gestational age at Birth:	gnosis: Gestational age at Birth: (weeks) Birth weight(kg)			
List of primary diagnosis is listed at the back of this page. Please select and circle the primary diagnosis from this list.				
Did the patient receive other medical therapies in last six mont	r Attendance □ Difficult Neonatal State of Asthma □ Other (explain): ironmental Pollutants/Tobacco Smoke hs? Ex; Bronchodilators, Steroids Oxygen (circle one) □ Y □ N  t: □ kgs □ lbs Date of Wt:			
Prescription and Orders  Synagis® (palivizumab) Is this the first dose? ☐ Yes ☐ No If no, date first dose given:				
Synagis (palivizumab) is this the first dose?   Yes   No if no, date first dose given:  Start ASAP   Next dose due:				
Administer 15mg/kg monthly IM November - March				
Pediatric Anaphylaxis				
Epinephrine 0.01ml/kg (max 0.3ml) of 1:1000 SUB Q or IM, and con	tact EMS or physician, as appropriate.			
Prescribing Physician Information				
Name (printed):	License #			
*PHYSICIAN SIGNATURE:	DATE:			

## Eligible for a Maximum of 5 doses (Age at onset of RSV season):

<sup>\*</sup>Premature infants <29 weeks 0 days gestational age

<sup>\*</sup>Infants/children <24 months of age who are profoundly immunocompromised

<sup>\*</sup>Infants/children ≤12 months of age with chronic lung disease of prematurity (CLDP) defined as gestational age <32 weeks, 0 days and required >21% oxygen for at least the first 28 days after birth

<sup>\*</sup>Infants/children in **second year of life** who continue to require treatment for chronic lung disease during the 6-month period before start of the 2<sup>nd</sup> RSV season with one of the following (supplemental oxygen, diuretic or chronic corticosteroid therapy)

<sup>\*</sup>Infants/children <12 months of age with hemodynamically significant congenital heart disease (CHD) who meet any of the following criteria: (acyanotic CHD receiving medication to control congestive heart failure and will require cardiac surgical procedures, moderate-to-severe pulmonary hypertension, cyanotic heart disease when decision is made in consultation with a pediatric cardiologist)

<sup>\*</sup>Infants/children ≤12 months of age with cystic fibrosis and with one of the following present (evidence of CLD or nutritional compromise)

<sup>\*</sup>Children in **second year of life** with cystic fibrosis with manifestations of severe lung disease and one of the following (previous hospitalization for pulmonary exacerbation in the first year of life, abnormalities on chest radiography or chest computed tomography that persists when stable, weight for length less than 10<sup>th</sup> percentile)

<sup>\*</sup>Infants/children ≤12 months of age with either congenital abnormalities of the airway or neuromuscular disease that comprises mobilization of respiratory secretions

	ICD10 codes for sy	nagis if criter	ia are met
Chronic pulmor	nary heart disease	Other congen	ital anomalies of circulatory system
127.0	Primary pulmonary hypertension	Q25.0	Patent ductus arteriosus
127.1	Kyphoscoliotic heart disease		Coarctation of aorta
127.2	Other secondary pulmonary hypertension/pulmonary hypertention NOS	Q25.1	Coarctation of aorta (preductal) (postductal)
127.82	Chronic pulmonary embolism	Q25.21	Interruption of aortic arch
127.89	Other specified pulmonary heart diseases	Q25.29	Other atresia of Aorta
150.9	Heart failure, unspecified	Q25.3	Supravalvular aortic stenosis
		Other congen	ital malformations of aorta
Chronic obstructive pulmonary disease and allied conditions		Q25.40	Congenital malformation of aorta unspecified
	Chronic Bronchitis	Q25.41	Absence and aplasia of aorta
J41.0	Simple chronic bronchitis	Q25.42	Hypoplasia of aorta
J41.1	Mucopurulent chronic bronchitis	Q25.42	Congenital aneurysm of aorta
041.1	Obstructive chronic bronchitis	Q25.44	Congenital dilation of aorta
J44.0	With acute bronchitis	Q25.45	Double aortic arch
J44.1	With (acute) exacerbation	Q25.46	Tortuous aortic arch
J44.9	Without exacerbation	Q25.47	Right aortic arch
J42	Unspecified chronic bronchitis	Q25.48	Anomalous origin of subclavian artery
UTL	Chronic obstructive asthma	Q25.49	Other congenital malformations of aorta
J44.0	With status asthmaticus	Q25.5	Atresia of pulmonary artery
J44.0	With (acute) exacerbation	Q25.6	Stenosis of pulmonary artery
J44.9	Chronic airway obstruction, not elsewhere classified	Q25.71	Coarctation of pulmonary artery
	nomialies & anomalies of cardiac septal closure	Q25.71	Congenital pulmonary arteriovenous malformation
	al anomalies of heart	Q25.72	Other congenital malformations of pulmonary artery
-	s of circulatory system	Q25.15	Anomalies of great veins
Q20.0	Common Arterial trunk	Q26.2	Total anomalous pulmonary venous connection
Q20.0	Transpostion of great vessles	Q26.3	Partial anomalous pulmonary venous connection
Q20.1	Double outlet right ventricle	Q26.9	Congenital malformation of great vein, unspecified
Q20.1	Double outlet left ventricle	Q27.0	Congenital absence and hypoplasia of umbilical artery
Q20.2 Q20.3	Discordant ventriculoarterial connection	Q33.3	Agenesis of lung
Q20.4	Common ventricle	Q33.6	Congenital hypoplasia and dysplasia of lung
Q20.4 Q20.5	Discordant atrioventricular connection	Q00.0	Other anomalies of lung
Q21.0	Ventricular septal defect	Q33.4	Congenital bronchiectasis
Q21.1	Ostium secundum type atrial septal defect	Q33.8	Other congenital malformation of lung
Q21.3	Tetralogy of Fallot	Q33.9	Congenital malformation of lung, unspecified
42	Endocardial cushion defects	400.0	Weeks of gestation (when criteria met)
Q21.2	Endocardial cushion defects	P07.21	Less than 23 completed weeks
Q21.2	Ostium primum defect	P07.22	23 completed weeks
Q20.8	Cor biloculare	P07.23	24 completed weeks
Q20.8	Other	P07.24	25 completed weeks
Q21.9	Unspecified defect of septal closure	P07.25	26 completed weeks
	al anomalies of heart	P07.26	27 completed weeks
	Anomalies of pulmonary valve	P07.31	28 completed weeks
Q22.0	Atresia, congenital	P07.32	29 completed weeks
Q22.1	Stenosis, congenital	P07.33	30 completed weeks
Q22.2	Congenital pulmonary valve insufficiency	P07.34	31 completed weeks
Q22.4	Congenital tricuspid atresia and stenosis	P07.35	32 completed weeks
Q22.5	Ebstein's anomaly	P07.36	33 completed weeks
Q23.0	Congenital stenosis of aortic valve	P07.37	34 completed weeks
Q23.1	Congenital insufficiency of aortic valve	P07.38	35 completed weeks
Q23.2	Congenital mitral stenosis	P07.39	36 completed weeks
Q23.3	Congenital mitral insufficiency		
Q23.4	Hypoplastic left heart syndrome	P27.1	Bronchopulmonary dysplasia originating in the perinatal period
		P27.8	Other chronic respiratory diseases originating in the perinatal period
Other specified	anomalies of heart	P27.9	Unspecified chronic respiratory disease originating in the perinatal period
Q24.1	Subaortic stenosis		, ,
Q24.2	Cor triatriatum	Others - (Plea	se indicate ICD10 CM code & accurate diagnosis)
Q24.3	Infundibular pulmonic stenosis	(	
	Coronary artery anomaly		
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Q24.5 Q24.6	Congenital heart block		
Q24.5 Q24.6 Q24.8	Congenital heart block Obstructive anomalies of heart, not elsewhere classified		