

Respiratory Syncytial Virus Enrollment Form

| | |
|--|------------------------------------|
| Phone: 1-800-454-3730 Fax referral to: 1-844-490-4870 | Date: _____ Need-by date: _____ |
|--|------------------------------------|

Ship to: Patient Office Other:

SECTION I — member and provider information

| | |
|--|-------------------------|
| 1. Member name (last, first, middle initial) | |
| 2. Member identification number | 3. Member DOB |
| 4. Prescriber name | 5. Prescriber NPI |
| 6. Prescriber address (street, city, state, ZIP+4) | |
| 7. Prescriber phone number | |
| 8. Billing provider name | 9. Billing provider NPI |

SECTION II — clinical information for all prior authorization requests

| | | |
|--|------------------------|----|
| 10. Was Synagis® administered when the child was hospitalized? <input type="checkbox"/> Yes <input type="checkbox"/> No | | |
| If yes, indicate the date(s) of administration in the space(s) provided. (No more than five doses will be authorized, inclusive of any hospital-administered doses.) | | |
| 1. | 2. | 3. |
| 11. Current weight — child (in kilograms) | 12. Date child weighed | |
| 13. Calculated dosage of Synagis (15 milligrams per kilogram of body weight) | | |
| 14. Case-specific diagnosis/ICD-10 | | |

Providers are required to complete *one* of Section III A, III B, III C, III D, III E or III F (depending on the child's medical condition) for a prior authorization request to be considered for approval.

SECTION III A — clinical information for chronic lung disease

| | |
|--|-----------------------------------|
| 15. The child has chronic lung disease of prematurity. <input type="checkbox"/> Yes <input type="checkbox"/> No | |
| 16. Did the child require oxygen at greater than 21 percent for at least the first 28 days after birth? <input type="checkbox"/> Yes <input type="checkbox"/> No | |
| 17. Indicate the child's gestational age at delivery (in weeks and days). | |
| Weeks | Days |
| 18. Check all therapies below the child has continuously used over the past six months. | |
| <input type="checkbox"/> Corticosteroid | <input type="checkbox"/> Diuretic |
| <input type="checkbox"/> Supplemental oxygen | |

SECTION III B — clinical information for congenital heart disease

19. The child is younger than 12 months of age at the start of the respiratory syncytial virus (RSV) season and has hemodynamically significant congenital heart disease. Yes
 No

SECTION III C — clinical information for cardiac transplant

20. The child is younger than 24 months of age at the start of the RSV season and is scheduled to undergo a cardiac transplantation during the RSV season. Yes
 No

SECTION III D — clinical information for pre-term infants

21. The child is younger than 12 months of age at the start of the RSV season and was born before 29 weeks' gestation (i.e., zero days through 28 weeks, six days).
 Yes No

Indicate the child's gestational age at delivery (in weeks and days).

Weeks

Days

SECTION III E — clinical information for pulmonary abnormalities and neuromuscular disease

22. The child is younger than 12 months of age at the start of the RSV season and has a neuromuscular disease or congenital abnormality that impairs the ability to clear secretions from the upper airway because of an ineffective cough. Yes
 No

If yes, indicate the disease or anomaly.

SECTION III F — clinical information for immunocompromised children

23. The child is younger than 24 months of age at the start of the RSV season and is profoundly immunocompromised due to the following:

| | | |
|---------------------------|------------------------------|-----------------------------|
| a. Solid organ transplant | <input type="checkbox"/> Yes | <input type="checkbox"/> No |
| b. Stem cell transplant | <input type="checkbox"/> Yes | <input type="checkbox"/> No |
| c. Receiving chemotherapy | <input type="checkbox"/> Yes | <input type="checkbox"/> No |
| d. AIDS | <input type="checkbox"/> Yes | <input type="checkbox"/> No |
| e. Other | <input type="checkbox"/> Yes | <input type="checkbox"/> No |

If other, indicate the cause of the child's immunodeficiency:

SECTION IV — authorized signature

24. Prescriber signature

25. Date signed

SECTION V — additional information

26. Indicate any additional information in the space provided. Additional diagnostic and clinical information explaining the need for the product requested may be included here.
