

**SURVEILLANCE OF CONGENITAL RUBELLA SYNDROME (CRS) - CASE INVESTIGATION FORM**  
 EPIDEMIOLOGY UNIT, MINISTRY OF HEALTH

The Medical Officer/Hospital and REE/MOH should carry out the investigation personally. Necessary data should be obtained from the mother of the new baby/BHT/Physician/investigation reports/diagnosis cards. Early investigation and return is essential.

Serial No:   /   /   /

**A. GENERAL**

1. Date of notification to MOH :   /   /     (dd/mm/yy)
2. Date of notification to Epidemiology Unit :   /   /     (dd/mm/yy)
3. Name of the reporting Institution / Hospital .....
4. Ward No: .....
5. BHT No: .....
6. Name of the hospital where the baby was born .....
7. Ward No: .....
8. BHT No: .....

**B. PARTICULARS OF PATIENT (Please (✓) appropriate box where applicable)**

9. Name of patient (BLOCK LETTERS) .....
10. Name of the parent/guardian .....
11. Residential Address: .....
12. Date of Birth :   /   /     (dd/mm/yy)

13. Age <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> Yrs      Months      Days	14. Sex <input type="checkbox"/> 1. Male <input type="checkbox"/> 2. Female	15. Ethnic group <input type="checkbox"/> 1. Sinhalese <input type="checkbox"/> 2. Tamil <input type="checkbox"/> 3. Moor <input type="checkbox"/> 4. Others <input type="checkbox"/> 9. Unknown	16. Mother's occupation ..... <li>17. District .....</li> <li>18. MOH area .....</li>
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**B. PRESENT ILLNESS /OUTCOME**

19. Date of detection of signs and symptoms of CRS: <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> d   d      m   m      y   y  20. Where did the patient detect first <input type="checkbox"/> 1. Government hospital <input type="checkbox"/> 2. Private hospital <input type="checkbox"/> 3. Medical Officer of Health <input type="checkbox"/> 4. Private practitioner <input type="checkbox"/> 5. Ayurvedic institution <input type="checkbox"/> 6. Other (specify) .....	21. Outcome of the event <input type="checkbox"/> 1. Still under treatment <input type="checkbox"/> 2. Died <input type="checkbox"/> 3. Transferred <input type="checkbox"/> 4. Discharged  22. Date of discharge, transfer or death (where relevant) <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> d   d      m   m      y   y	23. If transferred, name of hospital ..... <li>24. Was patient transferred from some other hospital                  Yes <input type="checkbox"/> / No <input type="checkbox"/></li> <li>25. If "yes", where was the patient transferred from ? .....</li>
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**C. CLINICAL DATA**

**Surveillance Case definition:**

Child <1 year of age with maternal history of Rubella infection and/or following signs and symptoms.

List A <input type="checkbox"/> 1. Cataract/s <input type="checkbox"/> 2. Congenital glaucoma <input type="checkbox"/> 3. Congenital heart disease <input type="checkbox"/> 4. Loss of hearing <input type="checkbox"/> 5. Pigmentary Retinopathy	List B <input type="checkbox"/> 1. Purpura <input type="checkbox"/> 2. Splenomegaly <input type="checkbox"/> 3. Microcephaly <input type="checkbox"/> 4. Mental Retardation <input type="checkbox"/> 5. Meningo-encephalitis <input type="checkbox"/> 6. Radiolucent bone disease <input type="checkbox"/> 7. Jaundice (within 24hr of delivery)	Laboratory data consistent with Congenital Rubella Infection (CRI) <input type="checkbox"/> positive result of rubella IgM	<b>For office use only</b> Compatible with the case definition. <input type="checkbox"/> 1. Yes <input type="checkbox"/> 2. No
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**D. LABORATORY FINDINGS**

26. Was blood taken for rubella serological investigations?  1. yes  2. no if no reason .....
27. Was specimens collected for rubella virus isolation ?  1. yes  2. no if no reason .....
28. If yes:

Investigation	Date of collection of specimen (dd/mm/yy)	Laboratory MRI/ other govt./ private/ not known	Results (mark NA if test results are not available and RP if pending)
1. maternal IgG persisting >6/52 in infant			
2. rubella specific IgM			
3. virus isolation / PCR			

**E. MATERNAL HISTORY**

29. Age of mother at time of delivery:  
  years
30. Did the mother have a rubella-like illness during the present pregnancy?  
 1. yes  
 2. no  
 3. not known
31. If yes, period of gestation at the time of illness  
  in weeks  
 not known
32. Which of the following symptoms and signs were present?  
 1. fever  
 2. rash  
 3. lymphadenopathy  
 4. conjunctivitis  
 5. arthritis/arthralgia  
 6. others (specify) .....
33. Was rubella serologically confirmed during pregnancy ?  
 1. yes  
 2. no  
 3. not known

**F. MOTHER'S IMMUNIZATION HISTORY**

34. Was the mother immunized for rubella?  
 1. yes  2. no  3. not known
35. If yes, date of vaccination:  
       
 d d m m y y  
 not known
36. Type of vaccine used:  
 1. Rubella  3. MR  
 2. MMR  4. Not known
37. Place of vaccination  
 1. MOH clinic  
 2. school  
 3. government hospital  
 4. private dispensary/surgery  
 5. private hospital  
 6. other (specify) .....
7. not known
38. If not immunized, reason:  
 1. medical contraindication  
 2. unaware of the need for vaccination  
 3. non-availability of vaccine  
 4. no faith in the vaccine  
 5. others (specify) .....
6. not known

**G. CONTACT HISTORY**

39. Was the mother in contact with a known or suspected case of rubella during the index pregnancy?  
 1. yes  
 2. no  
 3. not known
40. If yes, period of gestation in weeks:  
   
 not known

**FOR OFFICE USE**  
 Time between immunization and development of maternal infection  
  yrs   months

41. Remarks: .....

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Signature: ..... Name:.....

Date: ..... Designation: .....

For office use only	
<b>Final classification</b>	
Laboratory confirmed	<input type="checkbox"/>
Clinically confirmed	<input type="checkbox"/>
C R I	<input type="checkbox"/>