

Rubella Surveillance Worksheet

NAME _____ (last) (first)		ADDRESS (Street and No.) _____		Phone _____	Hospital Record No. _____											
This information will not be sent to CDC																
REPORTING SOURCE TYPE NAME _____ <input type="checkbox"/> physician <input type="checkbox"/> PH clinic ADDRESS _____ <input type="checkbox"/> nurse <input type="checkbox"/> laboratory ZIP CODE _____ <input type="checkbox"/> hospital <input type="checkbox"/> other clinic PHONE (____) _____ <input type="checkbox"/> other source type _____			SUBJECT ADDRESS CITY _____ SUBJECT ADDRESS STATE _____ SUBJECT ADDRESS COUNTY _____ SUBJECT ADDRESS ZIP CODE _____ LOCAL SUBJECT ID _____													
CASE INFORMATION																
Date of Birth ____-____-____ <small>month day year</small>		Sex M=male F=female <input type="checkbox"/>		Ethnic Group H=Hispanic/Latino N=Not Hispanic/Latino O=Other _____ U=Unknown <input type="checkbox"/>												
Race <input type="checkbox"/> American Indian/Alaskan Native <input type="checkbox"/> Asian <input type="checkbox"/> Black/African American <input type="checkbox"/> Native Hawaiian/Pacific Islander <input type="checkbox"/> White <input type="checkbox"/> Not asked <input type="checkbox"/> Refused to answer <input type="checkbox"/> Other <input type="checkbox"/> Unknown																
Country of Birth _____		Other Birth Place _____		Country of Usual Residence _____												
Age at Case Investigation _____		Age Unit* _____		Reporting County _____												
Reporting State _____																
Date Reported ____-____-____ <small>month day year</small>		Date First Reported to PHD ____-____-____ <small>month day year</small>		National Reporting Jurisdiction ____												
Date First Reported to County ____-____-____ (mm/dd/yyyy)		Earliest Date Reported to State ____-____-____ (mm/dd/yyyy)														
CASE INVESTIGATION STATUS CODE		<table border="1" style="width:100%; border-collapse: collapse;"> <tr> <td>Approved</td> <td>Deleted</td> <td>Notified</td> <td>Ready for review</td> <td>Reviewed</td> <td rowspan="2">Unknown</td> </tr> <tr> <td>Closed</td> <td>In progress</td> <td>Other (specify) _____</td> <td>Rejected</td> <td>Suspended</td> </tr> </table>				Approved	Deleted	Notified	Ready for review	Reviewed	Unknown	Closed	In progress	Other (specify) _____	Rejected	Suspended
Approved	Deleted	Notified	Ready for review	Reviewed	Unknown											
Closed	In progress	Other (specify) _____	Rejected	Suspended												
Case Class Status <input type="checkbox"/> Suspected <input type="checkbox"/> Confirmed <input type="checkbox"/> Unknown <input type="checkbox"/> Probable <input type="checkbox"/> Not a case		Case Investigation Start Date ____-____-____ (mm/dd/yyyy)														
CASE DETECTION METHOD	Laboratory report		Prenatal testing	Provider reported	Self-referral											
	Other _____		Prison entry screening	Routine physical	Unknown											
Confirmation Date ____-____-____ <small>month day year</small>																
CASE CONFIRMATION METHOD	Active surveillance		Lab diagnosis		No information given											
	Case/outbreak investigation		Lab reporting		Occupational disease surveillance											
	Clinical diagnosis		Local/state specified		Other (specify) _____											
	Epi-linked		Medical records review		Provider certified											
CLINICAL INFORMATION																
SIGNS/SYMPTOMS		Y	N	U												
Rash																
Onset Date		Duration		Age at Onset	Age Type Units*											
____-____-____ <small>month day year</small>		____ (days)		____	____											
Fever		Highest Measured Temperature		Tempertaure Units												
____-____-____ <small>month day year</small>		____.____		°Cel <input type="checkbox"/> °F <input type="checkbox"/>												
*Units a = year d = day mo = month wk = week unk = unknown																
Arthralgia		Y	N	U												
Conjunctivitis		Y	N	U												
Arthritis		Y	N	U												
Lymphadenopathy		Y	N	U												
Other _____		Y	N	U												
Unknown		Y	N	U												
ILLNESS		Onset Date		End Date	Diagnosis Date											
		____-____-____ <small>month day year</small>		____-____-____ <small>month day year</small>	____-____-____ <small>month day year</small>											
		Duration		Illness Duration Units*												
		____ (days)		____												
HOSPITALIZATION		Hospitalized?		Admit Date	Discharge Date											
		Y=yes N=no U=unknown <input type="checkbox"/>		____-____-____ <small>month day year</small>	____-____-____ <small>month day year</small>											
		Duration		Pregnancy Status												
		____ (days)		<input type="checkbox"/> Y=yes <input type="checkbox"/> N=no <input type="checkbox"/> U=Unknown												
INV920 Y N U Y N U Y N U																
COMPLICATIONS		Encephalitis		Other	Death?											
Thrombocytopenia		Unknown		Cause of Death _____												

PREGNANCY INFORMATION

Expected Delivery Date _____ (mm/dd/yyyy) Expected Place of Delivery _____

Trimester at onset of illness? ☐ First ☐ Second ☐ Third ☐ Unknown Number of weeks gestation at onset? ☐ ☐

Is there documentation of previous immunity testing? Y=yes N=no U=unknown ☐ Age at time of previous testing? ☐ ☐

Previous Immunity Testing Result

Positive	Significant rise in IgG
Negative	No significant rise in IgG
Indeterminate	Other
Pending	Not done
Unknown	

Year of previous rubella immunity test? ☐ ☐ ☐ ☐

Diagnosed with the condition before? Y=yes N=no U=Unknown ☐

Previous disease serologically confirmed? Y=yes N=no U=unknown ☐

Year of previous disease? _____ Age at previous diagnosis? ☐ ☐

Previous case diagnosed by: ☐ physician/healthcare provider ☐ parent ☐ other _____ Age Units† _____

†UNITS a = year d = day mo = month wk = week unk = unknown

PREGNANCY OUTCOME

What was the outcome of current pregnancy? ☐ Live birth with CRS ☐ Other _____ ☐ Unknown Autopsy Result _____

Age of fetus at time of pregnancy cessation: ☐ ☐ (weeks) Was an autopsy performed? Y=yes N=no U=unknown ☐

EXPOSURE AND IMPORTATION INFORMATION

Did symptom onset occur within 14-23 days of entering U.S. following travel or living outside the U.S.? Y=yes N=no U=unknown ☐

International Destination(s) of Recent Travel _____ Travel Return Date _____ (mm/dd/yyyy)
 _____ Travel Return Date _____ (mm/dd/yyyy)

Length of time in the U.S. since last travel: ☐ ☐ ☐ Length of time in U.S. units†: ☐ ☐ ☐

Country of Exposure _____ State or Province of Exposure _____

County of Exposure _____ City of Exposure _____

Import Status – US-Acquired 1=import-linked case 2=imported virus case 3=endemic case 4=unknown source case 5=other ☐

CASE DISEASE IMPORTED CODE

Indigenous	In state, out of jurisdiction
International	Yes, imported, but not able to determine source state/country
Out of state	Unknown

Imported Country _____

Imported State _____

Traceable to an international import? Y=yes N=no U=unknown ☐ Imported County _____ Imported City _____

TRANSMISSION SETTING

Athletics	Day care center	Hospital outpatient clinic	Other (specify)
College	Doctor's office	Hospital ward	Place of worship
Community	Home	International travel	School
Correctional facility	Hospital ER	Military	Work
			Unknown

Age & setting of case verified? Y=yes N=no U=unknown ☐ Epi-linked to a confirmed or probable case? Y=yes N=no U=unknown ☐

Was case patient a healthcare provider? Y=yes N=no U=unknown ☐ Part of an outbreak? Y=yes N=no U=unknown ☐

COMMENTS

LABORATORY TESTING

VPD Lab Message Reference Laboratory _____

VPD Lab Message Patient Identifier _____

VPD Lab Message Specimen Identifier _____

Lab testing done to confirm diagnosis? Y=yes N=no U=unknown ☐

Was a specimen sent to CDC? Y=yes N=no U=unknown ☐

Was case laboratory confirmed? Y=yes N=no U=unknown ☐

Test Type	Test Result	Test Result Quantitative	Test Method	Result Units	Date Specimen Collected <small>month day year</small>	Date Specimen Sent to CDC <small>month day year</small>	Date Specimen Analyzed <small>month day year</small>	Specimen Source	Specimen Type	Performing Lab Type
IgM (capture)										
IgM										
IgG EIA (acute)										
IgG EIA (conv)										
culture										
PCR										
other										
unknown										
Ab IF										
Ab latex										
genotype										

TEST RESULTS CODES

P=positive N=negative
X=not done E=pending
I=Indeterminate
NS=no significant rise in titer
PS=significant rise in titer
U=unknown

SPECIMEN TYPE CODES

1=entire throat 6=entire eye
2=intervertebral space 7=pharyngeal
3=skin structure 8=other (specify)
4=mouth region 9=unknown
5=lens of eye 10=nasal cavity

PERFORMING LABORATORY TYPE CODES

1=CDC lab 5=public health lab
2=commercial lab 6=VPD testing lab
3=hospital lab 8=other (specify)
4=other clinical lab 9=unknown

GENOTYPE CODES

1a 1F 2A
1B 1g 2B
1C 1H 2c
1D 1I other
1E 1J unknown

SPECIMEN SOURCE

2=blood 3=body fluid 4=BAL 8=cataract 9=CSF 11=DNA sample 15=NP aspirate 16=NP swab 17=NP washings 18=nucleic acid 19=oral fluid
20=oral swab 21=plasma 22=RNA sample 23=saliva 25=serum 36=throat swab 38=urine 40=viral isolate 41=other 42=unknown

[illegible]

Reason Not Vaccinated Per ACIP			
1 = religious exemption	6 = too young	11 = vaccine record incomplete/unavailable	16 = immigrant
2 = medical contraindication	7 = parent/patient refusal	12 = parent/patient report of previous disease	
3 = philosophical objection	8 = other _____	13 = parent/patient unaware of recommendation	<input type="checkbox"/>
4 = lab evidence of previous disease	9 = unknown	14 = missed opportunity	<input type="checkbox"/>
5 = MD diagnosis of previous disease	10 = parent/patient forgot to vaccinate	15 = foreign visitor	

[illegible]

<p>VACCINE TYPE CODES</p> <p>03=MMR (measles, mumps, rubella virus)</p> <p>04=M/R (measles & rubella virus)</p> <p>05=Measles (measles virus) OTH=other</p> <p>06=Rubella (rubella virus) 998=no vaccine administered</p> <p>07=Mumps (mumps virus) 999=unknown</p> <p>38=Rubella/mumps (rubella & mumps virus)</p> <p>94=MMRV (measles, mumps, rubella, & varicella virus)</p>	<p>VACCINE MANUFACTURER CODES</p> <p>MSD = Merck</p> <p>OTH = other (specify)</p> <p>UNK = unknown</p>	<p>VACCINE EVENT INFORMATION SOURCE CODES</p> <p>00=new immunization record</p> <p>01=historical information, source unspecified</p> <p>02=historical information, other provider 11=imm. info system (IIS)</p> <p>05=historical information, other registry OTH=other (specify)</p> <p>06=historical information, birth certificate UNK=unknown</p> <p>07=historical information, school record</p> <p>08=historical information, public agency</p> <p>09=historical information, patient or parent recall</p> <p>10=historical information, patient or parent written record</p>
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VACCINE HISTORY COMMENTS

CASE NOTIFICATION

CONDITION CODE		10200		Immediate National Notifiable Condition Y=yes N=no U=unknown <input type="checkbox"/>			Legacy Case ID _____	
State Case ID _____		Local Record ID _____		Jurisdiction Code ____		Binational Reporting Criteria _____		
Date First Verbal Notification to CDC ____ ____ ____ month day year					Date Report First Electronically Submitted ____ ____ ____ month day year			
Date of Electronic Case Notification to CDC ____ ____ ____ (mm/dd/yyyy)						MMWR Week _____		MMWR Year _____
Notification Result Status		<input type="checkbox"/> Final results		<input type="checkbox"/> Record coming as correction		<input type="checkbox"/> Results cannot be obtained		
Person Reporting to CDC NAME _____ (first) _____ (last)				Person Reporting to CDC Email _____ @ _____ Person Reporting to CDC Phone No. (____) _____				
Current Occupation _____				Current Occupation Standardized _____				
Current Industry _____				Current Industry Standardized _____				

CLINICAL CASE DEFINITION [†]

SUSPECTED

Any generalized rash illness of acute onset that does not meet the criteria for probable or confirmed rubella or any other illness.

PROBABLE

In the absence of a more likely diagnosis, an illness characterized by all of the following:

- Acute onset of generalized maculopapular rash; **and**
- Temperature greater than 99.0° F or 37.2° C, if measured; **and**
- Arthralgia, arthritis, lymphadenopathy, or conjunctivitis; **and**
- Lack of epidemiologic linkage to a laboratory-confirmed case of rubella; **and**
- Noncontributory or no serologic or virologic testing.

CONFIRMED

A case with or without symptoms who has laboratory evidence of rubella infection confirmed by one or more of the following laboratory tests:

- Isolation of rubella virus; or
- Detection of rubella-virus specific nucleic acid by polymerase chain reaction; or
- IgG seroconversion[†] or a significant rise between acute- and convalescent-phase titers in serum rubella IgG antibody level by any standard serologic assay; or
- Positive serologic test for rubella IgM antibody^{†*}

OR

An illness characterized by all of the following:

- Acute onset of generalized maculopapular rash; and
- Temperature greater than 99.0°F or 37.2°C; and
- Arthralgia, arthritis, lymphadenopathy, or conjunctivitis; and
- Epidemiologic linkage to a laboratory-confirmed case of rubella.

[†] Not explained by MMR vaccination during the previous 6-45 days.

*Not otherwise ruled out by more specific testing in a public health laboratory

OTHER INFORMATION

Serum rubella IgM test results that are false positives have been reported in persons with other viral infections (e.g., acute infection with Epstein-Barr virus [infectious mononucleosis], recent cytomegalovirus infection, and parvovirus infection) or in the presence of rheumatoid factor. Patients who have laboratory evidence of recent measles infection are excluded.

[†]CSTE Position Statement 12-ID-09 at <https://wwwn.cdc.gov/nndss/conditions/rubella/case-definition/2013/>