Annex I

Integrated measles and rubella case investigation form

Recommended basic set of data for case-based reporting in national surveillance system

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Case ID: Region:	District
Date of notification:/Date of investigation	on://_ Date of report://
Initial clinical diagnosis: 1. Clinical measles □ 2. Clinical rubella □ 3. Others □ 9. Unknown □	
Outbreak-related: 1. Yes □ 2. No □ 9. Unk	nown □ Outbreak ID:
A. Personal data and immunization status	
Name*: *WHO Europe does not collect this information – please provide only Case ID number	
Sex: 1. Male □ 2. Female □ 9. Unknown □	
Date of birth:_//if not available, age in yearsor for younger than a year, age in months	
Address*: *WHO Europe does not collect this information	
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For female cases	
Is case pregnant? 1. Yes □ 2. No □ If ye	es, gestation age:weeks
Vaccination status	
Measles: 1. Yes □ 2. No □ 3. Unknown □ If ye	es, no. of doses Last vaccination date:
Source of vaccination status: 1. Medical record □ 2. Parent or guardian □//	
Rubella: 1. Yes □ 2. No □ 3. Unknown □ If ye	es, no. of doses Last vaccination date:
Source of vaccination status: 1 Medical record \Box 2	Parent or quardian \Box
Source of vaccination status: 1. Medical record □ 2. Parent or guardian □ □ □//	
B. Clinical information	
5. Cimical information	
Maculopapular rash 1. Yes □ 2. No. □ 9. Unknown □	
Date of rash onset://Duration of rash (c	lays):
Other symptoms	Presence of Yes □ No. □
	complications
Fever Yes No Unknown	Pneumonia Yes □ No □ Unknown □
Coryza Yes 🗆 No 🗆 Unknown 🗆	Malnutrition Yes □ No □ Unknown □
Cough Yes No Unknown	Diarrhoea Yes □ No □ Unknown □
Conjunctivitis Yes ☐ No ☐ Unknown ☐	Encephalitis Yes No Unknown
Adenopathy or arthralgia Yes □ No □ Unknown □ or arthritis	Other (specify) Yes □ No □ Unknown □
Hospitalized: 1. Yes □ 2. No □ 9. Unknown □ Name of hospital:	
Clinical outcome: 1. Dead: date of death//2. Survived □ 3. Lost to follow-up □ 9. Unknown □	
Cause of death:	

C. Epidemiological investigation

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Did the patient have contact with confirmed case of measles (within 7-18 days) or rubella (within 12–23 days) prior to rash onset? 1. Yes □ 2. No □ 9. Unknown □ If yes:	
Who (case ID/name):	
Where (country/address):	
When (dates):	
Were there confirmed cases of measles and/or rubella reported in the area prior to this case? 1. Measles □ 2. Rubella □ 3. Both □ 4. No □ 9. Unknown □	
Did the patient travel within 7–23 days before onset of rash? 1. Yes □ 2. No □ 9. Unknown □ If yes: Where (country/address): When (dates):	
When (dates): Travel details:	
Is the case epidemiologically linked to imported confirmed case? 1. Yes If yes: Who (case ID/name): Where (country/address): When (dates):	
Was the case in contact with a pregnant woman since development of the symptoms? 1. Yes □ 2. No □ 9. Unknown □ If yes, please provide name and address	
D. Laboratory investigation	
Specimen collected: 1. Yes □ 2.No □ 3. Unknown □ If yes, type of specimen: Serum □ Saliva/oral fluid □ Nasopharyngeal swab □ Dry blood spot □ Urine □ EDTA whole blood □ Other □	
Date of specimen collection:// Date specimen sent to lab://	
Measles IgM: Not tested □ Positive □ Negative □ In process □ Indeterminate □ Rubella IgM: Not tested □ Positive □ Negative □ In process □ Indeterminate □	
Date of laboratory result (first validated result)://	
Measles virus detection: Not tested □ Positive □ Negative □ In process □ Genotype Rubella virus detection: Not tested □ Positive □ Negative □ In process □ Genotype	
E. Final classification	
0 Discarded □	
1 Measles – laboratory-confirmed □ 2 Measles – epidemiologically linked □ 3 Measles – clinical □	
6 Rubella – laboratory-confirmed □ 7 Rubella – epidemiologically linked □ 8 Rubella – clinical □	
Source of infection: 1. Imported □ 2. Endemic □ 3. Import-related □ 9 Unknown □	
Date of final classification://	
Investigated by: Name	
Position:	