MEASLES BASICS

Measles infectious period
From four days before rash onset through four days after rash onset (day of rash onset is day 0).

Measles exposure
Sharing the same airspace with a person infectious with measles (during the 4 days prior through the 4 days after their rash onset), e.g., same classroom, home, clinic waiting room, airplane etc., or were in these areas up to 1 hour after the infectious person left the area. Although CDC recommends using a 2 hour window, there is only one report in the literature of measles transmission >60 minutes after an infectious person has left the setting.

No minimum time period has been established for exposure, but it is presumed that longer exposures are more likely to result in measles transmission than brief, transient exposures.

When exposures have occurred in venues in which it is not possible to identify individuals, it is helpful to notify local health care providers so that they can be on the alert for possible cases. In addition, some local health jurisdictions issue press releases to notify the public.

Measles incubation period
The period from exposure to onset of prodrome is generally 8–12 days. In family studies, the average interval between the appearance of rash in the index case and in subsequent cases is 14 days (range 7-21 days).

The course of measles infection
Measles typically begins with a mild to moderate fever accompanied by cough, coryza, and conjunctivitis. Two to three days later, Koplik's spots, a characteristic sign of measles, may appear. At this time the fever spikes, often ≥104°F. At the same time, a red blotchy maculopapular rash appears, usually first on the face, along the hairline and behind the ears. This rash rapidly spreads downward to the chest and back and finally, to the thighs and feet. In approximately one week, the rash fades in the same sequence that it appeared.

Measles laboratory criteria for diagnosis
- Preferred: Detection of viral RNA by reverse transcription polymerase chain reaction (RT- PCR); or
- Acceptable: Serum* measles IgM antibody positive
- Acceptable: Isolation of measles virus; or
- Acceptable: Significant rise in serum* measles IgG 
  antibody between acute and convalescent titers.

*Capillary blood (finger or heel stick) can be used for serologic testing if venous blood cannot be obtained.

Please send specimens to a public health lab for testing; use of commercial labs may delay testing.

More information on testing, including capillary blood collection, is available at: http://www.cdph.ca.gov/HealthInfo/discond/Documents/CDPHMeaslesLabTesting2011-01.pdf

ASSESSING SUSPECT MEASLES CASES

- Consider measles in patients of any age who have a fever AND a rash
- In measles cases there must be some fever, even subjective fever, and the rash must start on the head or neck.
- Patients with measles usually have at least 1 or 2 of the “3 Cs” – cough, coryza and conjunctivitis.
- If measles is being considered, please contact your local health department immediately, see http://www.cdph.ca.gov/programs/cclho/Documents/LHD_CD_Contact_Info.doc


ASSESSING MEASLES CONTACTS
(refer to Table for more detail)

Immunity to measles
Contacts who are not classified as high-risk† can be presumed to be immune to measles for the purposes of measles case investigations if they:
- were born prior to 1957; or
- have written documentation with dates of receipt of at least one dose of measles-containing vaccine given on or after their first birthday in 1968 or later; or
- have documented IgG+ test for measles; or
- laboratory confirmation of previous disease; or
- served in the U.S. armed forces; or
- were born in the U.S. in 1970 or later and attended a U.S. elementary school;‡ or
- entered the U.S. in 1996 or later with an immigrant visa or have a green card.‡
† Additional evidence of immunity is required for exposed high-risk persons, e.g., healthcare personnel of any age, pregnant women, immunocompromised people, household contacts of a case, or persons in settings with known unvaccinated persons (e.g., infant care settings). Additional evidence of immunity may also be required during an outbreak. Immunity can be presumed if the exposed person:
  o has documentation of a positive measles IgG test; or
  o has documentation of two doses of measles vaccine given in 1968 or later, separated by at least 28 days, with the first dose on or after the first birthday

‡ Unless known to be unvaccinated for measles, e.g., having a medical contraindication to vaccination or being philosophically or religiously opposed to vaccinations.

PUBLIC HEALTH FOLLOW-UP

**MMR vaccine for postexposure prophylaxis**

MMR vaccine may be given <72 hours of last exposure to persons ≥6 months of age (although administration of IG is preferred in infants 6-11 months of age) with 1 or no documented doses of MMR, if not contraindicated.

**Immunoglobulin (IG) for postexposure prophylaxis**

IG may be given to exposed susceptible people ≤6 days of last exposure to prevent infection. However, susceptible persons who receive IG >6 days after the first exposure (when there are multiple exposure dates), should still be placed on quarantine.

- Infants <12 months of age should receive 0.5 mL/kg of body weight of intramuscular IG (IGIM); maximum dose = 15 mL.
- Unvaccinated children <30 kg (66 lbs) who are not eligible for MMR PEP should receive 0.5 mL/kg of body weight of IGIM; maximum dose = 15 mL.
- Pregnant women without evidence of measles immunity should receive 400 mg/kg of intravenous IG (IGIV).
- Severely immunocompromised persons‡, irrespective of evidence of measles immunity, should receive 400 mg/kg of IGIV.
- For persons already receiving IGIV therapy, administration of ≥400 mg/kg body weight at least one time in the <3 weeks before first measles exposure should be sufficient to prevent measles infection.
- For patients receiving subcutaneous IG (IGSC) therapy, administration of at ≥200 mg/kg body weight once weekly for 2 consecutive weeks before first measles exposure should be sufficient.
- Persons weighing >30 kg (66 lbs) will not receive an adequate dose of measles antibodies from IGIM. Therefore, there is no recommendation to administer IGIM to such persons.
- When there are multiple exposure dates, exclusion/quarantine should be recommended through day 21 of the last exposure.

- CDPH does not recommend extending quarantine or exclusion beyond 21 days after exposure in persons who received IG PEP. It is unknown if IG prolongs the incubation period. However, symptoms should be monitored for an additional 7 days and if symptoms occur ≤28 days of exposure, persons who received IG should self-isolate and contact their local health department.
- Nonimmune persons who received IG should not receive MMR vaccine earlier than 6 months after IGIM or 8 months after IGIV administration.
- Information on IG administration is available at: [http://www.cdph.ca.gov/HealthInfo/discond/Documents/CDPHIGforMeaslesPEP.pdf](http://www.cdph.ca.gov/HealthInfo/discond/Documents/CDPHIGforMeaslesPEP.pdf)
- One source of IG is FFF Enterprises in Temecula CA, which can be reached 24/7 at: 1-800-843-7477.

§ Per CDC and IDSA guidance: Patients with high-level immunosuppression include those:

- with combined primary immunodeficiency disorder (e.g., severe combined immunodeficiency);
- receiving cancer chemotherapy;
- on treatment for ALL within and until at least 6 months after completion of immunosuppressive chemotherapy;
- within 2 months after solid organ transplantation;
- who have received a bone marrow transplant until at least 12 months after finishing all immunosuppressive treatment, or longer in patients who have developed graft-versus-host disease;
- with HIV infection with a CD4 T-lymphocyte count <200 cells/mm³ (age >5 years) and percentage <15 (all ages) (some experts include HIV-infected persons who lack recent confirmation of immunologic status or measles immunity);
- receiving daily corticosteroid therapy with a dose ≥20 mg (or >2 mg/kg/day for patients who weigh <10 kg) of prednisone or equivalent for ≥14 days; and
- receiving certain biologic immune modulators, that is, a tumor necrosis factor-alpha (TNF-α) blocker or rituximab.

After HSCT, duration of high-level immunosuppression is highly variable and depends on type of transplant (longer for allogeneic than for autologous), type of donor and stem cell source, and post-transplant complications such as graft versus host disease (GVHD) and their treatments.

Please contact CDPH for consultation.

**Home quarantine/symptom watch period**

If quarantine implemented, it should Day 7 (CDC recommends day 5 for healthcare workers) after first exposure through day 21 after last exposure (day of exposure is day 0).

If symptoms consistent with measles develop, patient should be immediately isolated through day 4 after rash onset (day of rash onset is day 0). Exposed people should be instructed to isolate themselves and notify their local health department if symptoms occur.
EXPOSURES IN SPECIAL SETTINGS

Measles exposure in a childcare setting with infants

Infant childcare settings are considered high risk because unvaccinated infants can develop severe measles.

When there is a measles exposure in such a setting:

- Administer IGIM to unvaccinated infants <12 months of age:
  - Infants who receive IG ≤6 days of first exposure should be protected against that exposure and have ongoing protection for some time and do not need to be excluded from childcare or quarantined, but should not visit other settings with high-risk persons.
  - Infants who receive IGIM >6 days after first exposure should be excluded from childcare and quarantined from day 7 after first exposure through day 21 after last exposure.

- Administer MMR or IGIM to unvaccinated children >12 months of age:
  - Administer MMR vaccine if <72 hours of last exposure to measles or administer IGIM to children ≤6 lbs if ≥72 hours but ≤6 days of last exposure to measles.
  - Children who receive MMR <72 hours of first exposure or IGIM ≤6 days of first exposure should be protected against that exposure and have ongoing protection for some time and do not need to be excluded from childcare or quarantined, but should not visit other settings with high-risk persons.
  - Children who receive MMR vaccine ≥72 hours or IGIM >6 days after first exposure should be excluded from childcare and quarantined from day 7 after first exposure through day 21 after last exposure.

- Administer the second dose of MMR to children with 1 documented MMR dose (if >28 days from last dose):
  - Children who have received 1 dose of MMR vaccine ≥2 weeks before first exposure may attend the childcare setting, but should not visit other settings with high-risk persons. Performing measles IgG testing on such children to determine immunity can also be considered.

Other recommendations:

To prevent the possibility of sustained transmission, new enrollees who are susceptible to measles and have not yet attended should not start until after the end of the incubation period.

Please consult CDPH at 510-620-3737 for additional discussion on measles exposures in childcare settings.

Measles flight investigations

If a suspect measles case reports air travel during their infectious period, please do the following:

- Collect as much information about the flight as possible such as the flight number, times, destinations, and seat number.
- Collect information on whether there were traveling companions.

MEASLES TREATMENT

No specific antiviral therapy is available for measles. Measles virus is susceptible in vitro to ribavirin, which has been given by the intravenous and aerosol routes to treat severely affected and immunocompromised children with measles. However, no controlled trials have been conducted, and ribavirin is not approved by the U.S. Food and Drug Administration for treatment of measles. IV ribavirin (Virazole®) is available in the U.S. under an emergency investigational new drug protocol. Please contact CDPH if this product is requested.

Vitamin A. Vitamin A treatment of children with measles in developing countries has been associated with decreased morbidity and mortality rates. Low serum concentrations of vitamin A also have been found in children in the United States, and children with more severe measles illness have lower vitamin A concentrations.

The World Health Organization currently recommends vitamin A for all children with acute measles, regardless of their country of residence. Vitamin A for treatment of measles is administered once daily for 2 days, at the following doses:

- 200 000 IU for children 12 months or older;
- 100 000 IU for infants 6 through 11 months of age; and
- 50 000 IU for infants younger than 6 months.
- An additional (i.e., a third) age-specific dose should be given 2 through 4 weeks later to children with clinical signs and symptoms of vitamin A deficiency.

Even in countries where measles usually is not severe, vitamin A should be given to all children with severe measles (e.g., requiring hospitalization).

Aquasol A™ appears to be the only parenteral vitamin A product currently available in the United States.
### TABLE. RECOMMENDED FOLLOW-UP OF MEASLES CONTACTS

<table>
<thead>
<tr>
<th>Measles immunity assessment for low-risk contacts (NOT immunocompromised, infant &lt;12 months, pregnant, healthcare worker or household contact)</th>
<th>IgG testing</th>
<th>MMR PEP</th>
<th>IG PEP</th>
<th>Quarantine if no PEP</th>
<th>Exclusion if no PEP</th>
<th>Symptom watch</th>
</tr>
</thead>
<tbody>
<tr>
<td>Two documented doses of MMR vaccine (~1% will be susceptible)</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>Passive</td>
</tr>
<tr>
<td>Known to be measles IgG positive (&lt;1% will be susceptible)</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>Passive</td>
</tr>
<tr>
<td>Born before 1957 (5% will be susceptible)</td>
<td>If desired</td>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>Yes</td>
<td>Passive</td>
</tr>
<tr>
<td>Have 1 documented dose of MMR vaccine (5% will be susceptible)</td>
<td>If desired</td>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>Yes</td>
<td>Passive</td>
</tr>
<tr>
<td>History of measles disease (not documented)</td>
<td>Yes</td>
<td>Yes</td>
<td>No</td>
<td>Yes</td>
<td>Yes</td>
<td>Active</td>
</tr>
<tr>
<td>Measles IgG negative or known to be unvaccinated</td>
<td>-</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Active</td>
</tr>
<tr>
<td>Unknown or no documentation of vaccination or immune status, with presumption of immunity</td>
<td>If desired</td>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>Yes</td>
<td>Passive</td>
</tr>
<tr>
<td>Unknown or no documentation of vaccination or immune status, without presumption of immunity</td>
<td>Yes</td>
<td>Yes</td>
<td>No</td>
<td>Yes</td>
<td>Yes</td>
<td>Active</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Measles immunity assessment for high-risk contacts (immunocompromised, infant &lt;12 months, pregnant, healthcare worker or household contact)</th>
<th>IgG testing</th>
<th>MMR PEP</th>
<th>IG PEP</th>
<th>Quarantine if no PEP</th>
<th>Exclusion if no PEP</th>
<th>Symptom watch</th>
</tr>
</thead>
<tbody>
<tr>
<td>Unvaccinated infants &lt;12 months of age</td>
<td>No</td>
<td>No</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Active</td>
</tr>
<tr>
<td>Pregnant women without 2 documented MMR or serologic evidence of immunity</td>
<td>Yes</td>
<td>No</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Active</td>
</tr>
<tr>
<td>Severely immunocompromised people (see page 2)</td>
<td>No</td>
<td>No</td>
<td>Yes</td>
<td>See footnote</td>
<td>Yes</td>
<td>Active</td>
</tr>
<tr>
<td>Household or other contact with prolonged exposure without 2 documented MMR or serologic evidence of immunity</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Active</td>
</tr>
</tbody>
</table>

**Footnotes**
1. Postexposure prophylaxis (PEP) with MMR vaccine can be given <72 hours of exposure to persons without contraindications for the vaccine.
2. If measles IgG status is unknown, persons >12 months of age who may have been vaccinated or had disease and receive MMR vaccine as PEP should have blood drawn and tested for measles IgG at the time of MMR administration.
3. Contacts at high risk of severe infection (severely immunocompromised people, unvaccinated infants, and susceptible pregnant women) should receive IG (IM or IV) PEP ≤6 days of last exposure to measles. If it can be done rapidly, it is recommended that pregnant women be tested for measles IgG prior to administering IGIV if there is a possibility they may have received vaccine or had disease.
4. Quarantine for 21 days after last exposure unless the exposed person: is measles IgG positive, meets a presumption of immunity, has received IG (IM or IV) PEP ≤6 days of first exposure, or MMR <72 hours of first exposure. If symptoms consistent with measles develop, exposed person should be isolated. If there is concern about whether measles symptoms will be reported or if there will be compliance with quarantine, active monitoring with periodic calls to the exposed person to monitor for development of measles symptoms is recommended.
5. Exclude from high-risk settings (e.g., childcare facilities with infants and healthcare facilities) for 21 days after last exposure unless the exposed person is found to be measles IgG positive or to have two documented MMR doses. Some jurisdictions may choose to exclude from other settings with large numbers of unvaccinated persons.
6. If MMR vaccine is given >72 hours of first exposure or IG is given >6 days of first exposure, exclude from high-risk settings.
7. If patient has two documented MMR doses and an IgG negative result, base public health decisions on the two documented doses of MMR vaccine, i.e., presume immunity.
8. IGIM can be considered for persons in this category weighing <30 kg (66 lbs). There is no public health recommendation for IGIM administration in susceptible persons > 30 kg (66 lbs). If patient is >12 months MMR PEP is preferred if within 72 hours.
9. Immunity may be presumed in persons who have served in the U.S. Armed Forces; or were born in the U.S. in ≥1970 or later and attended a U.S. elementary school; or entered the U.S. in 1996 or later with an immigrant visa or have a green card, unless known to be unvaccinated.
10. If no documentation of 2 doses of MMR vaccine or measles IgG positivity is available.
11. If patient is IgG negative, or if patient has unknown status and testing cannot be completed by day 6 after exposure, administer IGIV.
12. CDPH should be consulted about severely immunocompromised measles contacts to assess the need for quarantine.