

MALARIA

1. **Agent:** Protozoan parasites *Plasmodium falciparum*, *P. malariae*, *P. ovale*, and *P. vivax*.
2. **Identification:**
 - a. **Symptoms:** Acute or subacute febrile disease, usually with episodes of chills and fever every 2-3 days, separated by afebrile periods. Malaria caused by *P. falciparum* may progress to jaundice, shock, renal failure, coma, and death.
 - b. **Differential Diagnosis:** Other febrile illnesses associated with international travel, e.g., brucellosis, typhoid fever, and yellow fever.
 - c. **Diagnosis:** Demonstration of parasites in thick or thin blood smears.
3. **Incubation:** Variable; 12 days for *P. falciparum*, 30 days for *P. malariae*, and 14 days for *P. ovale* and *P. vivax*. Inadequate or inappropriate prophylaxis may extend the incubation period.
4. **Reservoir:** Human.
5. **Source:** Infected mosquitoes of the genus *Anopheles*.
6. **Transmission:** Bite of infected mosquito, blood transfusion from infected persons, congenital and parenteral transmission.
7. **Communicability:**
 - a. **Mosquito infection:** When gametocytes are present in blood of patient.
 - b. **Parenteral Transmission:** When trophozoites are present in blood.
8. **Specific Treatment:**
 - a. *Plasmodium ovale*, *P. vivax*: Chloroquine for acute malaria, primaquine for prevention of relapses (sometimes called "radical cure"). "Terminal prophylaxis" refers to primaquine treatment after leaving regions endemic for these species.
 - b. *Plasmodium falciparum*, *P. malariae*: Chloroquine for non-resistant strains. Patients with resistant *P. falciparum* malaria may require alternative treatment; consult ACDC.
 - c. Infection by any species transmitted by transfusion, parenteral, or congenital route: chloroquine.

9. **Immunity:** Partial immunity for individuals with continuous exposure in endemic areas, e.g., Africa, Central America and Southeast Asia.

REPORTING PROCEDURES

1. **Reportable.** *California Code of Regulations*, Sections 2500, 2586.
2. **Report Form: MALARIA CASE SURVEILLANCE REPORT (DHS 8657, 9/02 fillable).**
3. **Epidemiologic Data:**
 - a. Residence in or travel to areas endemic for malaria 4 years prior to onset. List countries and cities, dates of stay and any prophylactic medication.
 - b. Transfusion of blood or blood products 2 years prior to onset. Include dates, places, lot numbers, and manufacturer. **Notify ACDC at once for assistance in follow-up.**
 - c. Use of parenteral drugs.
 1. Surveillance of travel contacts, and persons sharing intravenous drug paraphernalia for symptoms of malaria.
 2. Follow-up examination of asymptomatic mothers of infant cases, and asymptomatic infants born to mothers with malaria.

CONTROL OF CASE, CONTACTS & CARRIERS

Investigation not required by district staff. Advise ACDC regarding suspect cases; ACDC will supply diagnosing physician with appropriate form or investigate. Initiate investigation within 7 days of notification.

CASE: Isolation: None.

CONTACTS: No restrictions.

CARRIERS: Not applicable.

PREVENTION-EDUCATION

1. Appropriate chemoprophylaxis for travelers to areas endemic for malaria.
2. Avoid outdoor exposure during hours of peak mosquito activity, i.e., between dusk and dawn.

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3. Use mosquito repellent and protective clothing when traveling to areas with endemic malaria.
4. Exclude persons with malaria from blood donor programs for 3 years after becoming asymptomatic and after therapy stopped. Asymptomatic U.S. donors not on anti-malarial chemoprophylaxis may donate 6 months after returning from an endemic area.
5. IV drug users may acquire malaria by sharing paraphernalia.
6. Several episodes of locally acquired (autochthonous) malaria have been reported in several states since 1996. Vector mosquitoes (*An. quadrimaculatus* have a wide range in central and eastern USA, and *An. freeborni* in the Western USA.)

Remarks: Diagnostic titer is $\geq 1:64$. Elevated titers may persist after therapy without subsequent exposure; allow 1-2 months for results.

DIAGNOSTIC PROCEDURES

1. Microscopic:

Container: Hematology-differential (Slide holder).

Laboratory Form: Test Requisition and Report Form H-3021 or online request if electronically linked to the Public Health Laboratory.

Examination Requested: Malaria.

Material: Blood smears, 2 thick and 2 thin on standard slides.

Remarks: Obtain smears midway between febrile episodes, if possible.

2. **Serology:** IFA available through the CDC only if blood smears are negative. No testing of well persons. Send to State Department of Health Services Laboratory.

Container: State Special Serology.

Laboratory Form: State Special Serology (Lab 413).

Examination Requested: Malarial IFA.

Material: Clotted blood and negative blood smears.

Amount: 10 ml.

Storage: Refrigerate.