

# Adult Malaria

## Definition

Malaria is life-threatening mosquito-borne disease caused by parasitic Plasmodium species. Transmitted by the Anopheles mosquito in many tropical regions of the world, malaria causes the abrupt onset of fever, rigors, headache and prostration.

## Pathogens

***P. falciparum*** is most common and the most deadly species. It is most often acquired in sub-Saharan Africa, but is present worldwide in malaria regions.

***P. vivax*, *P. ovale*** and ***P. malariae*** cause milder disease, but ***P. vivax*** and ***P. ovale*** can lead to recurrent malaria if not treated appropriately.

***P. knowlesi*** is present only in Southeast Asia, but can be as severe as ***P. falciparum***.

## Risk Stratification and Therapy

**Uncomplicated malaria** - symptomatic malaria without evidence of severe disease or organ dysfunction.

**Severe or complicated malaria** - symptomatic malaria with hyperparasitemia (>2%) or evidence of organ damage/complications. Adjuvant therapies such as exchange transfusion should be considered after consultation with infectious diseases.

Indication For Therapy	Causative Organisms	Treatment Regimens
<b>Severe falciparum malaria</b> Criteria <ul style="list-style-type: none"> <li>History of recent possible exposure/no other recognized pathology OR</li> <li>Asexual forms of <i>P. falciparum</i> on blood smear</li> </ul> <b>PLUS</b> One or more of the following: <ul style="list-style-type: none"> <li>Impaired consciousness/coma</li> <li>Prostration with extreme weakness</li> <li>Severe normocytic anemia</li> <li>Acute Renal failure</li> <li>Pulmonary edema or ARDS</li> <li>Hypoglycemia</li> <li>Circulatory collapse, shock</li> <li>Spontaneous bleeding/DIC</li> <li>Repeated generalized convulsions</li> <li>Acidemia/acidosis</li> <li>Macroscopic hemoglobinuria</li> <li>Parasitemia &gt;2% in non-immune individuals</li> </ul>	<i>P. falciparum</i>	<b>First Line Therapy</b> <b>INFECTIOUS DISEASE CONSULT REQUIRED PRIOR TO ORDERING</b> <b>Artesunate<sup>(R)</sup> IV</b> 2.4 mg/kg per dose IV at 0, 12, 24 and 48 hours. First dose to be administered STAT. Give IV push over 1-2 min <sup>5</sup> . <b>THEN</b> Start one of the following 4 hours after the last dose of Artesunate: <b>Atovaquone<sup>†</sup> 250 mg/Proguanil 100 mg (Malarone)</b> 4 tabs PO daily x 3 days OR <b>Doxycycline<sup>††</sup> 100 mg PO BID x 7 days</b> OR <b>Clindamycin<sup>*</sup> 10 mg/kg IV load, then 5 mg/kg IV q8h x 7 days</b>  <b>Alternative Therapy</b> (First line in first trimester of pregnancy) <b>INFECTIOUS DISEASE CONSULT REQUIRED PRIOR TO ORDERING</b> <b>Quinine IV<sup>(R)</sup></b> Loading dose <sup>**</sup> : 5.8 mg/kg base (7 mg/kg quinine dihydrochloride) <sup>1</sup> Maintenance: 8.3 mg/kg base (10 mg/kg quinine dihydrochloride) <sup>2</sup>  When patient is able to swallow, switch to oral quinine tablets to complete 3-7 days of therapy <sup>***</sup> . For patients requiring >48 h of IV therapy, reduce maintenance dose by 1/3 to 1/2.  <b>PLUS</b> one of the following (either concurrently with quinine or immediately after): <b>Atovaquone<sup>†</sup> 250 mg/Proguanil 100 mg (Malarone)</b> 4 tabs PO daily x 3 days OR <b>Doxycycline<sup>††</sup> 100 mg PO BID x 7 days</b> OR <b>Clindamycin<sup>*</sup> 10 mg/kg IV load, then 5 mg/kg IV q8h x 7 days</b>

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Indication For Therapy	Causative Organisms	Treatment Regimens
<b>Uncomplicated falciparum malaria</b>	Chloroquine-sensitive falciparum	<b>Chloroquine Phosphate (Aralen 150 mg base/tab)</b> 2 tabs PO BID on Day 1 & 2, then 2 tabs PO on Day 3.
	Chloroquine-resistant falciparum	<b>Atovaquone<sup>†</sup> 250 mg/Proguanil 100 mg (Malarone)</b> 4 tabs PO daily x 3 days <b>OR</b> <b>Quinine sulfate</b> (250 mg base/300 mg salt/tab) 2 tabs PO TID x 7 days  <b>PLUS one</b> of the following (either concurrently with quinine or immediately after): <b>Doxycycline<sup>††</sup> 100 mg PO BID x 7 days</b> <b>OR</b> <b>Clindamycin* 300 mg PO base q6h x 7 days</b>
<b>Non-falciparum malaria</b> (For non-falciparum malaria acquired outside of New Guinea, chloroquine remains the drug of choice)	<i>P. ovale</i> <i>P. vivax</i> <i>P. malariae</i> <i>P. knowlesi</i>	<b>Chloroquine Phosphate (Aralen 150 mg base/tab)</b> 2 tabs PO BID on Day 1 & 2, then 2 tabs PO on Day 3

<sup>1</sup> In 100 mL of NS or D5W infused by infusion pump over 30 minutes.

<sup>2</sup> Diluted in 10 mL/kg of NS or D5W infused over 4 hours q8h

<sup>5</sup>Hemolysis can occur 1-3 weeks after treatment initiation. CBC should be followed weekly for 4 weeks after treatment initiation.

<sup>†</sup>Preferred agent unless patient received Malarone prophylaxis, is pregnant, or has a CrCl less than 30 mL/min.

<sup>††</sup>Contraindicated if age <8 years old or in pregnancy or breastfeeding.

\*Use clindamycin only if the patient is unable to take other alternatives (i.e. Malarone or doxycycline). Where IV therapy is recommended, may step-down to PO clindamycin 20 mg/kg/day divided QID once oral therapy is tolerated.

\*\*Do not give IV quinine loading dose if patient received quinine, quinidine or mefloquine within preceding 24 hours.

\*\*\*7 day duration of quinine therapy is recommended for *P. falciparum* infections acquired in Southeast Asia. 3 day therapy is recommended for all other regions.

(R) – This antimicrobial agent is **restricted**; Refer to Osler's antimicrobial restriction policies for more information