



# **NATIONAL DRUG POLICY ON MALARIA**

**2013**



**Directorate of National Vector Borne Disease Control  
Programme**

**(Directorate General of Health Services)**

**Ministry of Health and Family Welfare**

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## NATIONAL DRUG POLICY ON MALARIA (2013)

### Preamble

Malaria is one of the major public health problems of the country. Around 1.5 million laboratory confirmed cases of malaria are annually reported in India. Around 50% of the total malaria cases reported is due to *P.falciparum*. One of the reasons attributed to rise in proportion of *P.falciparum* cases is resistance to chloroquine, which was used for a long time as the first line of treatment of malaria cases. *P.falciparum* infections are known to lead to severe malaria, if timely treatment with effective drugs is not administered.

The National Drug Policy on Malaria was first formulated in 1982 and has subsequently been reviewed and revised periodically. The present National Drug Policy for Malaria (2013) has been drafted keeping in view the availability of more effective antimalarial drugs and drug resistance status in the country.

Early diagnosis and complete treatment is one of the key strategies of the National Malaria Control Programme. All fever cases clinically suspected of malaria should be investigated for confirmation of malaria by either microscopy or Rapid Diagnostic Test (RDT)<sup>1</sup>.

In high Pf predominant areas where it is not possible to get microscopy results within 24 hours, ASHAs/other community health volunteers/MPWs should be provided with rapid diagnostic kits and anti-malarials (including ACT) for early diagnosis and treatment of *P.falciparum* cases.

Effective treatment of malaria under the National Drug Policy aims at:

- Providing complete cure (clinical and parasitological) of malaria cases
- Prevention of progression of uncomplicated malaria into severe malaria and thereby reduce malaria mortality
- Prevention of relapses by administration of radical treatment
- Interruption of transmission of malaria by use of gametocytocidal drugs
- Preventing development of drug resistance by rational treatment of malaria cases.

### Treatment of uncomplicated malaria

1. It is stressed that all fever cases should be suspected of malaria after ruling out other common causes and should be investigated for confirmation of malaria by Microscopy or Rapid Diagnostic Kit (RDK) so as to ensure treatment with full therapeutic dose with appropriate drug to all confirmed cases.

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<sup>1</sup> Till 2012, Pf RDTs have been supplied under NVBDCP. From 2013, Bivalent RDT have been introduced

2. The malaria case management is very important for preventive serious cases and death due to malaria. So, the private healthcare providers should also follow the common National Guidelines for treatment of malaria as per the Drug Policy 2013
3. *P. vivax* cases should be treated with chloroquine for three days and Primaquine for 14 days. Primaquine is used to prevent relapse but is contraindicated in pregnant women, infants and individuals with G6PD deficiency.

Note: Patients should be instructed to report back in case of haematuria or high colored urine / cyanosis or blue coloration of lips and Primaquine should be stopped in such cases. Care should be taken in patients with anaemia.

4. *P. falciparum* cases should be treated with ACT (Artesunate 3 days + Sulphadoxine-Pyrimethamine 1 day). This is to be accompanied by single dose primaquine preferably on day 2.
5. However, considering the reports of resistance to partner drug SP In North Eastern States, the Technical Advisory Committee has recommended to use the Co-formulated tablet of ARTEMETHER( 20 mg) - LUMEFANTRINE (120 mg (ACT-AL) as per the age-specific dose schedule for the treatment of Pf cases in North Eastern States **(Not recommended during the first trimester of pregnancy and for children weighing < 5 kg)**
6. Production and sale of Artemisinin monotherapy has been banned in India
7. Pregnant women with uncomplicated *P. falciparum* should be treated as follows:
  - 1<sup>st</sup> Trimester: Quinine
  - 2<sup>nd</sup> & 3<sup>rd</sup> Trimester: ACT

**Note: Primaquine is contra indicated in pregnant woman**

8. In cases where parasitological diagnosis is not possible due to non-availability of either timely microscopy or RDT, suspected malaria cases will be treated with full course of chloroquine, till the results of microscopy are received. Once the parasitological diagnosis is available, appropriate treatment as per the species, is to be administered.
9. Presumptive treatment with chloroquine is no more recommended.
10. Resistance should be suspected if in spite of full treatment with no history of vomiting, diarrhoea, patient does not respond within 72 hours, clinically and parasitologically. Such cases not responding to ACT, should be treated with oral quinine with Tetracycline / Doxycycline. These instances should be reported to concerned District Malaria /State Malaria Officer/ROHFW for initiation of therapeutic efficacy studies.

## Treatment of *Vivax* Malaria

Diagnosis of vivax malaria may be made by the use of RDT (Bivalent) or microscopic examination of the blood smear. On confirmation following treatment is to be given:

### Drug schedule for treatment of *P vivax* malaria:

**1. Chloroquine:** 25 mg/kg body weight divided over three days i.e.  
10 mg/kg on day 1,  
10 mg/kg on day 2 and  
5 mg/kg on day 3.

**2. Primaquine\*:** 0.25 mg/kg body weight daily for 14 days.

**Primaquine is contraindicated in infants, pregnant women and individuals with G<sub>6</sub>PD deficiency.**

14 day regimen of Primaquine should be given under supervision.

### Dosage Chart for Treatment of *Vivax* Malaria

Age	Day 1		Day 2		Day 3		Day 4 to 14
	CQ (150 mg base)	PQ (2.5 mg)	CQ (150 mg base)	PQ (2.5 mg)	CQ (150 mg base)	PQ (2.5 mg)	PQ (2.5 mg)
Less than 1 yr	½	0	½	0	¼	0	0
1-4 years	1	1	1	1	½	1	1
5-8 years	2	2	2	2	1	2	2
9-14 years	3	4	3	4	1½	4	4
15 yrs or more*	4	6	4	6	2	6	6
Pregnancy	4	0	4	0	2	0	0

**Note: CQ 250mg tablet is having 150 mg base**

## **Treatment of *Falciparum* Malaria**

Diagnosis of falciparum malaria may be made by the use of RDT (Monovalent or Bivalent) or microscopic examination of the blood smear. It is imperative to start the treatment for falciparum malaria immediately on diagnosis. The treatment for falciparum malaria is as follows:

### **In Other States (other than North-Eastern States):**

#### **1. Artemisinin based Combination Therapy (ACT-SP)**

Artesunate (AS), available as 50 mg tablets are given for three days, and Sulfadoxine-Pyrimethamine (S-P) tablets, containing 500 mg Sulfadoxine and 25 mg pyrimethamine are given for one day, as shown in the dosage chart below.

All tablets for a day should be taken together, swallowed with water.

In addition, Primaquine (PQ Large) tablets should be given on the second day.

### **Dose schedule for Treatment of uncomplicated *P.falciparum* cases:**

#### **1. Artemisinin based Combination Therapy (ACT-SP)**

Artesunate 4 mg/kg body weight daily for 3 days Plus  
Sulfadoxine (25 mg/kg body weight) – Pyrimethamine (1.25 mg/kg body weight)  
on first day.

\* ACT is not to be given in 1st trimester of pregnancy.

#### **2. Primaquine: 0.75 mg/kg body weight on day 2.**

With the introduction of different coloured Blister Packs for different age groups, treatment by the field level staff has been made easy. The colour code for different age groups for Packing of Tablet ACT+SP has been given as follows:

**Dosage Chart for Treatment of *falciparum* Malaria with ACT-SP**

Age Group (Years)	1 <sup>st</sup> day		2 <sup>nd</sup> day		3 <sup>rd</sup> day
	AS	SP	AS	PQ	AS
0-1* Pink Blister	1 (25 mg)	1 (250 +12.5 mg)	1 (25 mg)	Nil	1 (25 mg)
1-4 Yellow Blister	1 (50 mg)	1 (500+25 mg each)	1 (50 mg)	1 (7.5 mg base)	1 (50 mg)
5-8 Green Blister	1 (100 mg)	1 (750+37.5 mg each)	1 (100 mg)	2 (7.5 mg base each)	1 (100 mg)
9-14 Red Blister	1 (150 mg)	2 (500+25 mg each)	1 (150mg)	4 (7.5 mg base each)	1 (150 mg)
15 & Above White Blister	1 (200 mg)	2 (750+37.5 mg each)	1 (200 mg)	6 (7.5 mg base each)	1 (200 mg)

\* SP is not to be prescribed for children <5 months of age and should be treated with alternate ACT

### In North-Eastern States (NE States):

1. ACT-AL Co-formulated tablet of ARTEMETHER( 20 mg) - LUMEFANTRINE (120 mg)

(Not recommended during the first trimester of pregnancy and for children weighing < 5 kg)

#### Recommended regimen by weight and age group

The packing size for different age groups based on Kg bodyweight.

Co-formulated tablet ACT-AL	5–14 kg ( > 5 months to < 3 years)	15–24 kg (≥ 3 to 8 years)	25–34 kg (≥ 9 to 14 years)	> 34 kg ( > 14 years)
Total Dose of ACT-AL	20 mg/ 120 mg twice daily for 3 days	40 mg /240 mg twice daily for 3 days	60 mg /360 mg twice daily for 3 days	80 mg /480 mg twice daily for 3 days
	<b>Pack size</b>			
No. of tablets in the Packing	6	12	18	24
Give	1 Tablet twice daily for 3 days	2 Tablets twice daily for 3 days	3 Tablets Twice daily for 3 days	4 Tablets Twice daily for 3 days
Colour of the pack	Yellow	Green	Red	White

\* ACT-AL is not to be prescribed for children weighting less than 5 kg.

**2. Primaquine:** 0.75 mg/kg body weight on day 2

### **Treatment of uncomplicated *P.falciparum* cases in pregnancy:**

**1st Trimester** : **Quinine** salt 10mg/kg 3 times daily for 7 days.

Quinine may induce hypoglycemia; pregnant women should not start taking quinine on an empty stomach and should eat regularly, while on quinine treatment.

**2<sup>nd</sup> and 3<sup>rd</sup> trimester:** **Area-specific ACT** as per dosage schedule given above.

i.e. **ACT-AL in North Eastern States**

**ACT-SP in Other States**

Primaquine (PQ) prevents transmission of falciparum malaria to others by its ability to kill gametocytes. PQ tablets should be taken after a meal; not on an empty stomach. Children less than the age of one year and pregnant women should not be given Primaquine. As pregnant women having falciparum malaria require different medicines, they should be directed to go to the nearest PHC or hospital immediately, without delay.



## Treatment of mixed infections (*P.vivax* + *P.falciparum*) cases:

All mixed infections should be treated with full course of ACT and Primaquine 0.25 mg per kg body weight daily for 14 days.

**In North-Eastern States:** Treat with: Age-specific ACT-AL for 3 days + Primaquine 0.25 mg per kg body weight daily for 14 days.

**In Other States:** SP-ACT 3 days + Primaquine 0.25 mg per kg body wt. daily for 14 days.

### Dosage Chart for Treatment of mixed (*vivax* and *falciparum*) Malaria with ACT-SP

Age	Day 1			Day 2		Day 3		Days 4-14
	AS tablet (50 mg)	SP tablet	PQ (2.5 mg)	AS tablet (50 mg)	PQ (2.5 mg)	AS tablet (50 mg)	PQ (2.5 mg)	PQ (2.5 mg)
Less than 1 yr	½	½	0	½	0	½	0	0
1-4 years	1	1	1	1	1	1	1	1
5-8 years	2	1 ½	2	2	2	2	2	2
9-14 years	3	2	4	3	4	3	4	4
15 yrs or more	4	3	6	4	6	4	6	6

## Treatment of *P. ovale* and *P. malariae*:

In India these species are very rarely found in few places. *P. ovale* should be treated as *P. vivax* and *P. malariae* should be treated as *P. falciparum*.

## Treatment of mixed infections:

All cases of mixed infection are to be treated as Pf as per the drug policy applicable in the area plus primaquine for 14 days

## Treatment of severe malaria cases

Severe malaria is an emergency and treatment should be given as per severity and associated complications which can be best decided by the treating physicians. Before admitting or referring patients, the attending doctor or health worker, whoever is able to do it, should do RDT and take blood smear; give a parenteral dose of artemisinin derivative or quinine in suspected cerebral malaria cases and send case sheet, details of treatment history and blood slide with patient. Parenteral artemisinin derivatives or quinine should be used irrespective of chloroquine resistance status of the area with one of the following options:

### Chemotherapy of severe and complicated malaria

Initial parenteral treatment for at least 48 hours: <b>CHOOSE ONE of following four options</b>	Follow-up treatment, when patient can take oral medication following parenteral treatment
<p><b>Quinine:</b> 20mg quinine salt/kg body weight on admission (IV infusion or divided IM injection) followed by maintenance dose of 10 mg/kg 8 hourly; infusion rate should not exceed 5 mg/kg per hour. Loading dose of 20mg/kg should not be given, if the patient has already received quinine.</p>	<p><b>Quinine</b> 10 mg/kg three times a day with: doxycycline 100 mg once a day <b>or</b> clindamycin in pregnant women and children under 8 years of age, - to complete 7 days of treatment.</p>
<p><b>Artesunate:</b> 2.4 mg/kg i.v. or i.m. given on admission (time=0), then at 12 h and 24 h, then once a day.</p> <p><b>or</b></p> <p><b>Artemether:</b> 3.2 mg/kg bw i.m. given on admission then 1.6 mg/kg per day.</p> <p><b>or</b></p> <p><b>Arteether:</b> 150 mg daily i.m for 3 days in adults only (not recommended for children).</p>	<p>Full oral course of Area-specific ACT:</p> <p><b>In NorthEastern states:</b> Age-specific ACT-AL for 3 days + PQ Single dose on second day</p> <p><b>In other states:</b> Treat with: ACT-SP for 3 days + PQ Single dose on second day</p>

**Note:** The parenteral treatment in severe malaria cases should be given for minimum of 24 hours once started (irrespective of the patient's ability to tolerate oral medication earlier than 24 hours).

After parenteral artemisinin therapy, patients will receive a full course of Area-specific oral ACT for 3 days. Those patients who received parenteral Quinine therapy should receive oral Quinine 10 mg/kg body weight three times a day for 7 days (including the days when parenteral Quinine was administered) plus Doxycycline 3 mg/kg body weight once a day or Clindamycin 10 mg/kg body weight 12-hourly for 7 days (Doxycycline is contraindicated in pregnant women and children under 8 years of age) or area-specific ACT as described.

**Note:**

- Pregnant women with severe malaria in any trimester can be treated with artemisinin derivatives, which, in contrast to quinine, do not risk aggravating hypoglycaemia.
- The parenteral treatment should be given for minimum of 48 hours
- Once the patient can take oral therapy, give:
- Quinine 10 mg/kg three times a day with doxycycline 100 mg once a day or clindamycin in pregnant women and children under 8 years of age, to complete 7 days of treatment, in patients started on parenteral quinine.
- Full course of ACT to patients started on artemisinin derivatives.
- Use of mefloquine should be avoided in cerebral malaria due to neuropsychiatric complications associated with it.

### **Some don'ts in severe malaria case management**

Do not use corticosteroids, give intravenous mannitol, use heparin as anticoagulant, administer adrenaline or overhydrate.

## Chemoprophylaxis

Chemoprophylaxis should be administered only in selective groups in high *P.falciparum* endemic areas. Use of personal protection measures including Insecticide Treated bed Nets (ITN) / Long Lasting Insecticidal Nets (LLIN) should be encouraged for pregnant women and other vulnerable population including travellers for longer stay. However, for longer stay of Military and Para-military forces in high *Pf* endemic areas, the practice of chemoprophylaxis should be followed wherever appropriate e.g. troops on night patrol duty and decisions of their Medical Administrative Authority should be followed.

### Short term chemoprophylaxis (up to 6 weeks)

**Doxycycline:** 100 mg once daily for adults and 1.5 mg/kg once daily for children (contraindicated in children below 8 years). The drug should be started 2 days before travel and continued for 4 weeks after leaving the malarious area.

**Note:** It is not recommended for pregnant women and children less than 8 years.

### Chemoprophylaxis for longer stay (more than 6 weeks)

**Mefloquine:** 250 mg weekly for adults and should be administered two weeks before, during and four weeks after exposure.

**Note:** Mefloquine is contraindicated in individuals with history of convulsions, neuropsychiatric problems and cardiac conditions. Therefore, necessary precautions should be taken and all should undergo screening before prescription of the drug.

**Note:** The treatment matrix for different situations like unavailability of Microscopy in 24 hours, Microscopy available, where Bi-valent RDT is available is given in Annexure-1.

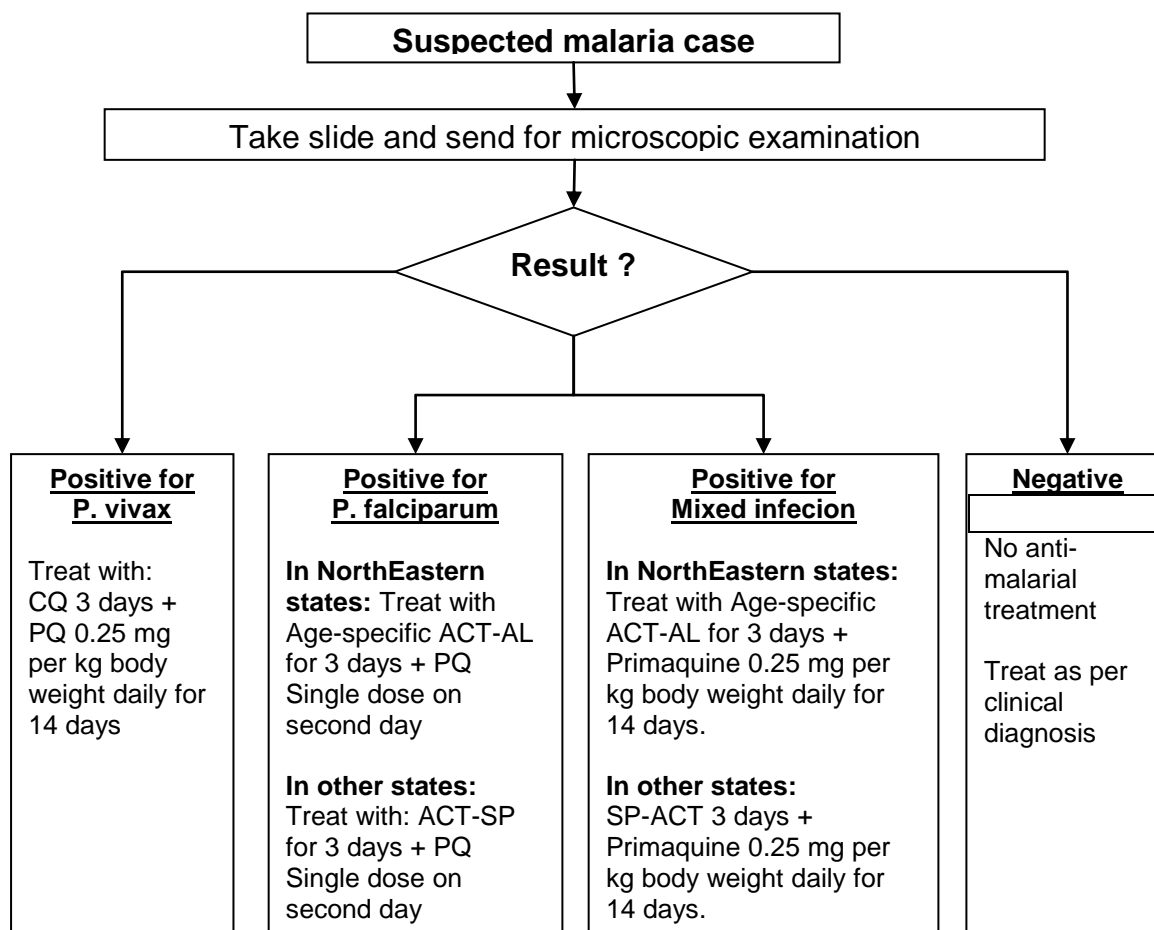
### References:

1. National Drug Policy for Malaria 2010, NVBDCP, MoH&FW, Govt. of India, 2010.
2. Operation Manual for Malaria Control for District level Officers. NVBDCP, 2008.
3. Record Note of the Meeting of Technical Advisory Committee, NVBDCP, 2013.
4. Website of National Vector Borne Disease Control Programme  
<http://www.nvbdc.gov.in/malaria-new.html>
5. World Health Organization (2006). WHO Guidelines for the Treatment of Malaria. Geneva World Health Organization

<http://www.who.int/malaria/docs/TreatmentGuidelines2006.pdf>

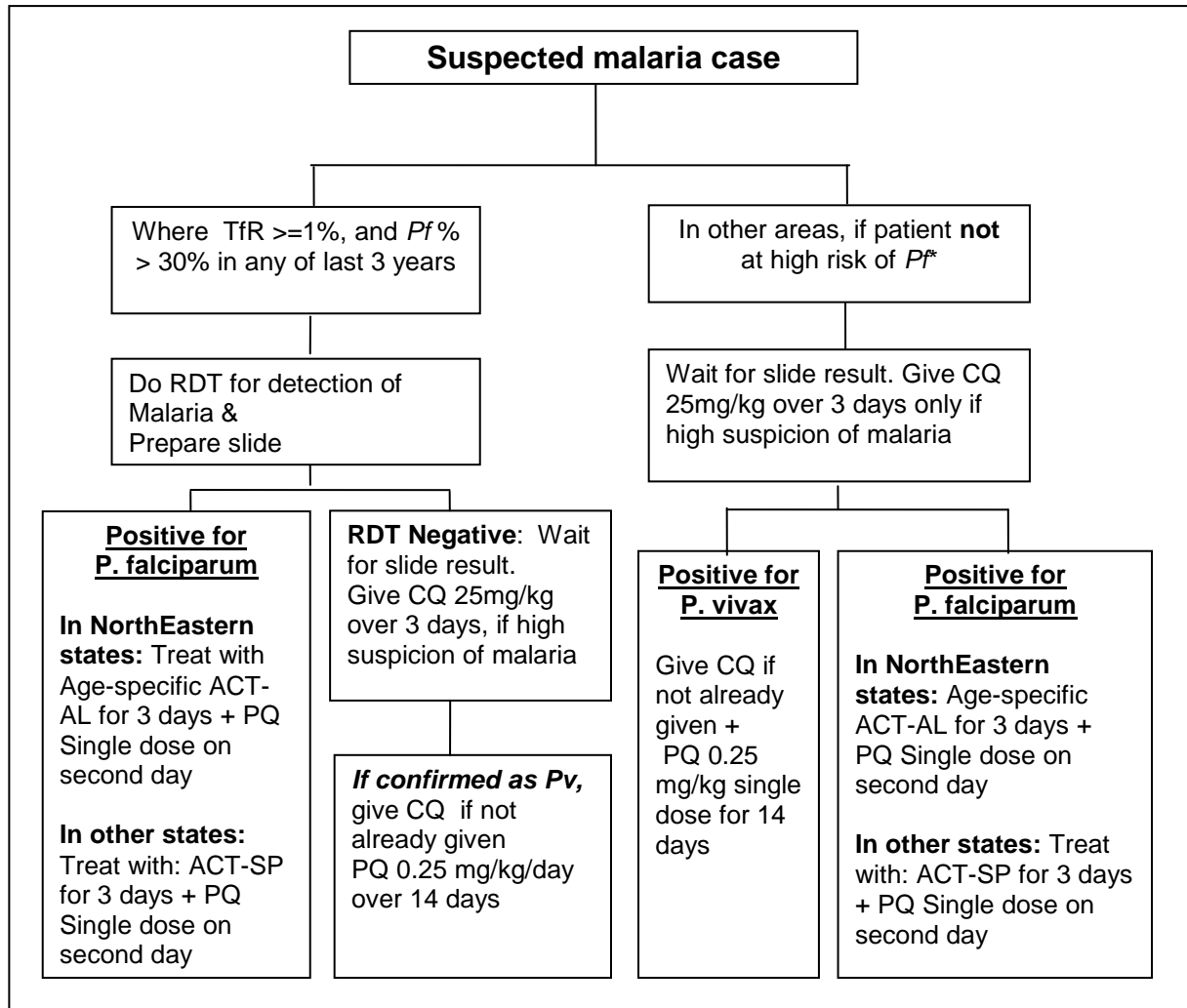
**DRUG SCHEDULE FOR TREATMENT OF MALARIA UNDER NVBDCP****Diagnosis and Treatment for Malaria****Diagnosis & Treatment**

All fever cases diagnosed as malaria by either RDT or microscopy should be promptly given effective treatment. The medicine chosen will depend upon whether the patient has vivax malaria or falciparum malaria as diagnosed by the blood test. The flow charts in different settings for diagnosis and drug selection for the treatment of malaria are as under:

**Where microscopy result is available within 24 hours**

**ACT-AL** - Artemisinin-based Combination Therapy- Artemether - Lumefantrine  
**ACT-SP**- Artemisinin-based Combination Therapy (Artesunate+Sulfadoxine-Pyrimethamine)  
**CQ** - Chloroquine  
**PQ** - Primaquine

**Where microscopy result is not available within 24 hours and Monovalent RDT is used**



TfR= Test falciparum rate

**Note:** if a patient has severe symptoms at any stage, then immediately refer to a nearest PHC or other health facility with indoor patient management or a registered medical doctor.

**Note:** PQ is contra-indicated in pregnancy and in children under 1 year (Infant).

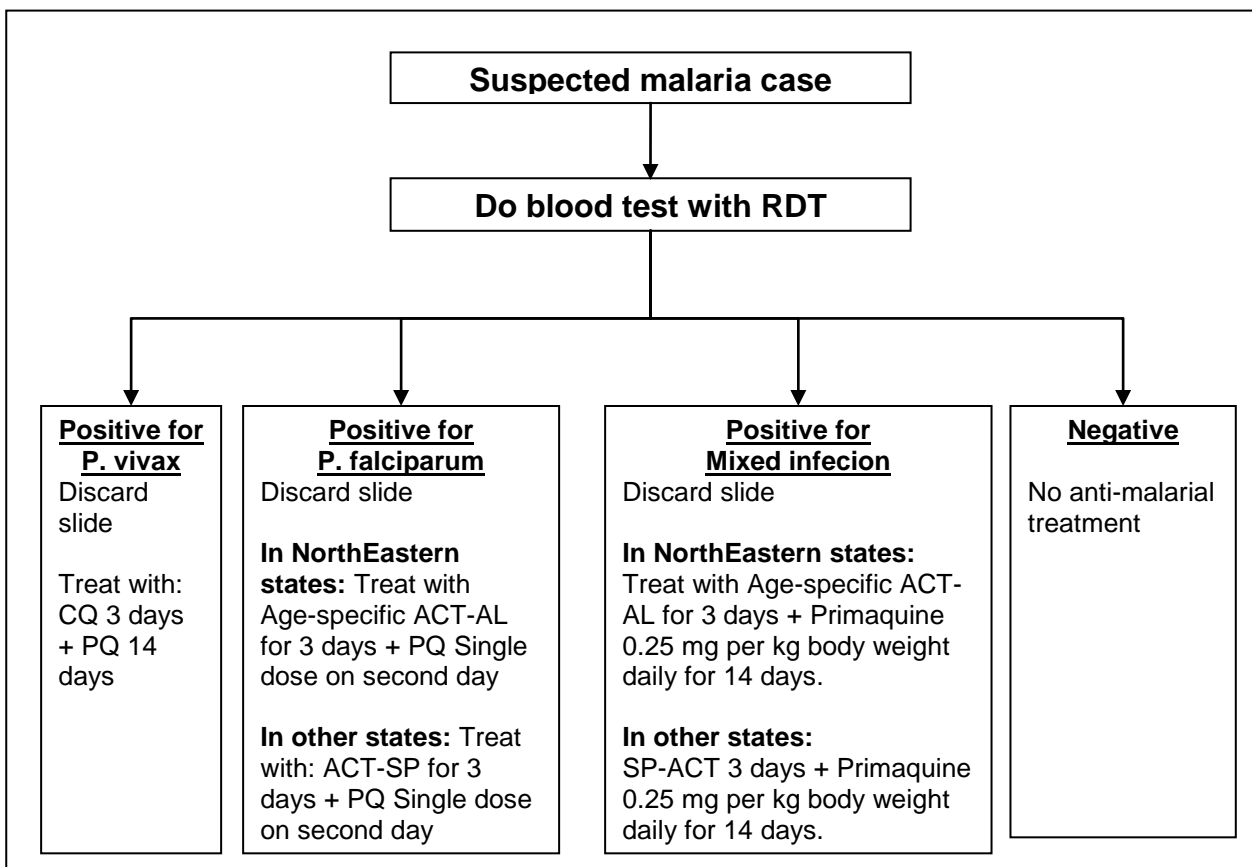
**ACT-AL** - Artemisinin-based Combination Therapy- Artemether - Lumefantrine

**ACT-SP**- Artemisinin-based Combination Therapy (Artesunate+Sulfadoxine-Pyrimethamine)

**CQ** - Chloroquine

**PQ** - Primaquine

**Where microscopy result is not available within 24 hours and Bivalent RDT is used**



- Note:** 1) However, if malaria is strongly suspected, prepare & send slide for microscopy  
2) If a patient has severe symptoms at any stage, then immediately refer to a nearest PHC or other health facility with indoor patient management or a registered medical doctor.  
3) PQ is contra-indicated in pregnancy and in children under 1 year (Infant).

**Note: PQ is contra-indicated in pregnancy and in children under 1 year (Infant).**

**ACT-AL** - Artemisinin-based Combination Therapy- Artemether - Lumefantrine

**ACT-SP-** Artemisinin-based Combination Therapy (Artesunate+Sulfadoxine-Pyrimethamine)

**CQ** - Chloroquine

**PQ** - Primaquine