

# Malaria study - bulaquine versus primaquine

<b>Submission date</b> 23/02/2006	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 03/03/2006	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 13/09/2017	<b>Condition category</b> Infections and Infestations	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

Prof Nilima Kshirsagar

### Contact details

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## Additional identifiers

### Protocol serial number

N/A

## Study information

### Scientific Title

Malaria study - bulaquine versus primaquine

### Acronym

BQ vs PQ

### **Study objectives**

To assess whether bulaquine has superior gametocytocidal activity versus primaquine for Plasmodium falciparum malaria

### **Ethics approval required**

Old ethics approval format

### **Ethics approval(s)**

Yes, protocol approved by the Ethics Committee of the Seth GS Medical College and KEM Hospital, Mumbai, India

### **Study design**

Randomised, parallel group

### **Primary study design**

Interventional

### **Study type(s)**

Treatment

### **Health condition(s) or problem(s) studied**

Uncomplicated P. falciparum malaria

### **Interventions**

Patients are randomised to receive either one of two gametocytocidal drugs: bulaquine 75 mg or primaquine 45 mg

### **Intervention Type**

Drug

### **Phase**

Not Applicable

### **Drug/device/biological/vaccine name(s)**

Bulaquine, primaquine

### **Primary outcome(s)**

Gametocytemia at days 8, 15, 22 and 29 of follow-up

### **Key secondary outcome(s))**

Gametocyte viability at days 8, 15, 22 and 29

### **Completion date**

31/12/2005

## **Eligibility**

### **Key inclusion criteria**

1. Aged more than 16 years
2. Uncomplicated Plasmodium falciparum malaria
3. Gametocyte count more than 55 /ul within 72 hours of diagnosis, irrespective of asexual parasitemia
4. Willing to give written, informed consent

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex**

All

**Key exclusion criteria**

1. Pregnancy
2. Lactation
3. Allergy to primaquine or bulaquine
4. Glucose-6-Phosphate Dehydrogenase (G6PD) deficiency
5. Co-infection with Plasmodium vivax

**Date of first enrolment**

01/06/2003

**Date of final enrolment**

31/12/2005

**Locations****Countries of recruitment**

India

**Study participating centre****Not provided at time of registration**

Dean, Professor and Head

Mumbai

India

400012

**Sponsor information**

## Organisation

Indian Council of Medical Research

## ROR

<https://ror.org/0492wrx28>

## Funder(s)

### Funder type

Government

### Funder Name

The project was undertaken as part of the Indian Council of Medical Research centre for Advanced Research in Clinical Pharmacology at the department of clinical pharmacology, Seth GS Medical College & KEM Hospital, Mumbai, India

## Results and Publications

### Individual participant data (IPD) sharing plan

### IPD sharing plan summary

Not provided at time of registration

### Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	results	01/02/2006		Yes	No
<a href="#">Other publications</a>		01/12/2006		Yes	No