

A research study of the role of chloroquine in treating patients with dengue

Submission date 10/11/2006	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 21/11/2006	Overall study status Completed	☐ Statistical analysis plan ☒ Results
Last Edited 05/02/2015	Condition category Infections and Infestations	☐ Individual participant data

Plain English summary of protocol

Background and study aims

This study aims to find out whether chloroquine reduces the number of dengue viruses in a patient with a dengue infection. Dengue fever is the most common mosquito-transmitted viral disease in humans. Severe forms of dengue infection can result in dengue hemorrhagic fever (DHF) and dengue shock syndrome (DSS). Currently, no vaccine or treatment exists except for supportive care. Studies have shown that the dengue virus needs a certain level of acidity (or pH) in order to enter into human cells. Chloroquine is a drug known to affect other viruses that require a pH-dependent step to enter human cells. Therefore, this study will test whether chloroquine shows potential to be a treatment for dengue-infected patients by reducing the number of viruses that infect cells.

Who can participate?

Eligible patients have uncomplicated signs of dengue fever (fever, headache, aches, rash) after a standard clinical examination, with a history of symptoms less than 5 days. They must be over 14 years of age and weigh more than 45 kg, with no previous history of hypersensitivity to chloroquine, and cannot be pregnant or be receiving therapy for other disorders.

What does the study involve?

Prior to recruitment, all patients will have a standard clinical examination, a chest x-ray and a blood test to count the number of dengue viruses. The patients will then be randomly allocated to either group A or group B. Group A patients will receive chloroquine treatment and group B patients will receive a placebo (dummy) tablet. The study will then compare dengue outcomes in patients receiving placebo versus chloroquine by measuring fever clearance time and the time until tests show no presence of dengue virus. All adverse events will be fully recorded including duration, severity, outcome and relationship to study drug. Liver function tests will be repeated at discharge in all patients.

What are the possible benefits and risks of participating?

The Oxford University Clinical Research Unit will provide the drugs and support for the trial, and all patients will receive standard dengue clinical examination, diagnosis and treatment. Chloroquine side effects include nausea, vomiting, diarrhea, abdominal cramps and headache.

Where is the study run from?

The study is run by researchers at the Oxford University Clinical Research Unit (Viet Nam) and the Hospital for Tropical Diseases (Ho Chi Minh City, Viet Nam).

When is the study starting and how long is it expected to run for?

The study began in July 2006 and ended in March 2008.

Who is funding the study?

The Wellcome Trust (UK).

Who is the main contact?

The Clinical Trials Unit at the Oxford University Clinical Research Unit - Viet Nam

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Contact information

Type(s)

Scientific

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Additional identifiers

Protocol serial number

061330

Study information

Scientific Title

A randomised, double-blind, placebo-controlled trial of chloroquine for treatment of dengue

Acronym

EF

Study objectives

The primary purpose of this protocol is to evaluate chloroquine as compared to placebo in the treatment of dengue with the hypothesis that chloroquine will decrease viral replication and therefore may confer a clinical advantage. This protocol will also attempt to define differences

in clinical manifestations, the relationship between chloroquine concentrations and viral dynamics, and the pathogenesis of dengue, which may help to improve the treatment of this disease.

Ethics approval required

Old ethics approval format

Ethics approval(s)

1. Hospital for Tropical Diseases, 20/04/2006
2. Oxford Tropical Research Ethical Committee, 23/02/2006, ref: 005-06

Study design

Double-blind randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Dengue fever

Interventions

Patients will receive placebo (starch) or 600 mg chloroquine on day one and two, then 300 mg on day three. Delivery is by oral ingestion.

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Chloroquine

Primary outcome(s)

The primary objective is to compare the antiviral efficacy of chloroquine in the treatment of dengue infections as assessed by negative Reverse Transcriptase (RT)-Polymerase Chain Reaction (PCR) detection of viral Ribonucleic Acid (RNA) in plasma and clearance of NS-1 from blood.

Key secondary outcome(s))

No secondary outcome measures

Completion date

30/03/2008

Eligibility

Key inclusion criteria

Any adult patient (either sex) with dengue who gives consent. We plan to enrol all patients with suspected dengue presenting within three days of illness onset.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

No consent

Date of first enrolment

01/07/2006

Date of final enrolment

31/01/2008

Locations**Countries of recruitment**

Viet Nam

Study participating centre

Oxford University Clinical Research Unit

Ho Chi Minh City

Viet Nam

District 5

Sponsor information**Organisation**

University of Oxford (UK)

ROR

<https://ror.org/052gg0110>

Funder(s)

Funder type

Charity

Funder Name

Wellcome Trust

Alternative Name(s)

Funding Body Type

Private sector organisation

Funding Body Subtype

International organizations

Location

United Kingdom

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?
Results article	results	10/08/2010		Yes	No