

Effect trial: Treatment of cryptococcal antigen-positive patients identified through screening using fluconazole plus flucytosine vs fluconazole alone

Submission date 10/02/2021	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 04/03/2021	Overall study status Ongoing	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 25/03/2026	Condition category Infections and Infestations	<input type="checkbox"/> Individual participant data <input checked="" type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

Over the last 10 years, many more people living with HIV in Africa have been able to start antiretroviral treatment (ART). However, about one-third of people living with HIV only seek care when they have advanced HIV disease (when their CD4 count falls below 200 cells/ μ l). The frequency of infections such as cryptococcal meningitis in this group is high. Cryptococcal meningitis, caused by a fungus, is the commonest form of meningitis in sub-Saharan Africa, resulting in about 180,000 deaths per year globally with over 75% of these occurring in sub-Saharan Africa.

Over half of the patients diagnosed with cryptococcal meningitis die within 10 weeks in sub-Saharan Africa, a death rate similar to Ebola virus disease. Fortunately, cryptococcal disease can be diagnosed earlier, before the development of life-threatening meningitis, using a simple blood test to detect a part of the fungal cell called cryptococcal antigen (CrAg). Studies have shown that 70% of people testing positive for CrAg in blood would progress to develop meningitis or die without a diagnosis if they are not given antifungal treatment. Routine screening for CrAg among people presenting to HIV centres, combined with oral antifungal (fluconazole) treatment for those who screen CrAg-positive, can therefore decrease the risk of cryptococcal meningitis and death. CrAg screening has now been recommended in the national HIV guidelines of 28 countries and screening programmes are being rolled out across sub-Saharan Africa.

However, there is now also evidence showing that CrAg-positive patients who are treated with fluconazole alone have a 2.5 to 3 times higher chance of dying compared to people with similar CD4 counts who test CrAg-negative, despite receiving this antifungal treatment. Up to 40% of people testing blood CrAg-positive may have asymptomatic (subclinical) cryptococcal meningitis diagnosed at the time of the screening test; these people need a combination of antifungal medicines to treat meningitis. Yet many patients decline the procedure to diagnose meningitis (lumbar puncture) and are thus not treated appropriately for meningitis.

Recent studies have shown that combined oral treatment of fluconazole plus flucytosine for 2 weeks is safe, has few unwanted side effects and works as well as the standard of 2 weeks of

intravenous amphotericin B plus flucytosine for people treated in hospital with symptomatic cryptococcal meningitis. This study aims to compare how well this oral combination of fluconazole plus flucytosine works compared to fluconazole alone (the current recommended treatment) in reducing 6-month mortality in asymptomatic people with advanced HIV disease and a CrAg-positive blood test in Tanzania, South Africa and Vietnam.

Who can participate?

HIV-infected adults (>18 years old) identified through routine laboratory screening at participating centres as CrAg-positive in blood and who are also cerebrospinal fluid CrAg-negative or who decline a lumbar puncture.

What does the study involve?

Eligible participants are randomly allocated to be treated with either fluconazole (1200 mg) plus flucytosine OR fluconazole (1200 mg) alone for 14 days. Fluconazole (800 mg daily) will be given to all participants for a further 8 weeks and fluconazole 200 mg/d thereafter as per national guidelines. Participants who are treated as outpatients will be seen in clinic on day 1 and 14 and contacted on days 3 and 9 by telephone for adherence counselling and at 1, 2.5 (10 weeks), 4 and 6 months to determine survival status. Mortality (death rates) will be compared between the two groups.

What are the possible risks and benefits of participating?

The ultimate goal of this project is to reduce mortality due to HIV-associated cryptococcal disease by ensuring universal access to affordable, safe, and effective treatment. If a combination therapy of fluconazole and flucytosine is shown to be safe and more effective than fluconazole alone, results from this study would lead to evidence-based changes in regional and international treatment guidelines, and provide an effective and practical treatment, which could be administered to outpatients, with the potential to prevent a significant number of HIV-related deaths.

Risks associated with the oral combination of fluconazole and flucytosine are expected to be low, with little toxicity observed for CM patients in a recent phase III randomised controlled trial (ACTA trial) treated with the same combination. However, the risk-benefit for flucytosine in asymptomatic out-patients is not defined, and flucytosine can be associated with neutropenia. Toxicity in study participants will be closely monitored with adjustments made as necessary and adverse events regularly reported to the Data Monitoring Committee.

An additional potential risk is a lack of adherence to the medications due to an increased daily pill count for the first 14 days. Participants will be counselled on the importance of adherence and will be seen face-to-face twice during induction phase treatment (on day 1 and 14) and called by telephone on days 3 and 9 to encourage adherence to medication and discuss any issues. As the intervention is given over the first 2 weeks, a pill count will be performed on day 14. If significant issues are identified for some participants, patient advocates (including CM survivors) will be asked to assist.

Where is the study run from?

1. Mycology Division, WITS Health Consortium (Based at the National Institute for Communicable Diseases) (South Africa)
2. St George's University of London (UK)

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

November 2020 to August 2026

Who is funding the study?

The study is funded through the Joint Global Health Trials (JGHT) Scheme with funding from the

UK - Department of Health and Social Care (DHSC), the Foreign, Commonwealth & Development Office (FCDO), the Medical Research Council (MRC), Wellcome and National Institute for Health Research (NIHR).

Who is the main contact?

1. Prof. Nelesh Govender
neleshg@nicd.ac.za
2. Dr Síle Molloy
smolloy@sgul.ac.uk
3. Michelle Eriksson
meriksson@witshealth.co.za

Contact information

Type(s)

Scientific

Contact name

Dr Síle Molloy

ORCID ID

<https://orcid.org/0000-0001-8847-2325>

Contact details

St George's University of London
London
United Kingdom
SW17 0RE
+44 (0)208 728 5613
smolloy@sgul.ac.uk

Type(s)

Scientific

Contact name

Prof Nelesh Govender

ORCID ID

<https://orcid.org/0000-0001-7869-9462>

Contact details

Centre for Healthcare-Associated Infections, Antimicrobial Resistance and Mycoses
National Institute for Communicable Diseases
Johannesburg
South Africa
2131
+27 (0)11 555 0353
neleshg@nicd.ac.za

Type(s)

Scientific

Contact name

Ms Michelle Eriksson

Contact details

Wits Health Consortium
31 Princess of Wales Terrace
Parktown
South Africa
2193
+27 823387948
meriksson@witshealth.co.za

Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

Protocol serial number

JRES 2020.0308

Study information

Scientific Title

Fluconazole plus flucytosine vs fluconazole alone for cryptococcal antigen-positive patients identified through screening: a phase III randomised controlled trial

Acronym

EFFECT - Efficacy of Flucytosine and Fluconazole as Early Cryptococcal Treatment

Study objectives

Pre-emptive combination treatment of fluconazole plus flucytosine for 2 weeks will be more efficacious in reducing 6-month all-cause mortality than standard treatment of fluconazole alone among CrAg-positive HIV-infected adults

Ethics approval required

Old ethics approval format

Ethics approval(s)

1. Approved 22/01/2021, St George's, University of London (SGUL) Research Ethics Committee (Joint Research and Enterprise Services, Ground Floor, Jenner Wing, St. George's, University of London, Cranmer Terrace, London, SW17 0RE, UK; +44 (0)208 266 6073; sgulrec@sgul.ac.uk), ref: 2020.0308
2. Approved 21/07/2021, National Institute for Medical Research (NIMR) REC (PO Box 9653, Dar es Salaam, Tanzania; +255 22 2121 400; ethics@nimr.or.tz), no ref provided
3. Approved 12/08/2021, University of Cape Town Faculty of Health Sciences Human REC (Room G50, Old Main Building, Groote Schuur Hospital, Observatory, Cape Town 7925, South Africa; +27 (021) 650 5184; hrec-enquiries@uct.ac.za), ref: 274/2021
4. Approved 21/05/2021, University of the Witwatersrand Human REC (Suite 189, Private Bag

x2600, Houghton, Johannesburg 2021, South Africa; +27 (0)11 275 9200; HREC-Medical. ResearchOffice@wits.ac.za), ref: 210305

5. Approved 22/07/2022, University of KwaZulu-Natal Biomedical REC (UKZN Research Office, Govan Mbeki Building, Westville Campus, University Drive, Durban 4001, South Africa; +27 (031) 260 1074; brec@ukzn.ac.za; ref: BREC/00002777/2021

6. Approved 30/06/2021, Walter Sisulu University Health Research Ethics and Biosafety Committee (Academic Health Services Complex of the Eastern Cape, Postgraduate Research and Training, Faculty of Health Sciences, Walter Sisulu University, Private Bag X1, WSU, South Africa 5117; +27 (0)47 502 2100; ggeorge@wsu.ac.za), ref: 067/2021

Study design

Open-label multicentre phase III randomized controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Asymptomatic cryptococcal antigen (CrAg)-positive individuals with advanced HIV infection

Interventions

Individual randomisation using SAS PROC PLAN via a permuted-block randomisation method stratified by site. Block sizes will vary.

1. Fluconazole 1200 mg/day plus flucytosine 25 mg/kg four times daily PO (intervention arm) for 2 weeks
2. Fluconazole alone 1200 mg/day PO (standard dose 'control arm') for 2 weeks

All participants will then receive fluconazole 800 mg/day to 10 weeks, and fluconazole 200 mg /day thereafter for a minimum of 12 months as per national guidelines. ART will be commenced on day 14 as per current international (WHO) and national guidelines.

Intervention Type

Drug

Phase

Phase III

Drug/device/biological/vaccine name(s)

Flucytosine, fluconazole

Primary outcome(s)

All-cause mortality, measured using patient medical records and interviews, at 6 months after randomisation

Key secondary outcome(s)

Current secondary outcome measures as of 24/07/2025:

1. Time to all-cause mortality within first 6 months measured using patient medical records and interviews, at 6 months.
2. All-cause mortality at 10 weeks measured using patient medical records and interviews, at 10 weeks
3. Time to all-cause mortality within first 10 weeks measured using patient medical records and interviews, at 10 weeks
4. Cryptococcal meningitis-free survival to 6-months measured using patient medical records, to 6 months
5. Incidence rate of symptomatic laboratory-confirmed cryptococcal meningitis, measured using patient medical records, measured to 6 months
6. Crypto-related mortality at 6-months measured using patient medical records, to 6 months
7. Tolerability and safety: proportions of patients developing clinical and laboratory-defined grade III/IV adverse events measured using patient medical records and interviews at 6 months
8. Efficacy outcomes by baseline CrAg titre and semi-quantitative assay score at 10 weeks and 6 months
9. Rate of disability at 6 months measured using patient medical records and interviews at 6 months
10. Health service costs per life-year saved, measured using participant and health service provider interviews, at 6 months

Previous secondary outcome measures:

1. Time to all-cause mortality, measured using patient medical records and interviews, within the first 6 months
2. All-cause mortality, measured using patient medical records and interviews, at 10 weeks
3. Cryptococcal meningitis-free survival, measured using patient medical records, to 6 months
4. Incidence rate of symptomatic laboratory-confirmed cryptococcal meningitis, measured using patient medical records, over 6 months
5. Tolerability and safety: proportions of patients developing clinical and laboratory-defined grade III/IV adverse events measured using patient medical records and interviews up to 21 days
6. Efficacy outcomes by baseline CrAg titre and semi-quantitative assay score at 10 weeks and 6 months
7. Health service costs per life-year saved, measured using participant and health service provider interviews, over 6 months

Completion date

31/08/2026

Eligibility

Key inclusion criteria

1. Consecutive patients aged >18 years
2. HIV-seropositive
3. CD4 count of <100 cells/ μ l
4. Serum/plasma Cryptococcal antigen (CrAg) test positive within the last 14 days
5. Cerebrospinal fluid (CSF) CrAg test negative or lumbar puncture not done (declined)
6. Willing to participate in the study

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Mixed

Lower age limit

18 years

Upper age limit

120 years

Sex

All

Total final enrolment

0

Key exclusion criteria

Current exclusion criteria:

Initial exclusion criteria:

1. Prior episode of cryptococcal meningitis
2. Pregnancy (confirmed by urine or serum pregnancy test) or breastfeeding
3. Previous serious reaction to flucytosine or fluconazole
4. Already taking high-dose fluconazole treatment (800-1200 mg/day) for ≥ 1 week
5. Contraindicated concomitant medications including cisapride and the class of antihistamines including terfenadine
6. HIV-seronegative
7. Clinical symptoms/ signs of symptomatic meningitis at any time since CrAg screening, i.e. a progressively severe headache OR a headache and marked nuchal rigidity OR a headache and vomiting OR seizures OR a Glasgow Coma Scale (GCS) score of < 15
8. Jaundice
9. CSF positive for cryptococcal meningitis (i.e. positive microscopy with India Ink, culture, or CrAg test) at any time between the CrAg test and screening for eligibility, while the patient remained without clinical symptoms/ signs of meningitis as described in 7 (late withdrawal criterion)

Late exclusion criteria: (after randomisation and day 1 visit)

10. DAIDS grade 4 abnormalities of platelets, neutrophil count or creatinine level on baseline bloods:

10.1. Platelets $< 25,000 \times 10^6/l$

10.2. Neutrophils $< 400 \times 10^6/l$

10.3. Creatinine $> 3.5 \times$ upper limit of normal (with standardised ULN of $114 \mu\text{mol}/l$, this includes any creatinine level $\geq 400 \mu\text{mol}/l$)

11. Microbiological evidence of CM on CSF if full CSF results not present at randomisation (e.g. screening CSF CrAg negative but culture on same sample later returns positive for CM)

Previous exclusion criteria:

1. Prior episode of Cryptococcal meningitis
2. Pregnancy (confirmed by urine or serum pregnancy test) or breastfeeding
3. Previous serious reaction to flucytosine or fluconazole
4. Already taking high-dose fluconazole treatment (800-1200 mg/day) for ≥ 1 week
5. Contraindicated concomitant medications including cisapride and the class of antihistamines including terfenadine
6. HIV-seronegative
7. Clinical symptoms/ signs of symptomatic meningitis at any time since CrAg screening, i.e. a progressively severe headache OR a headache and marked nuchal rigidity OR a headache and vomiting OR seizures OR a Glasgow Coma Scale (GCS) score of < 15
8. Jaundice
9. CSF positive for Cryptococcal meningitis (i.e. positive microscopy with India Ink, culture, or CrAg test) at any time between the CrAg test and screening for eligibility, while the patient remained without clinical symptoms/ signs of meningitis as described in 7 (late withdrawal criterion)

Date of first enrolment

08/05/2022

Date of final enrolment

31/10/2025

Locations

Countries of recruitment

South Africa

Tanzania

Viet Nam

Study participating centre

Khayelitsha Site B Clinic and Mitchell's Plain Hospital

Cnr Walter Sisulu & Steve Biko Road

Khayelitsha

Cape Town

South Africa

7784

Study participating centre

Chris Hani Baragwanath Academic Hospital

26 Chris Hani Rd

Diepkloof 319-lq

Johannesburg
South Africa
1864

Study participating centre

Helen Joseph Hospital

Rossmore
Johannesburg
South Africa
2092

Study participating centre

Klerksdorp/Tshepong Hospital

Jouberton
Klerksdorp
South Africa
2574

Study participating centre

King Edward VIII Hospital

Sydney Rd
Umbilo
Durban
South Africa
4013

Study participating centre

Harry Gwala Regional Hospital (formerly Edendale Hospital)

89 Selby Msimang Rd
Plessislaer
Pietermaritzburg
South Africa
3201

Study participating centre

Livingstone Hospital

Lindsay Rd, Industrial
Gqeberha
South Africa
6020

Study participating centre
Amana Regional Referral Hospital
Uhuru St, Ilala
Dar es Salaam
Tanzania

-

Study participating centre
Mwananyama Regional Referral Hospital
Kinondoni
Dar es Salaam
Tanzania

-

Study participating centre
Temeke Regional Referral Hospital
Temeke
Dar es Salaam
Tanzania

-

Study participating centre
Dora Nginza Hospital
Spondo St
Gqeberha
South Africa
6020

Sponsor information

Organisation
St George's, University of London

ROR
<https://ror.org/040f08y74>

Funder(s)

Funder type

Research organisation

Funder Name

Wellcome Trust

Alternative Name(s)

Wellcome, WT

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

United Kingdom

Funder Name

Medical Research Council

Alternative Name(s)

Medical Research Council (United Kingdom), UK Medical Research Council, Medical Research Committee and Advisory Council, MRC

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

Funder Name

Department of Health and Social Care

Alternative Name(s)

Department of Health & Social Care, DH

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

Funder Name

National Institute for Health Research

Alternative Name(s)

National Institute for Health Research, NIHR Research, NIHRresearch, NIHR - National Institute for Health Research, NIHR (The National Institute for Health and Care Research), NIHR

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

Funder Name

Foreign, Commonwealth and Development Office

Alternative Name(s)

Foreign, Commonwealth & Development Office, Foreign, Commonwealth & Development Office, UK Government, FCDO

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

Results and Publications

Individual participant data (IPD) sharing plan**IPD sharing plan summary**

Data sharing statement to be made available at a later date

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Protocol article		24/03/2026	25/03/2026	Yes	No
Study website	Study website	11/11/2025	11/11/2025	No	Yes