205071 - A phase IV, longitudinal, crosssectional, retrospective, ancillary epidemiology study of the EPI-MAL-005 study to evaluate the genetic diversity in the Plasmodium falciparum parasite circumsporozoite sequences before and after the implementation of the RTS,S/AS01E vaccine in malaria-positive subjects ranging from 6 months to less than 5 years of age (EPI-MALARIA-010 VS AME)

**First published:** 07/10/2021 **Last updated:** 05/09/2025





## Administrative details

#### **EU PAS number**

**EUPAS42948** 

### **Study ID**

45256

#### **DARWIN EU® study**

No

### **Study countries**

□ Ghana

\_\_\_\_ Kenya

#### **Study description**

The RTS,S/AS01E vaccine has been developed for routine immunization of infants and children living in malaria-endemic countries of Sub-Saharan Africa. The aim of this retrospective, ancillary epidemiology study is to monitor the genetic diversity in circumsporozoite sequences in the Plasmodium falciparum (P. falciparum) parasite in malaria-positive subjects aged 6 months to <5 years vaccinated or not with RTS,S/AS01E.

### **Study status**

**Finalised** 

### Research institutions and networks

## Institutions

## GlaxoSmithKline (GSK)

First published: 01/02/2024

**Last updated:** 01/02/2024



### **Amsterdam UMC**

First published: 01/02/2024

**Last updated:** 01/02/2024

Institution

**Educational Institution** 

Hospital/Clinic/Other health care facility

Kintampo Health Research Centre Kintampo, Ghana, KEMRI-Walter Reed Project Kombewa, Kenya, Broad Institute (BI), Harvard T.H. Chan School of Public Health (HSPH)

### Contact details

### **Study institution contact**

Call Center EU Clinical Trials
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Study contact

Vx.public disclosure global @gsk.com

### **Primary lead investigator**

Call Center EU Clinical Trials

**Primary lead investigator** 

# Study timelines

### Date when funding contract was signed

Planned: 07/10/2020 Actual: 07/10/2020

### Study start date

Planned: 08/10/2021 Actual: 08/10/2021

### **Date of final study report**

Planned: 13/08/2025 Actual: 13/08/2025

# Sources of funding

• Pharmaceutical company and other private sector

# More details on funding

GlaxoSmithKline

# Study protocol

gsk-205071-protocol-redact.pdf (817.71 KB)

# Regulatory

Was the study required by a regulatory body?

Yes

#### Is the study required by a Risk Management Plan (RMP)?

EU RMP category 3 (required)

# Methodological aspects

# Study type

# Study type list

### **Study topic:**

Disease /health condition

### Study type:

Non-interventional study

### Scope of the study:

Assessment of risk minimisation measure implementation or effectiveness Disease epidemiology

### Main study objective:

To monitor the genetic diversity in circumsporozoite sequences in the P. falciparum parasite population before and after vaccine implementation in children aged 6 months to <5 years.

# Study Design

### Non-interventional study design

Cross-sectional

# Study drug and medical condition

#### Medical condition to be studied

Malaria

# Population studied

#### Age groups

Infants and toddlers (28 days - 23 months)
Children (2 to < 12 years)

#### **Estimated number of subjects**

5600

# Study design details

#### **Outcomes**

Prevalence of P. falciparum haplotype infections among subjects infected or not with P. falciparum and frequency of P. falciparum haplotype infections among the individual malaria clones in subjects vaccinated or not with RTS,S/AS01E per study site. Prevalence and frequency of P.falciparumhaplo type infections by age group, gender and RTS,S/AS01Evaccinationstatusper study site, Trends in longitudinal prevalence of specific

P.falciparumhaplotypesamongsubjectsinfectedornotwith
P.falciparum,vaccinatedornotwith RTS,S/AS01E,Trends
inlongitudinalfrequencyofspecific P.falciparum haplotypes among the
individualmalariaclonesinsubjectsvaccinatedornotwithRTS,S/AS01E.

#### Data analysis plan

- The haplotype prevalence will be estimated by site, as the number of subjects infected with a specific P. falciparum haplotype, divided by the total number of

subjects. Thus, the denominator will be all the subjects aged 6 months to <5 years included in the EPI-MAL-010 study for each of the 2 sites considered: malaria positive and negative subjects based on malaria blood reading and/or NAAT.

- The haplotype frequency will be estimated by site, as the number of occurrences of a specific P. falciparum haplotype, divided by the total number of clones. Thus, in case of multiple infections with P. falciparum malaria, the same subject will contribute multiple times in the denominator. The frequency will be estimated using data only from subjects aged 6 months to <5 years, measured malaria positive by microscopy and/or NAAT, included in the EPI-MAL-010 study for each of the 2 sites considered.

## Data management

### **ENCePP Seal**

The use of the ENCePP Seal has been discontinued since February 2025.

The ENCePP Seal fields are retained in the display mode for transparency but are no longer maintained.

## Data sources

#### Data source(s)

Other data source

#### Data source(s), other

Retrospective, ancillary study, re-using samples of the EPI-MAL-005 study

### Data sources (types), other

Retrospective, ancillary study, re-using samples of the EPI-MAL-005 study

# Use of a Common Data Model (CDM)

### **CDM** mapping

No

# Data quality specifications

#### **Check conformance**

Unknown

### **Check completeness**

Unknown

### **Check stability**

Unknown

### **Check logical consistency**

Unknown

## Data characterisation

#### **Data characterisation conducted**

No