Diagnostic Performance and Comparison of Ultrasensitive and Conventional Rapid Diagnostic Test, Thick Blood Smear and Quantitative PCR for Detection of Low-Density *Plasmodium Falciparum* Infections During a Controlled Human Malaria Infection Study in Equatorial Guinea

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Abstract

Background

Progress towards malaria elimination has stagnated, partly because infections persisting at low parasite densities comprise a large reservoir contributing to ongoing malaria transmission and are difficult to detect. We compared the performance of an ultrasensitive rapid diagnostic test (uRDT) designed to detect low density infections to a conventional RDT (cRDT), expert microscopy using Giemsa-stained thick blood smears (TBS), and quantitative polymerase chain reaction (qPCR) during a controlled human malaria infection (CHMI) study conducted in malaria exposed adults (NCT03590340).

Methods

Blood samples were collected from healthy Equatoguineans aged 18-35 years beginning on day 8 after CHMI with 3.2x10^3 cryopreserved, infectious Plasmodium falciparum (Pf) sporozoites (PfSPZ Challenge, strain NF54) administered by direct venous inoculation. qPCR (18s ribosomal DNA), uRDT (Alere™ Malaria Ag Pf.), cRDT (Carestart Malaria Pf/PAN (PfHRP2/pLDH)), and TBS were performed daily until the volunteer became TBS positive and treatment was administered. qPCR was the reference for the presence of Pf parasites.

Results

279 samples were collected from 24 participants; 123 were positive by qPCR. TBS detected 24/123 (19.5% sensitivity [95% CI 13.1% – 27.8%]), uRDT 21/123 (17.1% sensitivity [95% CI 11.1% – 25.1%], cRDT 10/123 (8.1% sensitivity [95% CI 4.2% – 14.8%]); all were 100% specific. qPCR was the most sensitive test (p<0.001); TBS and uRDT were more sensitive than cRDT (TBS vs. cRDT p=0.015; uRDT vs. cRDT p=0.053), detecting parasitemias as low as 3.7 parasites/mL (p/mL) (TBS and uRDT) compared to 5.6 p/mL (cRDT) based on TBS density measurements. TBS, uRDT and cRDT did not detect any of the 70/123 samples positive by qPCR below 5.86 p/mL, the qPCR density corresponding to 3.7 p/mL by TBS. The median prepatent periods in days (ranges) were 14.5 (10-20), 18.0 (15-28), 18.0 (15-20) and 18.0 (16-24) for qPCR, TBS, uRDT and cRDT, respectively; qPCR detected parasitemia significantly earlier (3.5 days) than the other tests.

Conclusions

TBS and uRDT had similar sensitivities, both were more sensitive than cRDT, and neither matched qPCR for detecting low density parasitemia. uRDT could be considered an alternative to TBS in selected applications such as CHMI or field diagnosis where qualitative, dichotomous results for malaria infection might be sufficient.

Background
Over the past decade, malaria treatment and vector control interventions have significantly decreased malaria burden worldwide. The global incidence rate of malaria decreased between 2010 and 2018 from 71 to 57 cases per 1000 people at risk [1]. However, the World Malaria Report released at the end of 2020 documented a stagnation in reduction of annual malaria cases and deaths since 2015, with 229 million cases and 409,000 deaths in 2019 [1]. A significant challenge faced by malaria control and elimination projects is addressing transmission potential from low parasite density carriers with mild or no symptoms. Low parasite density carriers are estimated to account for 20-50% of human-to-mosquito transmission [2]. It would be beneficial to have a rapid test able to identify these infections.

Current widely deployed diagnostic tools such as conventional rapid diagnostic tests (cRDTs) are affordable and have user-friendly formats and function. In 2018, 259 million cRDTs were distributed mainly in Sub-Saharan Africa and utilized to examine suspected malaria cases [1]. cRDTs test for the presence of histidine-rich protein 2 (PfHRP2), an antigen specific to *Plasmodium falciparum* (Pf), and many iterations of the test also include a pan-malaria antigen (PAN) common to all 4 major malaria species such as lactate dehydrogenase (LDH). While cRDTs are affordable, provide quick and readable results, and require little training to operate, they cannot detect low-density infections, which can occur frequently in low transmission areas [2–4]. As a result, efforts are underway to develop advanced malaria rapid diagnostic tests that are more sensitive and effective at identifying low-density Pf infections [5, 6].

Laboratory-based techniques such as thick blood smear (TBS) and polymerase chain reaction (PCR) are considered to have greater sensitivity than cRDTs [7]. Historically, TBS has been the gold standard for malaria diagnosis. When performed by expert microscopists reading 0.5 µL of blood, detection may range typically between 10 and 50 parasites per microliter (p/µL), and under carefully controlled CHMI and TBS preparation conditions, expert microscopists can quantify parasite densities at the theoretical lower limit of detection for this blood volume, 2 p/µL. In contrast, cRDTs have reported detection limits of 100-200 p/µL of blood in field studies [8–10]. TBS has the additional advantages of diagnosing infections even in the presence of parasites carrying PfHRP2 deletions, a challenge that RDT manufacturers are currently facing [11, 12], and allowing detection of all species of malaria parasite. In recent years, the further development of the quantitative polymerase chain reaction (qPCR) method has enabled the detection of low parasite densities that frequently go undetected by cRDTs and TBS [13, 14]. Utilizing qPCR, reservoirs of low parasite density cases can be identified and treated, which is an essential component for elimination in low transmission areas [7, 15–17]. The drawback of both TBS and qPCR is the need for specialized laboratory equipment, materials and well-trained staff, which are often in short supply in low income countries, making them unfeasible for use under field conditions on a large scale [18, 19]. TBS can also lead to false negative results when performed by a non-competent microscopist, the parasitemia is low or there are multiple-species co-infections [20, 21]. An ideal diagnostic tool would combine the sensitivity of qPCR with the affordability and simplicity of the cRDT.

Ultrasensitive RDTs (uRDTs) have recently been developed and offered as a reliable diagnostic tool that can be used in clinical studies and in field operations [22–24]. Similar to cRDTs in form and function, uRDTs detect PfHRP2 in Pf, but at greater sensitivity, identifying densities as low as 0.1–1.0 p/µL in
culture-derived samples [6, 22]. A higher sensitivity RDT capable of detecting low parasite density infections could target these infections for treatment. It could also provide a rapid and accurate confirmation of parasite clearance following drug administration [25] or of protection against CHMI or against natural Pf exposure following vaccination with a malaria vaccine [26–28]. To date, limited literature exists systematically examining the performance of uRDTs in direct comparison to cRDTs, TBS and qPCR in samples collected from individuals with low parasite density [29, 30]. Existing investigations have been conducted predominantly in the field where it is difficult to monitor factors such as time to infection and the waxing and waning of parasite densities [23, 30–34].

A new approach to conducting controlled human malaria infection (CHMI) has become available over the past five years, based on administering aseptic, purified, cryopreserved, infectious Pf sporozoites (SPZ) (PfSPZ Challenge) [35]. In this model, PfSPZ are administered by syringe, replacing mosquito bite administration, and the study subjects are then monitored in the standard way [36]. The advantages are that CHMI can be performed by institutions without an insectary or without the need to import infected mosquitoes, the dose of PfSPZ can be standardized, and CHMI can be administered at any time without coordinated mosquito infections. CHMI using PfSPZ Challenge has recently been utilized to evaluate efficacy of antimalarial drugs and vaccines in malaria-naive and malaria pre-exposed populations [37–43].

CHMI using injectable PfSPZ provides an opportunity to assess malaria diagnostics under carefully controlled conditions in malaria exposed populations. The exact exposure time is known, the induced infections gradually increase in density, and the time of first detection and associated prepatent period can be precisely determined. The aim of our study was to systematically evaluate and compare uRDT performance against three other commonly used malaria diagnostic tools using whole blood samples collected daily from malaria pre-exposed individuals undergoing CHMI.

**Methods**

**Study site**

The Bioko Island Malaria Elimination Program (BIMEP) focuses on developing malaria vaccines and other interventions to decrease malaria-attributable morbidity and mortality on Bioko Island, Equatorial Guinea [44]. BIMEP performs various activities on Bioko such as in-depth epidemiological studies and clinical trials of the malaria vaccine candidates Sanaria® PfSPZ Vaccine and PfSPZ-CVac [42, 45, 46] to collect data on safety and efficacy to support vaccine licensure. In 2018, the BIMEP conducted a regimen optimization trial of PfSPZ Vaccine (ClinicalTrials.gov ID: NCT03590340) at the Baney Research Facility that involved 104 healthy Equatoguinean adults male and female, aged 18-35 years (Jongo SA, AJTMH, submitted)[47]. Study volunteers were recruited from Baney district and the city of Malabo, and were enrolled after providing informed consent. Homologous (the strain of Pf in PfSPZ Challenge, PfNF54, was the same as in the vaccine) CHMI with $3.2 \times 10^3$ PfSPZ was administered to 95 eligible individuals, six to
seven weeks after last vaccination and the observation period was conducted in a hotel at the La Paz Hospital beginning eight days after PfSPZ Challenge injection.

**Malaria Rapid Diagnostic Tests**

Malaria rapid diagnostic tests were performed with frozen venous whole blood samples anticoagulated with EDTA and stored at -80°C. Blood samples were allowed to thaw slowly and equilibrate to ambient temperature for at least 30 min before performing the RDT. The commercially available Carestart malaria Pf/PAN (PfHRP2/pLDH) Ag Combo, ACCESSBIO, USA (Lot# MR18F63, expiration: 30th Nov, 2020) and the Alere™ Malaria Ag Pf, Standard Diagnostic Inc., Republic of Korea (Lot# 05LDE001A, expiration: 20th Feb, 2020) were used throughout the study. Both tests have similar manufacturer instruction workflows, requiring 5 µL blood to be applied to the RDT, followed by addition of the assay buffer and incubation for 20 min. After incubation, diagnosis was determined by the appearance of lines in the test result window. A single control line was considered malaria negative, a line for both control and test were considered malaria positive, and no lines, neither control nor test, was considered an invalid test. Two to three readers were involved in determining each RDT result, the third added when there were discrepancies between the first two readers. The samples were analysed in batches of 10 by a first reader and the developed RDTs were then given to a second reader to confirm the reading of the first reader. Any discrepancy between first and second readers was resolved using a third reader within a period of 20 ± 1 min. Results were recorded in the database as binary figures with zero representing negatives and one representing positives. Five fresh known malaria positive and five fresh known negative samples were used for quality control for the performance of the assay and for training the technicians prior to conducting the study.

**Thick blood smear for quantification of Plasmodium falciparum.**

Two milliliters of fresh whole venous blood from study participants were used for the preparation of TBS. The TBS was prepared by evenly spreading 10 µL of fresh whole venous blood into a 1 cm x 2 cm rectangle. The smears were air dried, stained for 45 min using 4% Giemsa stain and rinsed with buffered water, pH 7.2. The slides were dried and read using a light microscope with a high-power field (immersion oil, 100x objective) of 0.18 mm diameter. Six x 1 cm passes equivalent to 0.54 µL of blood or 24 x 1 cm passes equivalent to 2.14 µL of blood for symptomatic volunteers were read before a TBS was declared positive or negative. The slides were read by two independent expert microscopists and any discrepancies were resolved by a third microscopist. For Giemsa stain quality control, known positive and negative thin blood smears were included at the beginning of the day and analysed for both parasites and cell staining color and quality according to a standard operating procedure. Only Giemsa stain that passed the quality control procedures was allowed to be used for the slide staining on that day. Microscopes were maintained on a daily basis.
Quantification of P. falciparum parasite density by polymerase chain reaction (qPCR)

DNA was extracted directly from 180 µL of freshly collected venous whole blood using Quick-DNA Miniprep kits (Zymo Research, Irvine, USA) and eluted with 50 µL of elution buffer as recommended by manufacturer. DNA samples were kept at −20°C until analysis using the Bio-Rad CFX96 Real-Time PCR System (Bio-Rad Laboratories, California, USA). The PlasQ assay previously described by Schindler et al. [48] was used for quantification of Plasmodium spp. and Pf parasites in the venous blood sample. This multiplex assay targets two independent Plasmodium genes namely the Pan-Plasmodium 18S rDNA sequence (Pspp18S) and the Pf-specific acidic terminal sequence of the var genes (PfvarATS). The human Ribonuclease P gene (HsRNaseP) was used as a DNA extraction and qPCR amplification control. All qPCR assays were run in duplicate and both non-template control (molecular grade nuclease-free water) and Pf 3D7 DNA were included in each PCR run as negative and positive controls, respectively. For the parasite density estimation, a serial dilution was made according to the 1st Word Health Organization (WHO) International Standard for Pf DNA Amplification Technique (NIBSC code: 04/176) to establish a calibration curve with the parasite densities ranging between 0.01 to 10,000 p/µL. The actual parasite density of the tested sample by qPCR was then estimated from the calibration curve's y-intercept and slope. The lower limit of detection for this qPCR assay was 50 copies/mL. The sample was considered Pf positive if each of the two replicates for both PfvarATS and 18S RNA gene targets had quantitation cycles (Cq) < 40 and Cq < 28 for qPCR amplification control (HsRNaseP). In case of a discrepancy between duplicates, the assay was repeated, with at least two positive replicates out of four considered a positive result. The final results were used for the qPCR-based estimate of parasite density.

Controlled human malaria infection (CHMI)

From October 2018 to March 2019, 95 healthy Equatoguinean adults underwent CHMI (Jongo SA, AJTMH, in press)[47]. Prior to CHMI, a full 3-day course of artemether lumefantrine treatment was given to all volunteers. Eligibility criteria for CHMI were met if volunteers had received a complete regimen of PfSPZ Vaccine and were negative for malaria infection at the time of CHMI. During the ward observation period, volunteers were monitored daily for Pf parasitemia starting on day 8 to detect the parasite early and prevent the development of symptoms. Two milliliters of venous whole blood were collected in EDTA tubes daily on days 8–20 post-infection and transported to the laboratory in cooling boxes (4-8°C) within 30 min of collection. One mL blood was used for examining malaria parasites positivity and density by TBS and qPCR within 4 h of collection and 1 mL was stored at -80°C for 8 months before retrospectively analysing samples using uRDT and cRDT. The standard artemether/lumefantrine treatment was given to subjects once malaria parasites were detected by TBS or on day 28 post CHMI for volunteers who remained negative throughout the post-CHMI follow-up period. Positive TBS results confirmed by qPCR were used as the end-point for initiating participant malaria treatment and termination of ward visits and further diagnostic sample collection. Volunteers diagnosed as malaria positive by TBS during 28 days of CHMI follow-up were considered eligible for participation in this malaria diagnostics study.
Sampling and statistical analyses

We aimed to demonstrate differences in performance sensitivity between TBS, cRDT, and uRDT methods in detecting low density malaria infection, using qPCR as the gold standard for sensitivity. To obtain sufficient samples, we used Epi Info 7 software to calculate the sample size assuming the following parameters: minimum sensitivity of 90% for a reference method (qPCR) and 80% for cRDT, a 95% confidence and 90% power to detect a maximum sensitivity difference of 10%. We used cRDT in determining sample size due to its wider application in the field [49]. This gave a minimum required sample size of 267 samples. Sensitivity of the various tests was compared using Fisher's exact test.

Participating individuals were to be observed from day 8 after challenge until the day of first positivity by TBS. Tests were not performed on samples collected after TBS diagnosis since volunteers were treated and positive results could still occur for RDTs and qPCR due to residual parasite material, which would confound results. Out of the eligible volunteers we planned to randomly select a subset of 24 individuals to meet sample size requirements while maintaining the distribution of parasite densities observed using qPCR.

We assessed the distributions of the complete set, sampled and unsampled subsets to ensure they presented similar structure. Sampled individuals provided a total of 279 observed individual sample time points.

For our analysis, a multiplex qPCR targeting Pspp18S and PfvarATS was designated as the reference for detection of infection against which TBS, uRDT and cRDT positivity could be compared. A two-tailed Fishers exact test was used to determine the significant differences between the sensitivities of the various diagnostic tests. All samples were included in our sensitivity analysis of diagnostics. Only positive samples by either TBS, uRDT and cRDT were included in our analyses of the overall geometric mean (geomean) of parasite density of positive results and the geomean of parasite density at time of first detection (prepatent period). If a TBS, uRDT or cRDT test was negative, the respective sample point was deemed not applicable for the geomean parasite density and time to first detection analysis. TBS and qPCR both provided density measurements but we considered TBS to be more reliable as it did not involve conversion from gene copy number using a reference standard.

Results were recorded by trained and qualified laboratory staff on case report forms (CRFs) during the CHMI ward observation period and later entered onto an Excel spreadsheet (Microsoft, Office 2019 Ver 16). All samples were assigned a sample specific number that was linked to each volunteer ID. No personal information was recorded on laboratory CRFs and for the laboratory staff the connection of each sample with the corresponding donor volunteer was not possible. Retrospective RDT results were recorded on the same Excel spreadsheet. The geomean and geomean confidence intervals of parasite densities were calculated using R 4.0.1. Sensitivity and 95% confidence intervals for all diagnostic methods were calculated in R 4.0.1 using the epiR package [50].
Results

Overview

A total of 48 volunteers were diagnosed positive for malaria by reference method (qPCR), qualifying them for inclusion in the malaria diagnostic comparison. Individuals had an average of 12 time points of observation-days (range 8 to 17), with each day-test-record representing an independent observation since each was obtained from a newly collected whole blood sample. Out of the eligible volunteers we randomly selected 24 (50%) individuals maintaining the distribution of parasite densities observed using qPCR. The subset of selected samples were evaluated against unselected samples by parasite density distribution and variance; and the selected subset was determined to be an appropriate representation (supplementary Figures1, 2 and 3; and supplementary Tables 1).

A total of 279 samples were collected from the 24 selected study participants; 123 and 156 samples were positive and negative for Pf infection by qPCR, respectively. All 156 samples negative for Pf by qPCR were also negative by TBS, uRDT and cRDT demonstrating 100% specificity for these tests. In total, 24 of 123 positive samples were detected by TBS, 21 by uRDT and 10 by cRDT, providing sensitivities of 19.5% (95% CI 13.1% – 27.8%), 17.1% (95% CI 11.1% – 25.1%), and 8.1% (95% CI 4.2% – 14.8%), respectively. qPCR detected more positives than any of the other tests (p<0.001) and TBS and cRDT were both more sensitive than cRDT (TBS vs. cRDT, p=0.015 by Fishers Exact two-tailed; uRDT vs. cRDT, p=0.053). TBS detected 61.9% (13/21) of uRDT positive infections, while uRDT detected 54.2% (13/24) of TBS positive infections. The uRDT detected 100% (10/10) of cRDT positive infections while TBS detected 90% (9/10). The cRDT detected 47.6% (10/21) of uRDT positive infections and 37.5% (9/24) of TBS positive infections. The summary of the findings are depicted in a Venn-diagram (Figure 1).

Parasite density by qPCR was calculated using a standard curve generated using a WHO reference sample that related copy number to density, and ranged from 0.14 – 603.8 p/µL, with a geomean of 2.57 p/µL. Using this scale TBS detected parasites in the range of 5.9 – 603.8 p/µL (geomean 97.6 p/µL) and uRDT detected parasites in the range of 0.8 – 603.8 p/µL (geomean 103.4 p/µL), compared to cRDT, which detected parasites in the range of 2.6 – 603.8 p/µL (geomean = 149.4 p/µL) (Table 2).
Table 1
Number of positive and negative samples, overall geomean and sensitivity of TBS, uRDT, and cRDT compared to the reference qPCR method. All cases were low parasite density infections that occurred during CHMI follow-up. Parasite densities of positive samples by qPCR ranged from 0.14 – 603.84 p/µL.

| Diagnostic test | TBS (+) | TBS (-) | uRDT (+) | uRDT (-) | cRDT (+) | cRDT (-) | Total |
|-----------------|---------|---------|----------|----------|----------|----------|-------|
| PCR Pf (+)      | 24      | 99      | 21       | 102      | 10       | 113      | 123   |
| PCR Pf (-)      | 0       | 156     | 0        | 156      | 0        | 156      | 156   |
| **Total**       | **24**  | **255** | **21**   | **258**  | **10**   | **269**  | **279**|
| **Range of positive samples (p/µL by qPCR)** | 5.9 – 603.8 | 0.8 – 603.8 | 2.6 – 603.8|
| **Geomean of positive samples (p/µL by qPCR)** | 97.6 | 103.4 | 149.4 |
| **Sensitivity (%)** | 19.5% (13.1 – 27.8) | 17.1% (11.1 – 25.1) | 8.1% (4.2 – 14.8) |

Parasite density by TBS ranged from 3.7 – 201.8 p/µL (geomean 12.81 p/µL). Using this scale, uRDT detected parasites in the range of 3.7 – 201.8 p/µL (geomean 20.6 p/µL), compared to cRDT, which detected parasites in the range of 5.6 – 201.8 p/µL (geomean = 30.2 p/µL) (Table 2).

Table 2
Number of positive and negative samples of uRDT and cRDT compared to TBS. All cases were low parasite density infections that occurred during CHMI follow-up. Parasite densities of positive samples by TBS ranged from 3.7 – 201.8 p/µL.

| Diagnostic test | uRDT (+) | uRDT (-) | cRDT (+) | cRDT (-) | Total |
|-----------------|----------|----------|----------|----------|-------|
| TBS (+)         | 13       | 11       | 9        | 15       | 24    |
| TBS (-)         | 8        | 247      | 1        | 254      | 255   |
| **Total**       | **21**   | **258**  | **10**   | **269**  | **279**|
| **Geomean of positive samples (p/µL by TBS)** | 20.6 | 30.2 |
| **Sensitivity (%)** | 54.2% (33.2 – 73.8) | 37.5% (19.6 – 59.2) |

Examining just infections positive by both qPCR and TBS, the geomean ratio established from parasite density of qPCR and TBS (qPCR/TBS) was 7.62 p/µL (Table 3). PCR detects gene copy number in a specimen and each Pf genome has at least 5-8 copies [51], and in addition qPCR can detect free DNA in a specimens. These variables make it difficult to calculate parasite density accurately using qPCR, even when using a standard curve to convert copy numbers to density based on a WHO reference sample. TBS
detects parasites, regardless of how many genes and nuclei are present, but is hindered by the possible loss of significant numbers of parasites during processing [52]. We suspect that all these factors contributed to the higher densities found using qPCR compared to TBS.

Table 3
Comparison of parasite densities established from qPCR and TBS using paired samples in which both tests were positive. All cases were low parasite density infections that occurred during CHMI follow-up. The Ratio was determined by dividing the geometric mean of qPCR by geometric mean of TBS (qPCR/TBS). N = 24

| qPCR (Pf/µL) | TBS (Pf/µL) | Ratio qPCR/TBS |
|-------------|-------------|----------------|
| GeoMean     | 97.57       | 12.81          | 7.62           |
| Range       | [5.86 - 603.84] | [3.70 - 201.80] | [0.07 - 53.84] |

Next, we stratified the TBS, uRDT and cRDT diagnostic test sensitivity by ranges of parasite density of qPCR (Table 4). None of the three diagnostic tests detected Pf infections below 1 p/µL as determined by qPCR. For TBS, the lowest parasite density detected as quantified by qPCR was 5.86 p/uL and as quantified by TBS was 3.7 p/uL. At parasite densities of 1 – 50 p/uL, uRDT and TBS appeared roughly equally sensitive and both appeared more sensitive than cRDT. At parasite densities of 51 – 100 p/µL by qPCR, uRDT appeared more sensitive (71% [95% CI: 29% – 96%]) compared to TBS and cRDT with, 29% (4% – 71%) and 29% (4% – 71%) sensitivity, respectively. However, numbers were small and differences in sensitivity amongst the tests across the density categories were not statistically significant.

Table 4
Comparison of TBS, uRDT and cRDT sensitivity stratified by parasite density (p/µL) as determined by qPCR. Sensitivities for each density interval can be compared by diagnostic method. All cases were low parasite density samples that occurred during CHMI follow-up and were 100% specific compared to qPCR.

| Group density (p/µL) | # samples qPCR (+) | TBS (+) | TBS sensitivity (95% CI) | uRDT | uRDT sensitivity (95% CI) | cRDT | cRDT sensitivity (95% CI) |
|----------------------|---------------------|---------|--------------------------|------|---------------------------|------|--------------------------|
| <1                   | 37                  | 0       | -                        | 0    | -                         | 0    | -                        |
| 1-10                 | 38                  | 1       | 3% (0 - 14)              | 1    | 3% (0 - 14)               | 1    | 3% (0 - 14)              |
| 11-50                | 25                  | 7       | 28% (12 - 49)           | 4    | 16% (5 - 36)             | 0    | -                        |
| 51-100               | 7                   | 2       | 29% (4 - 71)           | 5    | 71% (29 - 96)            | 2    | 29% (4 - 71)             |
| >100                 | 16                  | 14      | 88% (62 - 98)          | 11   | 69% (41 - 89)            | 7    | 44% (20 - 70)            |
Finally, we stratified the uRDT and cRDT diagnostic test sensitivity by ranges of parasite density measured by TBS (Table 5). At parasite densities range between 1-50 p/µL, uRDT had higher sensitivity compared to cRDT; 33% (95% CI 12-62) for 1-10 p/µL and 100% (95% CI 48-100) at 11-50 p/µL compared to 13% (95% CI 2-40) for 1-10 p/µL and 80% (95% CI 28-99) at 11-50 p/µL respectively. Above 50 p/µL, both uRDT and cRDT had the same sensitivity but again numbers were too small to allow a meaningful comparison.

Table 5

Comparison of uRDT and cRDT sensitivity stratified by parasite density (p/µL), as determined by TBS. Sensitivities for each density interval can be compared by diagnostic method. All cases were low parasite density samples that occurred during CHMI follow-up.

| Group density (p/µL) | # samples TBS (+) | uRDT (+) | uRDT sensitivity (95% CI) | cRDT (+) | cRDT sensitivity (95% CI) |
|----------------------|-------------------|----------|---------------------------|----------|---------------------------|
| <1                   | 0                 | 0        | -                         | 0        | -                         |
| 1 - 10               | 15                | 5        | 33% (12 – 62)             | 2        | 13% (2 – 40)              |
| 11 - 50              | 5                 | 5        | 100% (48 -100)            | 4        | 80% (28 – 99)             |
| 51 - 100             | 2                 | 2        | 100% (16 -100)            | 2        | 100% (16 -100)            |
| >100                 | 2                 | 1        | 50% (1 – 99)              | 1        | 50% (1 - 99)              |

We further examined the range and distribution of parasite densities of samples determined to be positive by qPCR (n=123), by TBS (n=24), uRDT (n=21) and cRDT (n=10) over the follow up period for the 24 volunteers who were TBS positive. TBS and uRDT recorded a trend for lower geometric parasite densities detected compared to cRDT, which did not reach statistical significance [p = 0.19 and p = 0.26, respectively] (Figure 2).

**Time to first detection**

To investigate the efficiency of cRDT and uRDT to detect asexual blood stage parasites during CHMI follow-up, we compared the median time to first detection of parasites (in days) by qPCR, TBS, uRDT and cRDT. There was no evidence to support differences in prepatent period when using TBS, uRDT and cRDT since these methods all reported a median of 18.0 days to first parasite detection. The median days to detection of asexual blood stage parasitemia by qPCR was 14.5, 3.5 days earlier than TBS, uRDT and cRDT (p < 0.001 log-rank test) (Figure 3).

**Discussion**

As progress against malaria is made, asymptomatic infections at lower parasite densities become a significant challenge for malaria control and elimination efforts due to their contribution to ongoing
transmission [2, 3, 53, 54]. Mass testing of a population with treatment of those found positive is one approach to address this problem [55]. However, it is difficult to diagnose low density parasitemias and the most sensitive and therefore the best method, qPCR, is expensive, requires special laboratories and skilled personnel. Thus, the development of an inexpensive rapid test with equivalent sensitivity would be of great benefit, especially as conventional rapid diagnostic tests (cRDTs) are significantly less sensitive. Other applications could also benefit from a simple, rapid test that is more sensitive than cRDTs, such as follow-up after CHMI, a procedure for evaluating vaccine and drug efficacy [42, 43, 56–59] and for exploring innate and acquired immunity [60–64], that is now gaining attention by investigators due to the availability of cryopreserved infectious sporozoites that can be used by any clinical center without the need for infectious mosquitoes [65–67]. In CHMI, detection of low parasite densities is useful because it allows the identification and treatment of positive study subjects earlier in the course of their parasitemia thereby preventing or ameliorating clinical manifestations. Therefore, on many fronts, there is a need to develop simpler, highly sensitive methods to diagnose low parasite densities that could augment the success of mass testing and treatment, promote epidemiological studies and simplify and lessen the costs associated with CHMI.

cRDTs have been a tremendous boon to diagnosing clinical malaria, where parasite densities are relatively high and the tests adequately sensitive. uRDT have recently been developed, and might extend the usefulness of RDTs particularly in low to moderate transmission areas, in pre-elimination settings, and in experimental uses such as CHMI follow-up. For example, it has been reported that a uRDT is significantly more sensitive than cRDT and TBS, detecting PfHRP2 at parasite densities as low as 0.1 – 1.0 p/µL in culture-derived samples [6, 22]. For this reason, we tested uRDTs and cRDT during follow-up in a CHMI trial, and compared their sensitivities to those of TBS, using qPCR as the reference standard.

We indeed found that both TBS and the uRDT were more sensitive than the cRDT. However, our data demonstrate that in samples from malaria-experienced subjects undergoing CHMI with parasites that contained PfHRP2, the sensitivity of uRDT was about 10-fold lower that reported for culture-derived samples. None of 37 specimens less than 1.0 p/uL by qPCR were identified. In 15 specimens that had 1-10 p/uL by TBS, uRDT identified 5 (33%), and cRDT 2 (13%). In 5 specimens with 11-50 parasites/uL by TBS, uRDT identified all 5 as positive (100%) and cRDT identified 4 (80%). Overall, uRDT and TBS gave similar results, and both tests were more sensitive than cRDTs in a setting of CHMI with PfHRP2-containing parasites.

Currently, CHMI requires highly trained clinical and laboratory staff including expert microscopists. Considering the comparable outcomes of TBS and uRDT in our study, uRDT could be considered to replace TBS microscopy, especially in settings with inexperienced microscopists. However, parasite density estimation using qPCR has become a standard method utilized in many malaria studies [68, 69] and has been particularly useful in CHMI follow-up where it can detect parasitemia earlier than TBS and allow treatment before signs and symptoms of clinical malaria develop [70–72]. Thus, uRDT would need to show advantages over TBS in the early identification of positive study subjects, as TBS itself is now being supplanted by qPCR. As expected, we showed a significant difference in prepatent period amongst
qPCR on the one hand and TBS, uRDT and cRDT on the other, confirming that qPCR is the most sensitive diagnostic method. We further demonstrated that the median time to first malaria parasite detection by qPCR was 3.5 days earlier (14.5 days) compared to the other tests, and also that TBS, uRDT and cRDT were substantially equivalent to each other, each providing a prepatent period of 18 days. Similar to our findings, the CHMI studies conducted in semi-immune participants [42, 56, 71, 73] and in malaria naïve participants [74] have reported comparable prepatent periods using qPCR as the reference method. The fact that in this setting of progressively rising parasitemias, the prepatent periods calculated by TBS, uRDT and cRDT were similar even though positive samples diagnosed by uRDT had a lower overall geometric mean of parasite density by qPCR than did TBS, suggests that uRDT may not have any particular advantage over TBS other than reduced costs and easier performance, or even over cRDT, as in our study the day of treatment would not have been affected had it been cRDT- rather than TBS-based.

In a field setting, the greater sensitivity of uRDTs over cRDTs could allow the detection of more asymptomatic carriers. We did not directly examine this question in our CHMI-based study. However, our results showed that despite uRDTs being hailed as a significant improvement in malaria diagnostics, leading to increased sensitivity and specificity, satisfactory RDT performance for parasite density infections <10 p/μL remains elusive. Field studies of mass testing and treatment are needed to further explore the potential contribution of uRDT in identifying and treating asymptomatic carriers with low parasite densities contributing to ongoing transmission.

One important consideration for evaluating RDTs based on the detection of PfHRP2 is the increase in prevalence of parasites carrying PfHRP2 deletions, not only in Southeast Asia [75], but within the study area as well [11]. In 2018, approximately 65% of all suspected malaria cases in public health facilities in sub-Saharan African were tested with RDTs (~150 million cases) [1]. During CHMI, we used a standardized infectious PfSPZ dose of PfSPZ Challenge (NF54), a parasite that expresses PfHRP2 to initiate the infection. Conducting a similar study in hospitals and field environment with important confounders such as a deleted PfHRP2 gene would likely have had different results.

**Limitations**

Anticoagulated (EDTA) fresh whole blood was used for to prepare samples for qPCR and TBS assessments. Anticoagulated (EDTA) cryopreserved (temperature of -80°C) whole blood held for 8 months and thawed was used to prepare samples for uRDT and cRDTs. It is possible, but unlikely that HRP2 degraded during storage. Whole blood samples were temperature monitored during storage. When proper procedures are followed for long-term storage of whole blood, the quality of DNA, RNA or HRP2 is not compromised [6, 76]. Considering our strict temperature monitoring and only thawing samples once for processing, the difference in quality of samples over time is unlikely to have been different.

Another limitation is the discrepancy in parasite densities estimated by qPCR and TBS. qPCR may have overestimated parasite density due to variable numbers of copies of the amplification target and the
persistence of nucleic acid from non-viable parasites (xxx), and TBS may have underestimated parasite density due to the loss of parasites during processing.

Because the research subjects were semi-immune, they may have had variable levels of anti-HRP2 antibodies, which could have affected results [77, 78]. Some may also have had ongoing infections at the time of clearance with artemether/lumefantrine prior to CHMI. Although we did not measure the level of HRP2 in the participants before and after CHMI, none of the individuals were RDT positive between days 8 and 14 after CHMI. Therefore, we believe that this factor did not affect the current performance comparison.

**List Of Abbreviations**

- **cRDT**: Conventional rapid diagnostic test
- **PfHRP2**: Histidine-rich protein 2
- **Pf**: *Plasmodium falciparum*
- **PAN**: Pan-malaria antigen
- **TBS**: Thick blood smear
- **p/µL**: parasites per microlitre
- **qPCR**: Quantitative polymerase chain reaction
- **uRDT**: Ultra-sensitive rapid diagnostic test
- **CHMI**: Controlled human malaria infection
- **DVI**: Direct venous inoculation
- **BIMEP**: Bioko Island malaria Elimination Project
- **CRFs**: Clinical report forms

**Declarations**

**Ethics approval and consent to participate**

Ethics approval was obtained to conduct the malaria vaccine trial from the Comité Ética Nacional de Guinea Ecuatorial (CENGE) in Equatorial Guinea, Ifakara Health Institute (IHI) in Tanzania, Ethikkommission Nordwest- und Zentralschweiz (EKNZ) in Switzerland, and Prime IRB in the USA. Included in this approval was the use of participant biological samples for future use if informed consent
was provided. No identifying information was accessible to laboratory staff and was not included in any database or analysis for this study.

**Consent for publication**

Not applicable

**Availability of data and materials**

The datasets used and/or analysis during the current study are available from the corresponding author on reasonable request.

**Competing interests**

The authors declare that they have no competing interests.

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This work was funded by the Bioko Island Malaria Elimination Program. The funders had no role in study design, data collection, data analysis, data interpretation, decision to publish, or preparation of the manuscript.

**Author's contributions**

SLH posed the research question and the applicability of the CHMI follow-up as a means to evaluate a potential uRDT application. MM and CD conceived and designed the evaluation plan for this study. MM, JR, AD, LA, EN, COD, TS, VU, AM, AH, MSAL, BP, and MAOE were responsible for collecting and processing volunteer samples. MRR, CCF, GAG, RC, ES, PFB, LWPC, BKLS, TLR, BM, MT, SA, CM, SLH, SJ sponsored and oversaw the management of the PfSPZ Vaccine trial and CHMI. SJ was the principle investigator. CD managed all laboratory activities and supervised this study. JCM managed quality control and quality assurance of all documentation. TCS programmed and conducted the statistical analysis of the data and generated all tables and figures. MM and TCS wrote the manuscript in consultation with PFB, SLH and TLR. All authors read and approved the final version of the manuscript.

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Summary Key Statement

TBS has been the classical approach to malaria diagnosis for clinical use, malaria control programs, research studies such as CHMI and field epidemiology. TBS can distinguish the five malaria species that infect humans, which cannot yet be achieved by using RDTs or a single reaction qPCR, and provides a reasonable estimate of parasite density. TBS, however, requires laboratories that support and maintain microscopes, staining solutions and human resources with the requisite microscopy skills. qPCR, with much greater sensitivity, is now supplanting TBS for many applications such as CHMI follow-up, but also requires a high level of laboratory capability and involves higher costs than TBS. cRDTs have therefore been a welcome addition to malaria diagnostics and in many places have supplanted TBS for the clinical diagnosis of malaria, where parasite densities are high, but have not been useful for applications requiring greater sensitivity. In this study, we compared qPCR, TBS, a cRDT to a new uRDT advertised as rivaling qPCR in sensitivity, to assess its value during CHMI follow-up, an application where early diagnosis and treatment is important to reduce the severity of adverse events. Our major conclusions were that for this specific application, while uRDT was better than cRDT, and approached TBS in sensitivity, it did not close the gap with respect to qPCR, and thus could be considered for replacing TBS only in CHMI studies unable to use qPCR or TBS due to resource limitations. The value of uRDT in field studies, particularly in mass testing and treatment, requires further study.

References

1. Organization WH: Geneva: 2019. World malaria report 2019. 2020.
2. Slater HC, Ross A, Ouédraogo AL, White LJ, Nguon C, Walker PGT, Ngor P, Aguas R, Silal SP, Dondorp AM, et al: Assessing the impact of next-generation rapid diagnostic tests on Plasmodium falciparum malaria elimination strategies. Nature 2015, 528:S94-S101.
3. Hemingway J, Shretta R, Wells TN, Bell D, Djimdé AA, Achee N, Qi G: Tools and strategies for malaria control and elimination: what do we need to achieve a grand convergence in malaria? Journal PLoS biology 2016, 14:e1002380.
4. Lo E, Zhou G, Oo W, Afrane Y, Githeko A, Yan G: Low parasitemia in submicroscopic infections significantly impacts malaria diagnostic sensitivity in the highlands of Western Kenya. PLoS One 2015, 10:e0121763.
5. Danwang C, Kirakoya-Samadoulougou F, Samadoulougou S: Assessing field performance of ultrasensitive rapid diagnostic tests for malaria: a systematic review and meta-analysis. Malaria Journal 2021, 20:245.
6. Das S, Peck RB, Barney R, Jang IK, Kahn M, Zhu M, Domingo GJ: Performance of an ultra-sensitive Plasmodium falciparum HRP2-based rapid diagnostic test with recombinant HRP2, culture parasites, and archived whole blood samples. Malaria Journal 2018, 17:118.
7. Mathison BA, Pritt BS: Update on Malaria Diagnostics and Test Utilization. Journal of Clinical Microbiology 2017, 55:2009–2017.
8. Adams M, Joshi SN, Mbambo G, Mu AZ, Roemmich SM, Shrestha B, Strauss KA, Johnson NE, Oo KZ, Hlaing TM, et al: An ultrasensitive reverse transcription polymerase chain reaction assay to detect asymptomatic low-density Plasmodium falciparum and Plasmodium vivax infections in small volume blood samples. *Malaria Journal* 2015, **14**:520.

9. Bousema T, Okell L,Felger I, Drakeley C: Asymptomatic malaria infections: detectability, transmissibility and public health relevance. *Nature Reviews Microbiology* 2014, **12**:833–840.

10. Laban NM, Kobayashi T, Hamapumbu H, Sullivan D, Mharakurwa S, Thuma PE, Shiff CJ, Moss WJ, for Southern Africa International Centers of Excellence for Malaria R: Comparison of a PfHRP2-based rapid diagnostic test and PCR for malaria in a low prevalence setting in rural southern Zambia: implications for elimination. *Malaria Journal* 2015, **14**:25.

11. Berzosa P, González V, Taravillo L, Mayor A, Romay-Barja M, García L, Ncogo P, Riloha M, Benito A: First evidence of the deletion in the pfhrp2 and pfhrp3 genes in Plasmodium falciparum from Equatorial Guinea. *Malaria Journal* 2020, **19**:99.

12. Koita OA, Doumbo OK, Ouattara A, Tall LK, Konaré A, Diakité M, Diallo M, Sagara I, Masinde GL, Doumbo SN, et al: False-Negative Rapid Diagnostic Tests for Malaria and Deletion of the Histidine-Rich Repeat Region of the hrp2 Gene. *The American Journal of Tropical Medicine and Hygiene* 2012, **86**:194–198.

13. Berzosa P, de Lucio A, Romay-Barja M, Herrador Z, González V, García L, Fernández-Martínez A, Santana-Morales M, Ncogo P, Valladares B, et al: Comparison of three diagnostic methods (microscopy, RDT, and PCR) for the detection of malaria parasites in representative samples from Equatorial Guinea. *Malaria Journal* 2018, **17**:333.

14. Hofmann NE, Gruenberg M, Nate E, Ura A, Rodriguez-Rodriguez D, Salib M, Mueller I, Smith TA, Laman M, Robinson LJ, Felger I: Assessment of ultra-sensitive malaria diagnosis versus standard molecular diagnostics for malaria elimination: an in-depth molecular community cross-sectional study. *The Lancet Infectious Diseases* 2018, **18**:1108–1116.

15. Amaral LC, Robortella DR, Guimarães LFF, Limongi JE, Fontes CJF, Pereira DB, de Brito CFA, Kano FS, de Sousa TN, Carvalho LH: Ribosomal and non-ribosomal PCR targets for the detection of low-density and mixed malaria infections. *Malaria Journal* 2019, **18**:154.

16. Canier L, Khim N, Kim S, Sluydts V, Heng S, Dourng D, Eam R, Chy S, Khean C, Loch K, et al: An innovative tool for moving malaria PCR detection of parasite reservoir into the field. *Malaria Journal* 2013, **12**:405.

17. Naeem MA, Ahmed S, Khan SA: Detection of asymptomatic carriers of malaria in Kohat district of Pakistan. *Malaria Journal* 2018, **17**:44.

18. Mabey D, Peeling RW, Ustianowski A, Perkins MD: Diagnostics for the developing world. *Nature Reviews Microbiology* 2004, **2**:231–240.

19. Urdea M, Penny LA, Olmsted SS, Giovanni MY, Kaspar P, Shepherd A, Wilson P, Dahl CA, Buchsbaum S, Moeller G, Hay Burgess DC: Requirements for high impact diagnostics in the developing world. *Nature* 2006, **444**:73–79.
20. Ohrt C, Sutamihardja MA, Tang D, Kain KC: Impact of microscopy error on estimates of protective efficacy in malaria-prevention trials. The Journal of infectious diseases 2002, 186:540–546.

21. Payne D: Use and limitations of light microscopy for diagnosing malaria at the primary health care level. J Bulletin of the World Health Organization 1988, 66:621.

22. Das S, Jang IK, Barney B, Peck R, Rek JC, Arinaitwe E, Adrama H, Murphy M, Imwong M, Ling CL, et al: Performance of a High-Sensitivity Rapid Diagnostic Test for Plasmodium falciparum Malaria in Asymptomatic Individuals from Uganda and Myanmar and Naive Human Challenge Infections. The American Journal of Tropical Medicine and Hygiene 2017, 97:1540–1550.

23. Landier J, Haohankhunnatham W, Das S, Konghahong K, Christensen P, Raksuansak J, Phattharakokoedburn P, Kajeechiwa L, Thwin MM, Jang IK, et al: Operational Performance of a Plasmodium falciparum Ultrasensitive Rapid Diagnostic Test for Detection of Asymptomatic Infections in Eastern Myanmar. Journal of Clinical Microbiology 2018, 56:e00565-00518.

24. Yeung S, McGregor D, James N, Kheang ST, Kim S, Khim N, Ly P, Sovannaroath S, Witkowski B: Performance of Ultrasensitive Rapid Diagnostic Tests for Detecting Asymptomatic Plasmodium falciparum. The American Journal of Tropical Medicine and Hygiene 2020, 102:307–309.

25. Acquah FK, Donu D, Obboh EK, Bredu D, Mawuli B, Amponsah JA, Quartey J, Amoah LE: Diagnostic performance of an ultrasensitive HRP2-based malaria rapid diagnostic test kit used in surveys of afebrile people living in Southern Ghana. Malaria Journal 2021, 20:125.

26. Manjurano A, Omolo JJ, Lyimo E, Miyaye D, Kishamawe C, Matemba LE, Massaga JJ, Chalamalucha J, Kazyoba PE: Performance evaluation of the highly sensitive histidine-rich protein 2 rapid test for Plasmodium falciparum malaria in North-West Tanzania. Malaria Journal 2021, 20:58.

27. McMorrow ML, Aidoo M, Kachur SP: Malaria rapid diagnostic tests in elimination settings—can they find the last parasite? Clinical Microbiology and Infection 2011, 17:1624–1631.

28. Wu L, van den Hoogen LL, Slater H, Walker PGT, Ghani AC, Drakeley CJ, Okell LC: Comparison of diagnostics for the detection of asymptomatic Plasmodium falciparum infections to inform control and elimination strategies. Nature 2015, 528:S86-S93.

29. Lin JT, Saunders DL, Meshnick SR: The role of submicroscopic parasitemia in malaria transmission: what is the evidence? Trends in Parasitology 2014, 30:183–190.

30. Plucinski MM, Candrinho B, Dimene M, Colborn J, Lu A, Nace D, Zulliger R, Rogier E: Assessing Performance of HRP2 Antigen Detection for Malaria Diagnosis in Mozambique. Journal of Clinical Microbiology 2019, 57:e00875-00819.

31. Galata B, Mayor A, Gupta H, Balanza N, Jang IK, Nhamussua L, Simone W, Cisteró P, Chidimatembue A, Munguambe H, et al: Field performance of ultrasensitive and conventional malaria rapid diagnostic tests in southern Mozambique. Malaria Journal 2020, 19:451.

32. Unwin VT, Ahmed R, Noviyanti R, Puspitasari AM, Utami RAS, Trianty L, Lukito T, Syafruddin D, Poespoprodjo JR, Santana-Morales MA, et al: Use of a highly-sensitive rapid diagnostic test to screen for malaria in pregnancy in Indonesia. Malaria Journal 2020, 19:28.
33. Slater HC, Ross A, Felger I, Hofmann NE, Robinson L, Cook J, Gonçalves BP, Björkman A, Ouedraogo AL, Morris U, et al: The temporal dynamics and infectiousness of subpatent Plasmodium falciparum infections in relation to parasite density. *Nature Communications* 2019, **10**:1433.

34. Felger I, Maire M, Bretscher MT, Falk N, Tiaden A, Sama W, Beck H-P, Owusu-Agyei S, Smith TA: The dynamics of natural Plasmodium falciparum infections. *PloS one* 2012, **7**:e45542.

35. Mordmüller B, Supan C, Sim KL, Gómez-Pérez GP, Ospina Salazar CL, Held J, Bolte S, Esen M, Tschan S, Joanny F, et al: Direct venous inoculation of Plasmodium falciparum sporozoites for controlled human malaria infection: a dose-finding trial in two centres. *Malaria Journal* 2015, **14**:117.

36. Laurens MB, Duncan CJ, Epstein JE, Hill AV, Komisar JL, Lyke KE, Ockenhouse CF, Richie TL, Roestenberg M, Sauerwein RW, et al: A consultation on the optimization of controlled human malaria infection by mosquito bite for evaluation of candidate malaria vaccines. *Vaccine* 2012, **30**:5302–5304.

37. Epstein JE, Paolino KM, Richie TL, Sedegah M, Singer A, Ruben AJ, Chakravarty S, Stafford A, Ruck RC, Eappen AG, et al: Protection against Plasmodium falciparum malaria by PfSPZ Vaccine. *JCI insight* 2017, **2**:e89154-e89154.

38. Lyke KE, Ishizuka AS, Berry AA, Chakravarty S, DeZure A, Enama ME, James ER, Billingsley PF, Gunasekera A, Manoj A, et al: Attenuated PfSPZ Vaccine induces strain-transcending T cells and durable protection against heterologous controlled human malaria infection. *PNAS* 2017, **114**:2711–2716.

39. Seder RA, Chang L-J, Enama ME, Zephir KL, Sarwar UN, Gordon IJ, Holman LA, James ER, Billingsley PF, Gunasekera A, et al: Protection Against Malaria by Intravenous Immunization with a Nonreplicating Sporozoite Vaccine. 2013, **341**:1359–1365.

40. Ishizuka AS, Lyke KE, DeZure A, Berry AA, Richie TL, Mendoza FH, Enama ME, Gordon IJ, Chang LJ, Sarwar UN, et al: Protection against malaria at 1 year and immune correlates following PfSPZ vaccination. *Nat Med* 2016, **22**:614–623.

41. Mordmuller B, Surf R, Lagler H, Chakravarty S, Ishizuka AS, Lalremruata A, Gmeiner M, Campo JJ, Esen M, Ruben AJ, et al: Sterile protection against human malaria by chemoattenuated PfSPZ vaccine. *Nature* 2017, **542**:445–449.

42. Jongo SA, Urbano V, Church LWP, Oloto A, Manock SR, Schindler T, Mtoro A, KC N, Hamad A, Nyakarungu E, et al: Immunogenicity and Protective Efficacy of Radiation-Attenuated and Chemo-Attenuated PfSPZ Vaccines in Equatoguinean Adults *J The American Journal of Tropical Medicine and Hygiene* 2021, **104**:283–293.

43. Jongo SA, Church LWP, Mtoro AT, Schindler T, Chakravarty S, Ruben AJ, Swanson PA, Kassim KR, Mphina M, Tumbo A-M, et al: Increase of Dose Associated With Decrease in Protection Against Controlled Human Malaria Infection by PfSPZ Vaccine in Tanzanian Adults. *Clinical Infectious Diseases* 2020, **71**:2849–2857.

44. Billingsley PF, Maas CD, Oloto A, Schwabe C, García GA, Rivas MR, Hergott DEB, Daubenberger C, Saverino E, Chaouch A, et al: The Equatoguinean Malaria Vaccine Initiative: From the Launching of a
45. Nchama VUNN, Said AH, Mtoro A, Bidjimi GO, Owono MA, Maye ERM, Mangue MEO, Okomo GNN, Pasialo BEN, Ondo DM, et al: *Incidence of Plasmodium falciparum malaria infection in 6-month to 45-year-olds on selected areas of Bioko Island, Equatorial Guinea*. *Malaria Journal* 2021, 20:322.

46. Olotu A, Urbano V, Hamad A, Eka M, Chemba M, Nyakarungu E, Raso J, Eburi E, Mandumbi DO, Hergott D, et al: *Advancing Global Health through Development and Clinical Trials Partnerships: A Randomized, Placebo-Controlled, Double-Blind Assessment of Safety, Tolerability, and Immunogenicity of PfSPZ Vaccine for Malaria in Healthy Equatoguinean Men*. *The American journal of tropical medicine and hygiene* 2018, 98:308–318.

47. Jongo SA, Church LWP, Nchama VUNN, Hamad A, Chuquiyauri R, Kassim KR, Athuman T, Deal A, KC N, Mtoro A, et al: *Multi-Dose Priming Regimens of PfSPZ Vaccine: Safety and Efficacy Against Controlled Human Malaria Infection in Equatoguinean Adults*. *American Journal of Hygiene and Tropical Medicine* 2021.

48. Schindler T, Robaina T, Sax J, Bieri JR, Mpina M, Gondwe L, Acuche L, Garcia G, Cortes C, Maas C, Daubenberger C: *Molecular monitoring of the diversity of human pathogenic malaria species in blood donations on Bioko Island, Equatorial Guinea*. *Malaria Journal* 2019, 18:9.

49. Hajian-Tilaki K: *Sample size estimation in diagnostic test studies of biomedical informatics*. *Journal of Biomedical Informatics* 2014, 48:193–204.

50. Mark Stevenson TN, Cord Heuer, Jonathon Marshall, Javier Sanchez, Ron Thornton, Jeno Reiczigel, Jim Robinson-Cox, Paola Sebastiani, Peter Solymos, Kazuki Yoshida, Geoff Jones, Sarah Pirikahu, Simon Firestone, Ryan Kyle, Johann Popp, Mathew Jay, Charles Reynard: epiR: *Tools for the Analysis of Epidemiological Data*. 2020.

51. Mercereau-Puijalon O, Barale J-C, Bischoff E: *Three multigene families in Plasmodium parasites: facts and questions*. *International Journal for Parasitology* 2002, 32:1323–1344.

52. Bejon P, Andrews L, Hunt-Cooke A, Sanderson F, Gilbert SC, Hill AVS: *Thick blood film examination for Plasmodium falciparum malaria has reduced sensitivity and underestimates parasite density*. *Malaria Journal* 2006, 5:104.

53. Hawkins K, Burton R, LaBarre P: *Diagnostics to support malaria elimination: Choosing an appropriate biomarker to target the subclinical Plasmodium falciparum transmission reservoir*. In *IEEE Global Humanitarian Technology Conference (GHTC 2014)*; 10-13 Oct. 2014 2014: 561-568.

54. Mosha JF, Sturrock HJW, Greenhouse B, Greenwood B, Sutherland CJ, Gadalla N, Atwal S, Drakeley C, Kibiki G, Bousema T, et al: *Epidemiology of subpatent Plasmodium falciparum infection: implications for detection of hotspots with imperfect diagnostics*. *Malaria Journal* 2013, 12:221.

55. Okell LC, Griffin JT, Kleinschmidt I, Hollingsworth TD, Churcher TS, White MJ, Bousema T, Drakeley CJ, Ghani AC: *The Potential Contribution of Mass Treatment to the Control of Plasmodium falciparum Malaria*. *PLOS ONE* 2011, 6:e20179.
56. Jongco SA, Shekalaghe SA, Church LWP, Ruben AJ, Schindler T, Zenklusen I, Rutishauser T, Rothen J, Tumbo A, Mkindi C, et al: Safety, Immunogenicity, and Protective Efficacy against Controlled Human Malaria Infection of Plasmodium falciparum Sporozoite Vaccine in Tanzanian Adults. Am J Trop Med Hyg 2018, 99:338–349.

57. Sulyok Z, Fendel R, Eder B, Lorenz F-R, Kc N, Kamahl M, Lalamruata A, Nguyen TT, Held J, Adjadi FAC, et al: Heterologous protection against malaria by a simple chemoattenuated PfSPZ vaccine regimen in a randomized trial. Nature Communications 2021, 12:2518.

58. Sulyok M, Rückle T, Roth A, Mürbeth RE, Chalon S, Kerr N, Samec SS, Gobeau N, Calle CL, Ibáñez J, et al: DSM265 for Plasmodium falciparum chemoprophylaxis: a randomised, double blinded, phase 1 trial with controlled human malaria infection. The Lancet Infectious Diseases 2017, 17:636–644.

59. Murphy SC, Duke ER, Shipman KJ, Jensen RL, Fong Y, Ferguson S, Janes HE, Gillespie K, Seilie AM, Hanron AE, et al: A Randomized Trial Evaluating the Prophylactic Activity of DSM265 Against Preerythrocytic Plasmodium falciparum Infection During Controlled Human Malarial Infection by Mosquito Bites and Direct Venous Inoculation. The Journal of Infectious Diseases 2017, 217:693–702.

60. Mpina M, Maurice NJ, Yajima M, Slichter CK, Miller HW, Dutta M, McElrath MJ, Stuart KD, De Rosa SC, McNevin JP, et al: Controlled Human Malaria Infection Leads to Long-Lasting Changes in Innate and Innate-like Lymphocyte Populations. 2017, 199:107–118.

61. Kapulu MC, Njuguna P, Hamaluba M, Kimani D, Ngoi JM, Musembi J, Ngoto O, Otieno E, Billingsley PF, Team C-SS: Naturally acquired immunity among Kenyan adults suppresses the West African P. falciparum NF54 strain in controlled human malaria infection (CHMI). 2020:2020.2008.2011.20172411.

62. Lell B, Mordmüller B, Dejon Agobe J-C, Honkpehedji J, Zinsou J, Mengue JB, Loembe MM, Adegnika AA, Held J, Lalamruata A, et al: Impact of Sickle Cell Trait and Naturally Acquired Immunity on Uncomplicated Malaria after Controlled Human Malaria Infection in Adults in Gabon. The American journal of tropical medicine and hygiene 2018, 98:508–515.

63. Achan J, Reuling IJ, Yap XZ, Dabira E, Ahmad A, Cox M, Nwakanma D, Tetteh K, Wu L, Bastiaens GJH, et al: Serologic Markers of Previous Malaria Exposure and Functional Antibodies Inhibiting Parasite Growth Are Associated With Parasite Kinetics Following a Plasmodium falciparum Controlled Human Infection. Clinical Infectious Diseases 2019, 70:2544–2552.

64. Kapulu MC, Njuguna P, Hamaluba M, Kimani D, Ngoi JM, Musembi J, Ngoto O, Otieno E, Billingsley PF, Controlled Human Malaria Infection in Semi-Immune Kenyan Adults Study T: Safety and PCR monitoring in 161 semi-immune Kenyan adults following controlled human malaria infection. JCI insight 2021, 6:e146443.

65. Roestenberg M, Bijker EM, Sim BKL, Billingsley PF, James ER, Bastiaens GJH, Teirlinck AC, Scholzen A, Teelen K, Arens T, et al: Controlled human malaria infections by intradermal injection of cryopreserved Plasmodium falciparum sporozoites. The American journal of tropical medicine and hygiene 2013, 88:5–13.
66. Laurens MB, Berry AA, Travassos MA, Strauss K, Adams M, Shrestha B, Li T, Eappen A, Manoj A, Abebe Y, et al: Dose-Dependent Infectivity of Aseptic, Purified, Cryopreserved Plasmodium falciparum 7G8 Sporozoites in Malaria-Naive Adults. *The Journal of Infectious Diseases* 2019, 220:1962–1966.

67. Richie TL, Billingsley PF, Sim BK, James ER, Chakravarty S, Epstein JE, Lyke KE, Mordmuller B, Alonso P, Duffy PE, et al: Progress with Plasmodium falciparum sporozoite (PfSPZ)-based malaria vaccines. *Vaccine* 2015, 33:7452–7461.

68. Kamau E, Alemayehu S, Feghali KC, Saunders D, Ockenhouse CF: Multiplex qPCR for Detection and Absolute Quantification of Malaria. *PLOS ONE* 2013, 8:e71539.

69. Koepfl C, Nguitragool W, Hofmann NE, Robinson LJ, Ome-Kaius M, Sattabongkot J, Felger I, Mueller I: Sensitive and accurate quantification of human malaria parasites using droplet digital PCR (ddPCR). *Scientific Reports* 2016, 6:39183.

70. Hodgson SH, Douglas AD, Edwards NJ, Kimani D, Elias SC, Chang M, Daza G, Seilie AM, Magiri C, Muia A, et al: Increased sample volume and use of quantitative reverse-transcription PCR can improve prediction of liver-to-blood inoculum size in controlled human malaria infection studies. *Malaria Journal* 2015, 14:33.

71. Walk J, Schats R, Langenberg MCC, Reuling IJ, Teelen K, Roestenberg M, Hermsen CC, Visser LG, Sauerwein RW: Diagnosis and treatment based on quantitative PCR after controlled human malaria infection. *Malaria Journal* 2016, 15:398.

72. Hodgson SH, Juma E, Salim A, Magiri C, Kimani D, Njenga D, Muia A, Cole AO, Ogwang C, Awuondo K, et al: Evaluating controlled human malaria infection in Kenyan adults with varying degrees of prior exposure to Plasmodium falciparum using sporozoites administered by intramuscular injection. 2014, 5.

73. Rothen J, Murie C, Carnes J, Anupama A, Abdulla S, Chemb A, Mpina M, Tanner M, Lee Sim BK, Hoffme SL, et al: Whole blood transcriptome changes following controlled human malaria infection in malaria pre-exposed volunteers correlate with parasite prepatent period. *PLOS ONE* 2018, 13:e0199392.

74. Friedman-Klabanoff DJ, Laurens MB, Berry AA, Travassos MA, Adams M, Strauss KA, Shrestha B, Levine MM, Edelman R, Lyke KE: The Controlled Human Malaria Infection Experience at the University of Maryland. *Am J Trop Med Hyg* 2019, 100:556–565.

75. Verma AK, Bharti PK, Das A: HRP-2 deletion: a hole in the ship of malaria elimination. *The Lancet Infectious Diseases* 2018, 18:826–827.

76. Färnert A, Arez AP, Correia AT, Björkman A, Snounou G, do Rosário V: Sampling and storage of blood and the detection of malaria parasites by polymerase chain reaction. *Transactions of the Royal Society of Tropical Medicine and Hygiene* 1999, 93:50–53.

77. Poti KE, Sullivan DJ, Dondorp AM, Woodrow CJ: HRP2: Transforming Malaria Diagnosis, but with Caveats. *Trends in Parasitology* 2020, 36:112–126.

78. Rogier E, Plucksinski M, Lucchi N, Mace K, Chang M, Lemone JF, Candrinho B, Colborn J, Dimbu R, Fortes F, et al: Bead-based immunoassay allows sub-picogram detection of histidine-rich protein 2
from Plasmodium falciparum and estimates reliability of malaria rapid diagnostic tests. *PLOS ONE* 2017, 12:e0172139.

**Figures**

![Venn diagram](image)

**Figure 1**

Similarities and discrepancies in detection of *P. falciparum* cases by different diagnostic methods Venn-diagram showing distribution of positive results according to the diagnostic test used and qPCR as a reference method. All samples were assorted by thick blood smear (TBS), ultrasensitive rapid diagnostic test (uRDT), and conventional rapid diagnostic test (cRDT). All cases were low parasite density Pf infections that occurred during CHMI.
Figure 2

Distribution and Geomean of parasite densities detected by TBS, uRDT and cRDT. Distribution of parasite density by qPCR of all malaria positive samples by TBS, uRDT, and cRDT. Parasite density was measured by qPCR which served as the reference method. Red dots represents the geomean, and error bars represent the geometric 95% confidence intervals of each respective diagnostic group. Significance values were calculated using two-tailed Wilcoxon-test.
Figure 3

Time to first Plasmodium falciparum asexual blood stage parasites according to diagnostic method Kaplan-Meier plot of the number of infections detected by time to first detection (days) for qPCR, TBS, uRDT and cRDT. All cases were low parasite density Pf infections that occurred during CHMI. P-value < 0.001 calculated using the log-rank test.

Supplementary Files

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