Open access **Protocol**

BMJ Open Machine learning of blood haemoglobin and haematocrit levels via smartphone conjunctiva photography in Kenyan pregnant women: a clinical study protocol

Haripriya Sakthivel,^{1,2} Sang Mok Park,¹ Semin Kwon,¹ Eunice Kaguiri,^{3,4} Elizabeth Nyaranga,^{3,4} Jung Woo Leem,¹ Shaun G Hong,¹ Peter J Lane,⁵ Edwin O Were,^{3,4} Martin C Were,^{5,6,7} Young L Kim ¹ 1,8,9

To cite: Sakthivel H, Park SM, Kwon S, et al. Machine learning of blood haemoglobin and haematocrit levels via smartphone conjunctiva photography in Kenyan pregnant women: a clinical study protocol. BMJ Open 2025;15:e097342. doi:10.1136/ bmjopen-2024-097342

Prepublication history for this paper is available online. To view these files, please visit the journal online (https://doi. org/10.1136/bmjopen-2024-097342).

Received 30 November 2024 Accepted 23 April 2025



@ Author(s) (or their employer(s)) 2025. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ Group.

For numbered affiliations see end of article.

Correspondence to

Dr Edwin O Were: eowere@mu.ac.ke. Dr Martin C Were: martin.c.were@vumc.org and Dr Young L Kim; youngkim@purdue.edu

ABSTRACT

Introduction Anaemia during pregnancy is a widespread health burden globally, especially in low- and middleincome countries, posing a serious risk to both maternal and neonatal health. The primary challenge is that anaemia is frequently undetected or is detected too late, worsening pregnancy complications. The gold standard for diagnosing anaemia is a clinical laboratory blood haemoglobin (Hgb) or haematocrit (Hct) test involving a venous blood draw. However, this approach presents several challenges in resource-limited settings regarding accessibility and feasibility. Although non-invasive blood Hgb testing technologies are gaining attention, they remain limited in availability, affordability and practicality. This study aims to develop and validate a mobile health (mHealth) machine learning model to reliably predict blood Hgb and Hct levels in Black African pregnant women using smartphone photos of the conjunctiva.

Methods and analysis This is a single-centre, crosssectional and observational study, leveraging existing antenatal care services for pregnant women aged 15 to 49 years in Kenya. The study involves collecting smartphone photos of the conjunctiva alongside conventional blood Hgb tests. Relevant clinical data related to each participant's anaemia status will also be collected. The photo acquisition protocol will incorporate diverse scenarios to reflect real-world variability. A clinical training dataset will be used to refine a machine learning model designed to predict blood Hgb and Hct levels from smartphone images of the conjunctiva. Using a separate testing dataset, comprehensive analyses will assess its performance by comparing predicted blood Hgb and Hct levels with clinical laboratory and/or finger-prick readings. Ethics and dissemination This study is approved by the Moi University Institutional Research and Ethics Committee (Reference: IREC/585/2023 and Approval Number: 004514), Kenya's National Commission for Science, Technology, and Innovation (NACOSTI Reference: 491921) and Purdue University's Institutional Review Board (Protocol Number: IRB-2023-1235). Participants will include emancipated or mature minors. In Kenya, pregnant women aged 15 to 18 years are recognised

STRENGTHS AND LIMITATIONS OF THIS STUDY

- ⇒ Unmodified smartphone cameras and machine learning approaches are used to non-invasively predict blood haemoglobin (Hgb) and haematocrit (Hct) levels from an easily accessible site-the conjunctiva.
- ⇒ Development and validation of the model are tailored to predict blood Hab and Hct levels in a quantitative manner similar to clinical laboratory testing, rather than detecting anaemia as a binary outcome.
- ⇒ Study population is specifically designed to address healthcare disparities impacting Black African preg-
- ⇒ Target gestation includes all three trimesters with approximately equal representation from each trimester.
- ⇒ Due to the observational nature of the study, there is no intervention administered.

as emancipated or mature minors, allowing them to provide informed consent independently. The study poses minimal risk to participants. Findings and results will be disseminated through submissions to peer-reviewed journals and presentations at the participating institutions. including Moi Teaching and Referral Hospital and Kenya's Ministry of Health. On completion of data collection and modelling, this study will demonstrate how machine learning-driven mHealth technologies can reduce reliance on clinical laboratories and complex equipment, offering accessible and scalable solutions for resource-limited and at-home settings.

INTRODUCTION

The prevalence of anaemia remains high, affecting nearly one-quarter of the global population (1.92 billion) in 2021. 1-3 It is especially predominant among women of reproductive age in low- and middle-income countries, impacting 45% of pregnant and





40% of non-pregnant women. ⁴⁵ In East Africa, it is estimated that 42% of pregnant women are anaemic. ⁶ In Kenya, cases among pregnant women surged from $55\,539$ in 2016 to $295\,642$ in 2019. At the country's largest maternity unit, 57% of women in their second and third trimesters were affected by anaemia. ⁸ Even in the USA, more than 40% of females aged 12 to 21 years are estimated to have iron deficiency or iron-deficiency anaemia. ⁹

Anaemia is a major contributor to maternal and neonatal mortality. Moderate to severe anaemia exacerbates critical conditions such as haemorrhage and sepsis during pregnancy. Anaemia-associated pregnancy complications include preterm labour, low birth weight, stillbirth and neonatal mortality, all of which increase the risk of adverse outcomes for both mothers and newborns. Maternal anaemia during pregnancy can have long-term consequences on a child's neurocognitive development. Importantly, interventions are available to address anaemia even in resource-limited settings, including dietary modifications with iron-rich foods, supplementation with iron, folic acid, vitamin B_{12}^{13-16} and blood transfusion in cases of severe anaemia. The

Anaemia management during pregnancy relies on the ability to quantitatively assess blood haemoglobin (Hgb) and haematocrit (Hct) levels in a timely manner. The main challenge in resource-limited settings is that anaemia during pregnancy is often not detected or is detected too late. The World Health Organization (WHO) recommends at least one blood Hgb test per trimester. Unfortunately, women in these settings often lack access to recommended diagnostic testing. For instance, in Kenya, only 17% of women had access to minimally adequate delivery care with routine antenatal tests. ¹⁸ Other countries in sub-Saharan Africa and South Asia face similar challenges. However, there are only a limited number of studies using non-invasive or point-of-care (POC) blood Hgb tests specifically for pregnant women in general. ^{19–22}

The gold standard for diagnosing anaemia is a clinical laboratory blood Hgb test to measure Hgb content in the blood (grams per decilitre). ^{23–26} However, venous blood draw-based Hgb tests have several limitations, including the need for specialised equipment (haematology analyser), pain, discomfort, risk of haematoma, infection and iatrogenic blood loss. ²⁷ Non-invasive and cost-effective blood Hgb testing technologies remain limited. ^{28–32} For example, Masimo and OrSense devices require expensive specialised equipment available only in advanced hospital settings. ^{33–35} Alternatively, POC blood analysers that use capillary blood sampling (finger-prick testing) (eg, Abbott i-STAT, HemoCue and VERI-Q) are commercially available, but require environmentally sensitive cartridges with short shelf lives. ^{36 37}

Non-invasive POC blood Hgb assessment technologies have received considerable attention, 38 including HemaApp, 39 fingernail mobile app, 40-42 fingertip devices, 43 44 lip mucosal imaging 45 and palpebral conjunctiva smartphone imaging. 46-59 Specifically, the palpebral conjunctiva, a common site for assessing paleness

and anaemia, offers the advantages of easy, non-contact imaging without surface pressure. ⁶⁰ Its underlying microvasculature is unaffected by skin pigmentation (eg, melanocytes), removing the need for personal calibration. ⁶¹ In addition, the conjunctiva may not be easily recognisable, providing enhanced privacy protection. ⁶² ⁶³

Objectives and hypothesis

The primary objective of this study is to develop and validate a mobile health (mHealth) computational model using machine learning to accurately and precisely predict blood Hgb and Hct levels in Black African pregnant women using photos of the conjunctiva acquired by a smartphone camera. The central hypothesis is that blood Hgb levels can be reliably predicted from redgreen-blue (RGB) images of the conjunctiva in a noninvasive manner with performance comparable to POC finger-prick testing. First, we will capture high-quality conjunctiva photos under diverse photo acquisition settings from pregnant women across all three trimesters, encompassing a broad range of Hgb and Hct levels. Second, we will refine the mHealth prediction model and compare the predictions with conventional blood Hgb and Hct testing methods. Given the physiological changes during pregnancy that vary by trimester, this study emphasises acquiring data from all stages to ensure reliable predictions.

METHODS: PARTICIPANTS, STUDY PROCEDURES AND OUTCOMES

Study design

This is a single-centre, non-interventional, cross-sectional and observational study involving the acquisition of photos of the conjunctiva alongside conventional blood Hgb and Hct tests. Relevant clinical data related to the participant's anaemia status will also be collected. The blood Hgb and Hct values computed from the mHealth prediction model will not be used to guide interventions or diagnostics. All data collection will take place during a single study visit. Thus, a retention plan is not required.

Setting and recruitment

Figure 1 outlines the setting, enrolment and data collection. The primary clinical setting is the Maternal Child Health (MCH) clinic at Moi Teaching and Referral Hospital (MTRH) in Eldoret, Kenya, in collaboration with the Academic Model Providing Access to Healthcare (AMPATH). MTRH is the second-largest referral hospital in Kenya. The MCH clinic at MTRH has 20 obstetricians and over 40 residents who care for approximately 900 pregnant women per month. AMPATH also provides a framework for sustainable research and scalable healthcare access. AMPATH is a partnership between the Moi University School of Medicine, MTRH and a consortium of US institutions.

Study participants

Our study will recruit volunteer pregnant women receiving antenatal care at the MCH clinic, targeting

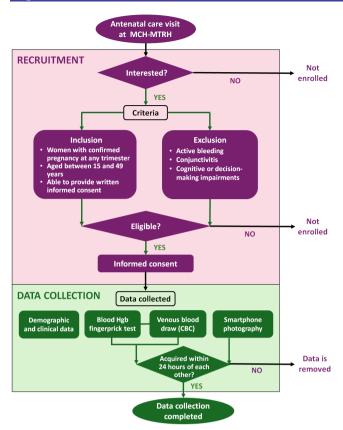


Figure 1 Flowchart of recruitment, enrolment and data collection. This single-centre, cross-sectional and observational study leverages existing antenatal care services for pregnant women aged 15 to 49 years at the Maternal Child Health (MCH) clinic at Moi Teaching and Referral Hospital (MTRH) in Eldoret, Kenya. CBC, complete blood count; Hgb, haemoglobin.

600 participants, with approximately 200 women per trimester, aged 15 to 49 years. Because the mHealth prediction model for blood Hgb computation relies on machine learning, conventional statistical methods are not directly applicable for estimating power and sample size. However, our estimates are conservative based on the previous study at MTRH. 49 For 200 participants per trimester, the 95% confidence intervals (CIs) for the correlation coefficient between the mHealth and clinical laboratory blood Hgb levels are expected to range from 0.09 to 0.13, assuming a correlation coefficient of 0.85. Similarly, the 95% CI for the intraclass correlation coefficient (ICC) will range from 0.07 to 0.13, assuming an expected ICC of 0.85. To mitigate the risk of overfitting, a separate masked testing dataset comprising 30% of the total data will be used. This testing dataset will be independent of the training dataset, which consists of the remaining 70% of the data.

Inclusion and exclusion criteria

The study inclusion criteria (figure 1) are as follows:

- 1. Women with confirmed pregnancy at any gestational stage (first, second or third trimester).
- 2. Aged 15 to 49 years.

3. Able to provide written informed consent.

Participants will be excluded if they have hypotension, active or ongoing bleeding, conjunctivitis (or visible conjunctival inflammation), trauma or infection affecting the eyes or eyelids, or if laboratory blood Hgb and Hct testing may be delayed beyond 24 hours after photography.

Overall procedure

If the patient agrees to participate in this study, study personnel will provide simple instructions on how to gently pull down the inner eyelid using the participant's index finger. Then, the study personnel will hold a colour reference chart on the patient's forehead and capture photos of both the left and right eyes using three different smartphone models. The total time required for imaging is approximately 5 minutes. Clinical data will also be collected, including laboratory Hgb and Hct values from blood samples drawn within 24 hours before or after the conjunctiva photo timestamp. The study personnel will complete a clinical data collection sheet and attach the results of the clinical laboratory test. All photos and associated data will be submitted through a customised data collection application (app).

METHODS: DATA COLLECTION, MANAGEMENT AND ANALYSIS Timepoints for data collection

All data collection for the study will take place during a single visit, lasting approximately 10 minutes, with no follow-up required. During the visit, consented participants' baseline demographic and clinical data will be recorded on a study form. Smartphone photography and venous blood draw and/or finger-prick testing will then be performed. To ensure data reliability, smartphone photography must occur within 24 hours of the blood draw. Once data collection is complete, all information will be uploaded to a customised data collection app linked to a Health Insurance Portability and Accountability Act (HIPAA)-compliant server, where it will be organised and processed for analysis.

Demographic and clinical data

Demographic data collected from participants will include date of birth, marital status and highest level of education completed. Clinical data will cover details of the current pregnancy, obstetric history, medical and surgical history (including blood pressure), family history and antenatal profile.

Clinical laboratory blood Hgb test and/or finger-prick test

We will assess the results of complete blood count (CBC) tests, specifically measuring clinical laboratory-based blood Hgb and Hct levels using the Beckman Coulter AcT 5diff or a similar device from samples collected on the same day. CBC tests will be conducted at a clinical reference laboratory certified by the College of American Pathologists' External Quality Assurance Program.



In addition, capillary blood sampling using VERI-Q will be performed either before or after photographing the conjunctiva.

Smartphone conjunctiva photographing

Photos acquired with a digital (or smartphone) camera exhibit different colours depending on smartphone models, image formats and light conditions.^{64°65} To develop an mHealth prediction model that is accurate under diverse data acquisition conditions, the photo acquisition protocol will incorporate a custom-made colour reference chart,⁵⁴ different smartphone models (Google Pixel 5, Samsung Galaxy A52 and Samsung Galaxy S21) and file formats. The colour reference chart, roughly the size of a business card, is designed to support colour recovery with reduced dependence on photo acquisition settings by being physically captured within each photo. Instead of commercially available colour reference charts (eg, Macbeth ColorChecker, ColorChecker Classic Mini), we will use a custom-designed colour chart that can be mass-printed with a standard inkjet printer. Due to sanitation requirements and participant tracking, disposable colour charts are necessary. However, the high cost of commercially available options makes them impractical for single-use applications.

For Samsung Galaxy S21, both DNG (also known as RAW) and JPEG formats will be generated using Pro Mode. Google Pixel 5 and Samsung Galaxy A52 will use a third-party app (Adobe Lightroom or Halide Mark) to capture photos in DNG format. As a key specification of smartphone cameras, we evaluated the spatial resolution of the three different smartphone models using the edge

method in a laboratory setting. The measurements were conducted at a typical distance of 100–150 mm between the camera and the participant's eye. Google Pixel 5 has a spatial resolution of 137 μ m, while Samsung Galaxy S21 has a spatial resolution of 172 μ m. Despite being a lower-end smartphone, the Samsung Galaxy A52 has a spatial resolution of 108 μ m.

Data collection mobile app

We developed a mobile app for Android to facilitate the collection and transfer of photos (figure 2). This app is specifically designed to ensure proper use of the colour reference chart during photo acquisition. It allows study personnel to upload photos taken with the smartphone camera, requiring them to complete form fields before selecting photos from the smartphone gallery for upload. All data, including the photos collected, are stored on a HIPAA-compliant cloud server and can be securely accessed through a high-security portal. Importantly, the data collection app is designed to efficiently handle large photo files in the DNG (RAW) format. The DNG format reduces non-linear rendering and image compression.⁶⁵ With a 10-bit colour depth in each RGB channel, DNG allows for 2^{10×3} combinations of RGB values. In contrast, JPEG with an 8-bit colour depth $(2^{8\times3}\approx16.77 \text{ million})$ colours) involves non-linear filtering and image compression. As a result, DNG photos are substantially larger than JPEG photos. Recent smartphone models support direct access to the DNG format either in the default camera settings or through third-party applications (eg, Adobe Lightroom or Halide Mark). The app also enables photo uploads even when network connectivity is interrupted; it

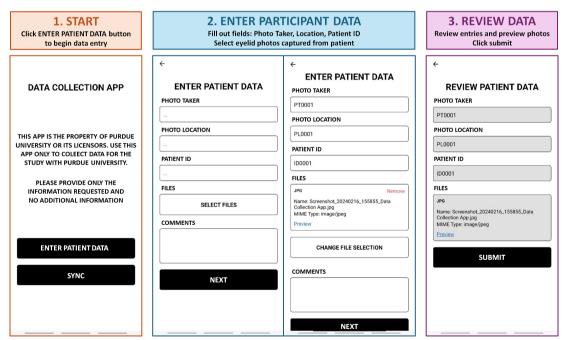


Figure 2 Representative screenshots of the customised data collection mobile app for Android smartphones. The app's flow and processes are designed to efficiently manage large photo files. The user interface enables study personnel to upload photos to a Health Insurance Portability and Accountability Act (HIPAA)-compliant cloud server, which can be securely accessed through a high-security portal.



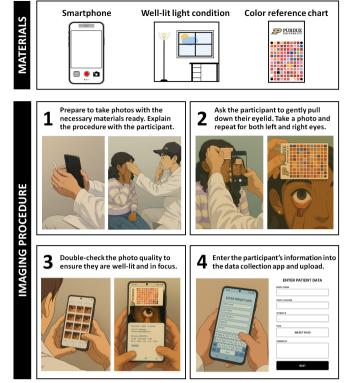


Figure 3 Photo acquisition instructions and procedure. Preparation materials and instructional examples guide study personnel in quickly capturing high-quality photos of the participant's conjunctiva. The original demonstration photos, featuring the authors performing the procedure, are further rendered using ChatGPT.

automatically uploads photos from a temporary folder in the background, one photo at a time, to reduce the data payload. Once all photos are uploaded, the temporary folder is deleted from the device.

Smartphone photographing procedure

Figure 3 summarises the photo acquisition protocol.

- 1. Direct the participant to sit facing the ceiling light source. Adjust the room brightness if necessary to ensure clear photos without shadows or glares.
- 2. Write the participant identifier (ID) and date on the colour reference chart to distinguish photos.
- 3. Ask the participant to remove glasses or any objects that may obstruct the forehead.
- 4. Rehearse pulling down the inner eyelid with the participant to ensure adequate and accurate exposure.
- 5. Hold the chart against the participant's forehead with one hand. Instruct the participant to use their fingertips to pull down the inner eyelid.
- 6. Ensure the colour reference chart is horizontally aligned with the participant's eye and visible in the camera view.
- 7. While holding the chart with one hand, use the other hand to operate the smartphone.
 - a. Ask the participant to look up at the ceiling while exposing the conjunctiva.

- b. Include both the entire colour reference chart and the conjunctiva within the frame.
- c. The colour reference chart must remain horizontal.
- d. Avoid covering the chart with fingers or casting shadows on it.
- e. Keep the chart flat without bending.
- Ensure consistent lighting; the colour reference chart and conjunctiva should have similar brightness.
- 8. Use the smartphones in this sequence: Samsung Galaxy A52, Google Pixel 5 and Samsung Galaxy S21.
 - a. Capture four photos of the left conjunctiva with each smartphone.
 - b. Capture four photos of the right conjunctiva with each smartphone.
- 9. Input the participant's information and upload the photos to the data collection app (figure 2).

Model refinement and optimisation

The mHealth prediction model will be refined and optimised for the target population, as it has not yet been tailored to this group. The current version of the mHealth model comprises four submodules 47 49 52-54 67-69:

- 1. Colour correction: extracts absolute colour values of the conjunctiva, ensuring consistency across different smartphone models and light conditions.⁶⁸
- 2. Automated segmentation: automatically identifies and delineates the conjunctival region of interest. ⁶⁷ ⁶⁹
- 3. Hyperspectral learning (also referred to as spectral reconstruction, spectral super-resolution or spectral reflectance estimation): reconstructs high-resolution spectral data from RGB values of photos captured by smartphone cameras. 49 53
- 4. Blood Hgb and Hct content computation: estimates blood Hgb and Hct levels using the reconstructed hyperspectral data. 47 49

To mitigate the risk of overfitting in the blood Hgb and Hct content computation, photo data will be divided into training (70% of participants) and testing (30%) datasets based on participant IDs. The photos from the same participants will be assigned exclusively to either the training or testing datasets to prevent data leakage. Cross-validation will be conducted to evaluate the model's performance across different subsets of the data. It should be noted that colour correction, automated segmentation and hyperspectral learning are not subjected to training, as these processes were already completed using separate data from our previous studies. 47 49 52-54 67-69 This compound machine learning model integrates domain knowledge (eg, tissue optics and computer vision) into the learning process. Notably, this is designed to mitigate the constraints of relatively limited data. It allows the model to be trained effectively with a limited clinical dataset, addressing the limitations of purely data-driven methods. 70 71



Performance evaluation

To assess the performance of the mHealth model compared with clinical laboratory blood Hgb, Hct and/or finger-prick blood Hgb values, we will perform the following analyses using a testing dataset or cross-validation methods.

- 1. Linear correlation analysis: Quantifies the strength of the relationship between mHealth and clinical laboratory blood Hgb and Hct values.
- 2. Bland–Altman analysis: Uses multiple measurement pairs to evaluate whether mHealth blood Hgb and Hct values align reliably with clinical laboratory results, returning bias and 95% limits of agreement.
- 3. ICC analysis: Assesses the reliability of mHealth blood Hgb and Hct values, focusing on reproducibility—the ability of different users to obtain consistent results. Given that multiple smartphones will be used to capture photos from the same participant, we will emphasise inter-reliability (reproducibility), which measures variation across different users evaluating the same group of participants.
- 4. Paired t-tests: Determines whether blood Hgb and Hct values obtained from the left and right conjunctivae are statistically identical.

In addition, we will follow the STARD (Standards for Reporting of Diagnostic Accuracy Studies) guideline^{72 73} for assessing the diagnostic performance of our mHealth prediction model as well as the TRIPOD+AI (Transparent Reporting of a multivariable prediction model for Individual Prognosis Or Diagnosis + Artificial Intelligence) guideline⁷⁴ for reporting our machine learning-based prediction study.

ETHICS AND DISSEMINATION Ethics approval and consent

This study is approved by the Moi University Institutional Research and Ethics Committee (Reference: IREC/585/2023 and Approval Number: 004514), Kenya's National Commission for Science, Technology, and Innovation (NACOSTI Reference: 491921) and Purdue University's Institutional Review Board (Protocol Number: IRB-2023-1235). Our study involves recruiting participants from vulnerable populations, specifically pregnant women, including some who are emancipated or mature minors. In Kenya, pregnant women aged 15 to 18 years are considered emancipated or mature minors, allowing them to provide informed consent independently, without parental involvement. The informed consent form is available in both English and Swahili, the native and widely spoken language in Kenya. Study personnel responsible for communicating with participants are fluent in both languages to ensure clear and effective communication.

Confidentiality, data storage and security

All study data will be stored and accessed in compliance with HIPAA and the Kenya Data Protection Act, 2019.

Specifically, photos will be labelled with the participant ID, smartphone model and left/right. Demographic and clinical information recorded on paper forms by site personnel will be scanned using the smartphone. Data will be uploaded via a custom data collection app developed for this study. This app transmits data to a secure Amazon Web Services server, which is HIPAA-compliant. Access to the server is restricted to study investigators and authorised personnel. Computer records will be stored on password-protected systems, and paper records will be secured in locked cabinets accessible only to authorised study personnel.

Dissemination

We will disseminate results through publications in peer-reviewed journals and presentations at the participating institutions, including Moi Teaching and Referral Hospital, and Kenya's Ministry of Health. This study primarily focuses on developing a machine learning model for blood Hgb and Hct assessments. Our next steps are to scale the project towards developing a minimally viable product—a functional mobile app for bloodless, quantitative blood Hgb assessment—for larger clinical trials. Building on further collaboration with healthcare philanthropy organisations, we plan to evaluate the effectiveness and implementation of the mHealth prediction model through pilot studies and real-world applications.

Patient and public involvement

Patients or the public were not involved in the design, conduct, reporting, or dissemination plans of this research.

DISCUSSION

Digital health technologies have experienced rapid growth and are now widely adopted across various clinical settings. In particular, photos captured with mobile devices (eg, smartphones and tablets) have emerged as pivotal tools in digital health applications, including telemedicine and mHealth. 75–78 Clinical photos are instrumental in healthcare diagnostics, monitoring and management, especially in at-home healthcare and resource-limited settings where traditional equipment may be scarce. Consequently, healthcare professionals increasingly regard smartphones and tablets as indispensable components of modern healthcare practice. However, guidelines on conducting clinical studies using high-quality clinical photos from mobile devices are often not available.

This protocol paper outlines a clinical study exploring the use of smartphone cameras as diagnostic tools. Building on prior research demonstrating the diagnostic potential of clinical photography, this study leverages smartphone technology to improve access to high-quality clinical images. Furthermore, advancements in machine learning and artificial intelligence enhance the diagnostic accuracy of photo-based analyses. The protocols



and procedures described here aim to extend the reach of diagnostic imaging in low-resource environments, where traditional diagnostic tools are often inaccessible. These methods may also be applicable to other clinical studies requiring high-quality imaging.

Author affiliations

¹Weldon School of Biomedical Engineering, Purdue University, West Lafayette, Indiana, USA

²The Charles Draper Stark Laboratory, Cambridge, Massachusetts, USA

³Academic Model Providing Access to Healthcare, Eldoret, Kenya

⁴Division of Obstetrics and Gynecology, Moi University College of Health Sciences, Eldoret, Kenya

⁵Vanderbilt Institute for Global Health, Vanderbilt University Medical Center, Nashville, Tennessee, USA

⁶Department of Biomedical Informatics, Vanderbilt University Medical Center, Nashville, Tennessee, USA

⁷Department of Medicine, Vanderbilt University Medical Center, Nashville, Tennessee, USA

⁸Regenstrief Center for Healthcare Engineering, Purdue University, West Lafayette, Indiana, USA

⁹Purdue Institute for Cancer Research, Purdue University, West Lafayette, Indiana, USA

Acknowledgements We sincerely acknowledge the faculty and staff at MTRH and AMPATH for their invaluable support of this study. We also extend our heartfelt gratitude, in advance, to the pregnant women who graciously chose to participate in this research.

Contributors HS, SMP, SK, JWL and YLK developed the data acquisition protocol and instructions and designed the mobile app for data collection. EK and EN provided feedback on the data acquisition process and collected data in the clinical setting. HS, SMP, SK, JWL, SGH and YLK reviewed data quality and provided feedback to the Kenya team. PJL and JWL assisted with study management and logistics. EOW, MCW and YLK obtained ethics approval. EOW led the clinical aspects, including subject enrolment, as the site PI for this study. MCW and YLK revised the protocol and study design. YLK conceptualised the study and provided mentorship and academic supervision. HS generated the figures. HS and YLK wrote the manuscript, with JWL, EOW and MCW providing feedback. YLK is the lead corresponding author and guarantor. For figure 3, the original demonstration photos, featuring the authors performing the procedure, are further rendered using ChatGPT. This is to follow the BMJ guideline on the usage of identifiable photos.

Funding This work was supported by the National Institutes of Health grant R01EB033788, National Institutes of Health grant R21TW012486, National Institutes of Health grant R33TW012486, National Institutes of Health Technology Accelerator Challenge Prize, and Global Health Labs (Gates Ventures). HS is a Draper Scholar funded by The Charles Stark Draper Laboratory, Inc.

Competing interests YLK is a founding member of HemaChrome LLC.

Patient and public involvement Patients and/or the public were not involved in the design or conduct of this research study.

Patient consent for publication Consent obtained directly from patient(s).

Provenance and peer review Not commissioned; externally peer reviewed.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/.

ORCID iD

Young L Kim http://orcid.org/0000-0003-3796-9643

REFERENCES

1 Safiri S, Kolahi A-A, Noori M, et al. Burden of anemia and its underlying causes in 204 countries and territories, 1990-2019: results

- from the Global Burden of Disease Study 2019. *J Hematol Oncol* 2021:14:185.
- 2 Correa-Agudelo E, Kim H-Y, Musuka GN, et al. The epidemiological landscape of anemia in women of reproductive age in sub-Saharan Africa. Sci Rep 2021;11:11955.
- 3 GBD 2021 Anaemia Collaborators. Prevalence, years lived with disability, and trends in anaemia burden by severity and cause, 1990-2021: findings from the Global Burden of Disease Study 2021. *Lancet Haematol* 2023;10:e713–34.
- 4 Kinyoki D, Osgood-Zimmerman AE, Bhattacharjee NV, et al. Anemia prevalence in women of reproductive age in low- and middle-income countries between 2000 and 2018. Nat Med 2021;27:1761–82.
- 5 Alem AZ, Efendi F, McKenna L, et al. Prevalence and factors associated with anemia in women of reproductive age across low- and middle-income countries based on national data. Sci Rep 2023;13:20335.
- 6 Liyew AM, Tesema GA, Alamneh TS, et al. Prevalence and determinants of anemia among pregnant women in East Africa; A multi-level analysis of recent Demographic and Health Surveys. PLoS One 2021;16:e0250560.
- 7 Odhiambo JN, Sartorius B. Mapping of anaemia prevalence among pregnant women in Kenya (2016-2019). BMC Pregnancy Childbirth 2020;20:711.
- 8 Okube OT, Mirie W, Odhiambo E, et al. Prevalence and factors associated with anaemia among pregnant women attending antenatal clinic in the second and third trimesters at Pumwani Maternity Hospital, Kenya. OJOG 2016;06:16–27.
- 9 Weyand AC, Chaitoff A, Freed GL, et al. Prevalence of Iron Deficiency and Iron-Deficiency Anemia in US Females Aged 12-21 Years, 2003-2020. JAMA 2023;329:2191–3.
- 10 Daru J, Zamora J, Fernández-Félix BM, et al. Risk of maternal mortality in women with severe anaemia during pregnancy and post partum: a multilevel analysis. Lancet Glob Health 2018;6:e548–54.
- 11 Ohuma EO, Jabin N, Young MF, et al. Association between maternal haemoglobin concentrations and maternal and neonatal outcomes: the prospective, observational, multinational, INTERBIO-21st fetal study. Lancet Haematol 2023;10:e756–66.
- 12 Sundararajan S, Rabe H. Prevention of iron deficiency anemia in infants and toddlers. *Pediatr Res* 2021;89:63–73.
- 13 Peyrin-Biroulet L, Williet N, Cacoub P. Guidelines on the diagnosis and treatment of iron deficiency across indications: a systematic review. Am J Clin Nutr 2015;102:1585–94.
- 14 De Franceschi L, Iolascon A, Taher A, et al. Clinical management of iron deficiency anemia in adults: Systemic review on advances in diagnosis and treatment. Eur J Intern Med 2017;42:16–23.
- 15 Ning S, Zeller MP. Management of iron deficiency. *Hematology Am Soc Hematol Educ Program* 2019;2019:315–22.
- 16 Raut AK, Hiwale KM. Iron Deficiency Anemia in Pregnancy. Cureus 2022;14:e28918.
- 17 Linder GE, Ipe TS. Pregnancy and postpartum transfusion. *Ann Blood* 2022;7:12.
- 18 Sharma J, Leslie HH, Kundu F, et al. Poor Quality for Poor Women? Inequities in the Quality of Antenatal and Delivery Care in Kenya. PLoS One 2017;12:e0171236.
- 19 Hadar E, Raban O, Bouganim T, et al. Precision and accuracy of noninvasive hemoglobin measurements during pregnancy. J Matern Fetal Neonatal Med 2012;25:2503–6.
- Yoshida A, Saito K, Ishii K, et al. Assessment of noninvasive, percutaneous hemoglobin measurement in pregnant and early postpartum women. Med Devices (Auckl) 2014;7:11–6.
- 21 Ahankari AS, Fogarty AW, Tata LJ, et al. Assessment of a non-invasive haemoglobin sensor NBM 200 among pregnant women in rural India. *BMJ Innov* 2016;2:70–7.
- Mills K, Vermeer JM, Berry WE, et al. Determining the validity of non-invasive spot-check hemoglobin co-oximetry testing to detect anemia in postpartum women at a tertiary care centre, a prospective cohort study. BMC Pregnancy Childbirth 2023;23:479.
- 23 Medicare, Medicaid and CLIA programs; regulations implementing the Clinical Laboratory Improvement Amendments of 1988 (CLIA)--HCFA. Final rule with comment period. Fed Regist 1992;57:7002–186.
- 24 Keng TB, De La Salle B, Bourner G, et al. Standardization of haematology critical results management in adults: an International Council for Standardization in Haematology, ICSH, survey and recommendations. Int J Lab Hematol 2016;38:457–71.
- 25 Pasricha S-R, Colman K, Centeno-Tablante E, et al. Revisiting WHO haemoglobin thresholds to define anaemia in clinical medicine and public health. *Lancet Haematol* 2018;5:e60–2.
- 26 Clinical laboratory improvement amendments of 1988 (Clia) proficiency testing regulations related to analytes and acceptable performance. Fed Regist 2019;84.



- 27 Marn H, Critchley JA. Accuracy of the WHO Haemoglobin Colour Scale for the diagnosis of anaemia in primary health care settings in low-income countries: a systematic review and meta-analysis. Lancet Glob Health 2016;4:e251–65.
- 28 Guo T, Patnaik R, Kuhlmann K, et al. Smartphone dongle for simultaneous measurement of hemoglobin concentration and detection of HIV antibodies. Lab Chip 2015;15:3514–20.
- 29 Smart LR, Ambrose EE, Raphael KC, et al. Simultaneous point-ofcare detection of anemia and sickle cell disease in Tanzania: the RAPID study. Ann Hematol 2018;97:239–46.
- 30 Perez-Plazola MS, Tyburski EA, Smart LR, et al. AnemoCheck-LRS: an optimized, color-based point-of-care test to identify severe anemia in limited-resource settings. BMC Med 2020;18:337.
- 31 An R, Huang Y, Man Y, et al. Emerging point-of-care technologies for anemia detection. Lab Chip 2021;21:1843–65.
- 32 Ahn HS, Lenet T, Gilbert RWD, et al. Accuracy of point-of-care testing devices for haemoglobin in the operating room: metaanalysis. BJS Open 2024;8:zrad148.
- 33 Moore LJ, Wade CE, Vincent L, et al. Evaluation of noninvasive hemoglobin measurements in trauma patients. Am J Surg 2013;206:1041–7.
- 34 Kim S-H, Lilot M, Murphy LS-L, et al. Accuracy of continuous noninvasive hemoglobin monitoring: a systematic review and metaanalysis. Anesth Analg 2014;119:332–46.
- 35 Hiscock R, Kumar D, Simmons SW. Systematic review and metaanalysis of method comparison studies of Masimo pulse cooximeters (Radical-7[™] or Pronto-7[™]) and HemoCue® absorption spectrometers (B-Hemoglobin or 201+) with laboratory haemoglobin estimation. Anaesth Intensive Care 2015;43:341–50.
- 36 Briggs C, Kimber S, Green L. Where are we at with point-of-care testing in haematology? *Br J Haematol* 2012;158:679–90.
- 37 Jaggernath M, Naicker R, Madurai S, et al. Diagnostic Accuracy of the HemoCue Hb 301, STAT-Site MHgb and URIT-12 Pointof-Care Hemoglobin Meters in a Central Laboratory and a Community Based Clinic in Durban, South Africa. PLoS One 2016;11:e0152184.
- 38 Hasan MK, Aziz MH, Zarif MII, et al. Noninvasive Hemoglobin Level Prediction in a Mobile Phone Environment: State of the Art Review and Recommendations. JMIR Mhealth Uhealth 2021;9:e16806.
- 39 Wang EJ, Li W, Zhu J, et al. Noninvasive hemoglobin measurement using unmodified smartphone camera and white flash. 2017 39th Annual International Conference of the IEEE Engineering in Medicine and Biology Society (EMBC); Seogwipo.
- 40 Mannino RG, Myers DR, Tyburski EA, et al. Smartphone app for non-invasive detection of anemia using only patient-sourced photos. Nat Commun 2018;9:4924.
- 41 Young MF, Raines K, Jameel F, et al. Non-invasive hemoglobin measurement devices require refinement to match diagnostic performance with their high level of usability and acceptability. PLoS One 2021;16:e0254629.
- 42 Haggenmüller V, Bogler L, Weber A-C, et al. Smartphone-based point-of-care anemia screening in rural Bihar in India. Commun Med (Lond) 2023;3:38.
- 43 Amrutha AM, Sidenur B, P S B, et al. Estimation of haemoglobin using non-invasive portable device with spectroscopic signal application. Sci Rep 2024;14:8697.
- 44 Sahoo KC, Sinha A, Sahoo RK, et al. Diagnostic Validation and Feasibility of a Non-invasive Haemoglobin Screening Device (EzeCheck) for "Anaemia Mukt Bharat" in India. Cureus 2024;16:e52877.
- 45 Donmez TB, Mansour M, Kutlu M, et al. Anemia detection through non-invasive analysis of lip mucosa images. Front Big Data 2023;6:1241899.
- 46 Kim O, McMurdy J, Jay G, et al. Combined reflectance spectroscopy and stochastic modeling approach for noninvasive hemoglobin determination via palpebral conjunctiva. Physiol Rep 2014;2:e00192.
- 47 Kim T, Choi SH, Lambert-Cheatham N, et al. Toward laboratory blood test-comparable photometric assessments for anemia in veterinary hematology. J Biomed Opt 2016;21:107001.
- 48 Dimauro G, Guarini A, Caivano D, et al. Detecting clinical signs of anaemia from digital images of the palpebral conjunctiva. IEEE Access 2019;7:113488–98.
- 49 Park SM, Visbal-Onufrak MA, Haque MM, et al. mHealth spectroscopy of blood hemoglobin with spectral super-resolution. Optica 2020;7:563–73.
- 50 Ghosal S, Das D, Udutalapally V, et al. sHEMO: Smartphone spectroscopy for blood hemoglobin level monitoring in smart anemia-care. *IEEE Sensors J* 2021;21:8520–9.

- 51 Suner S, Rayner J, Ozturan IU, et al. Prediction of anemia and estimation of hemoglobin concentration using a smartphone camera. PLoS One 2021;16:e0253495.
- 52 Park SM, Ji Y, Kwon S, *et al.* Association of noninvasive peripheral blood hemoglobin assessments with venous blood draws among sickle cell patients. *Blood* 2022;140:7832–3.
- 53 Ji Y, Park SM, Kwon S, et al. mHealth hyperspectral learning for instantaneous spatiospectral imaging of hemodynamics. PNAS Nexus 2023;2:pgad111.
- 54 Park SM, Ji Y, Kwon S, et al. Remote blood hemoglobin monitoring with hyperspectral color truthing for advancing sickle cell care. Blood 2023:142:2277.
- 55 Asare JW, Appiahene P, Donkoh ET, et al. Iron deficiency anemia detection using machine learning models: a comparative study of fingernails, palm and conjunctiva of the eye images. Engineering Reports 2023:5.
- 56 Hu XY, Li YJ, Shu X, et al. A new, feasible, and convenient method based on semantic segmentation and deep learning for hemoglobin monitoring. Front Med (Lausanne) 2023;10:1151996.
- 57 Dimauro G, Griseta ME, Camporeale MG, et al. An intelligent non-invasive system for automated diagnosis of anemia exploiting a novel dataset. Artif Intell Med 2023;136:102477.
- 58 Zhao L, Vidwans A, Bearnot CJ, et al. Prediction of anemia in realtime using a smartphone camera processing conjunctival images. PLoS One 2024;19:e0302883.
- 59 Chen Y, Hu X, Zhu Y, et al. Real-time non-invasive hemoglobin prediction using deep learning-enabled smartphone imaging. BMC Med Inform Decis Mak 2024;24:187.
- 60 Jacquet-Lagrèze M, Magnin M, Allaouchiche B, et al. Is handheld video microscopy really the future of microcirculation monitoring? Crit Care 2023;27:352.
- 61 Lin JY, Fisher DE. Melanocyte biology and skin pigmentation. *Nature New Biol* 2007;445:843–50.
- 62 Nakayama LF, de Matos JCRG, Stewart IU, et al. Retinal Scans and Data Sharing: The Privacy and Scientific Development Equilibrium. Mayo Clin Proc Digit Health 2023;1:67–74.
- 63 Maloney M, Bradley E. Recognition of external eye photos: HIPAA implications. *Invest Ophthalmol Vis Sci* 2013;54.
- 64 Ji Y, Kwak Y, Park SM, et al. Compressive recovery of smartphone RGB spectral sensitivity functions. Opt Express 2021;29:11947–61.
- 65 Delbracio M, Kelly D, Brown MS, et al. Mobile Computational Photography: A Tour. Annu Rev Vis Sci 2021;7:571–604.
- 66 Visbal Onufrak MA, Konger RL, Kim YL. Telecentric suppression of diffuse light in imaging of highly anisotropic scattering media. Opt Lett 2016;41:143–6.
- 67 Hong SG, Park SM, Kwon S, et al. Smartphone conjunctiva photography for malaria risk stratification in asymptomatic school age children. NPJ Digit Med 2025;8:151.
- 68 Park SM, Kwon S, Ji Y, et al. Machine reading and recovery of colors for hemoglobin-related bioassays and bioimaging. *Sci Adv* 2025.
- 69 Hong SG, Park SM, Kwon S, et al. Radiomic identification of anemia features in monochromatic conjunctiva photographs in school-age children. Biophoton Discovery 2025;2:022303.
- children. *Biophoton Discovery* 2025;2:022303.
 Karniadakis GE, Kevrekidis IG, Lu L, *et al.* Physics-informed machine learning. *Nat Rev Phys* 2021;3:422–40.
- 71 von Rueden L, Mayer S, Beckh K. Informed machine learning a taxonomy and survey of integrating prior knowledge into learning systems. *IEEE Trans Knowl Data Eng* 2023;35:614.
- 72 Cohen JF, Korevaar DA, Altman DG, et al. STARD 2015 guidelines for reporting diagnostic accuracy studies: explanation and elaboration. BMJ Open 2016;6:e012799.
- 73 Sounderajah V, Ashrafian H, Golub RM, et al. Developing a reporting guideline for artificial intelligence-centred diagnostic test accuracy studies: the STARD-Al protocol. BMJ Open 2021;11:e047709.
- 74 Collins GS, Moons KGM, Dhiman P, et al. TRIPOD+Al statement: updated guidance for reporting clinical prediction models that use regression or machine learning methods. BMJ 2024;385:e078378.
- 75 Hunt B, Ruiz AJ, Pogue BW. Smartphone-based imaging systems for medical applications: a critical review. J Biomed Opt 2021;26:040902.
- 76 Takahashi M, Koike R, Nagasawa K, et al. Development of telemedicine tools with an emphasis on visual observation. Artif Life Robot 2022;27:38–47.
- 77 Wemyss TA, Nixon-Hill M, Outlaw F, et al. Feasibility of smartphone colorimetry of the face as an anaemia screening tool for infants and young children in Ghana. PLoS One 2023;18:e0281736.
- 78 Zoltie T, Blome-Eberwein S, Forbes S, et al. Medical photography using mobile devices. BMJ 2022;378:e067663.