Sero-prevalence of syphilis and associated factors among pregnant women in Ethiopia: a systematic review and meta-analysis

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Abstract

Background: Syphilis remained a major cause of reproductive morbidity and poor pregnancy outcomes in developing countries. Previously, studies showed inconsistent results and failed to show the actual picture of the diseases in Ethiopia. Thus, the aim of this meta-analysis was, first, to determine the updated pooled prevalence of syphilis among pregnant women in Ethiopia and, second, to assess its associated factors.

Methods: A comprehensive search was made on PubMed, Google scholar, Science Direct, and African Journals Online databases to identify relevant articles. A random effects model was used to estimate pooled syphilis prevalence and odds ratio (OR) with the respective 95% confidence intervals (CIs) using STATA 14 statistical software. I² statistics and Egger’s regression test in conjunction with funnel plot was used to determine heterogeneity and publication bias among included studies respectively.

Result: We identified 13 suitable studies in this analysis. Accordingly, the pooled prevalence of syphilis among pregnant women in Ethiopia was 2.32% (95% CI, 1.68–2.97). Specifically, syphilis prevalence was 2.53% (95% CI, 1.92–3.14%) and 1.90% (95% CI, 0.40–3.40%) as per the treponemal and non-tyreponemal diagnostic test, respectively. On the other hand, regional analysis indicated that 4.06% (95% CI, 2.86–5.26) in Southern Nations Nationalities and Peoples (SNNP), 2.16% (95% CI, 1.57–2.75) in Amhara and 1.46% (95% CI, 0.69–2.23) in Oromia region. Being married (OR, 0.37 (95% CI, 0.12–0.91%)) was less likely to develop syphilis. On the other hand, women with history of multiple sexual partner (OR, 2.98 (95% CI, 1.15–7.70)) and women with history of previous sexually transmitted infection (STI) (OR, 4.88 (95% CI, 1.35–17.62)) have higher risk to develop syphilis. Besides, the pooled syphilis-HIV coinfection was 0.80% (95% CI, 0.60–1.01%).

Conclusion: This study provides evidence of relatively high prevalence of syphilis among pregnant women in Ethiopia. Therefore, it is recommended to further ramping up of current intervention measures to prevent future generations.

Systematic review registration: PROSPERO CRD4202021650

Keywords: Syphilis, Aero-prevalence, Pregnant women, Meta-analysis, Ethiopia

Background
Syphilis is a sexually transmitted infection (STI) caused by the spirochete Treponema pallidum, and it continues to be a main public health problem worldwide [1]. It spreads primarily through sexual contact and vertical transmission and can rarely be spread through blood
transfusion. Syphilis can be successfully controlled by effective public health measures due to the availability of a sound diagnostic test and effective and economical treatment options [2]. However, if syphilis is left untreated, it can lead to devastating fetal outcomes [3].

Pregnant women are sexually active and are at risk of STI, including syphilis [4]. Globally, 36 million people are infected with syphilis, out of which 2 million are pregnant women. More than half of infected women transmit the infection to their babies resulting in adverse pregnancy outcomes including early fetal death, stillbirth, preterm birth, low birth weight, neonatal death, and congenital infection in infants [5].

Syphilis remained a major cause of reproductive morbidity and poor pregnancy outcomes in developing countries [4]. In sub-Saharan Africa, syphilis sero-prevalence ranges from 4 to 15%, and can cause adverse outcomes in 50–80% of pregnancies [6]. Ethiopia is among the top three sub-Saharan countries with the highest numbers of adverse pregnancy outcomes attributed to syphilis [7]. Furthermore, studies have demonstrated that 21% children born from seropositive mothers in Ethiopia developed signs of syphilis. Besides, stillbirth and abortion rates of syphilis diagnosed women were almost double relative to the general population [8].

Previously established systematic review and meta-analysis conducted on the prevalence of syphilis among pregnant women in Ethiopia includes only five studies with smaller sample size and did not report syphilis prevalence based on the diagnostic test modality. Besides, it failed to demonstrate syphilis-HIV co-infection and predictors of syphilis sero-positivity [9]. Thus, in the absence of concrete and inclusive evidence in STI endemic settings including HIV and syphilis, this systematic review and meta-analysis was conducted to determine the updated pooled prevalence of syphilis among pregnant women in Ethiopia. Moreover, syphilis sero-reactivity associated factors and syphilis-HIV co-infection was also determined in this study to guide public health intervention and control measures.

Methods and materials

Reporting and study protocol registration

This review was conducted according to the requirements of the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) statement [10] (Additional file 1). This study was registered in International Prospective Register of Systematic Reviews (PROSPERO) database with protocol number, CRD42020211650.

Search strategy and information sources

An inclusive literature search was made from September 1 to 30, 2020, on PubMed, Google scholar, Science Direct, and African Journals Online databases based on Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) statement [10]. The following key words were used for PubMed database searching: [“Syphilis” or “Treponema pallidum” AND (magnitude OR prevalence OR seroprevalence) AND “pregnant women” OR “pregnant” AND “Ethiopia”]. In addition, literatures and reference lists of relevant articles were also retrieved to find additional studies.

Inclusion and exclusion criteria

All articles fulfilling the following conditions were screened and subsequently assessed for eligibility. Published studies conducted only in Ethiopia and reporting prevalence of syphilis among pregnant women, published in English language up to the end of September 2020. Studies with a clear description of participants’ involved and state the number of participants tested for syphilis, and articles with a clear number and/or prevalence of syphilis cases were also considered. Nevertheless, citations without abstracts and/or full-text, review articles, conference abstracts, editorials, duplicate studies, commentaries, trend analyses, studies which include non-pregnant women and those which do not report syphilis prevalence were excluded from the review. Similarly, given the known synergy between HIV and syphilis [11], studies conducted prior to HIV antiretroviral availability (before January 2005 in Ethiopia) [12] were also excluded as this would affect syphilis prevalence.

Outcome of interest

The primary outcome of this study was the prevalence of syphilis among pregnant women. Moreover, the results have been stratified by diagnostic test (treponemal vs non-treponemal testing) and different geographical regions in Ethiopia. Secondly, we have also determined factors associated with syphilis sero-positivity: marital status (married vs not cohabiting; not cohabiting involves: single, widowed, and divorced), previous STI (present vs absent), and previous history of multiple sexual partner (present vs absent). In addition, the prevalence of syphilis-HIV co-infection was also determined.

Study selection, quality assessment, and data extraction

The title and abstract of studies were screened after removing duplicates. Full-text review was conducted for articles found to be relevant by title and abstract to identify potential articles for inclusion in this meta-analysis. The quality of included studies was evaluated by using Joanna Briggs Institute (JBI) quality assessment checklist for prevalence studies [13]. Based on the JBI checklist, studies with a quality score of 50% and above were considered high quality and involved in the analysis.
Extracted data includes the following descriptive information: author and publication year, study area/region, study period, laboratory methods employed to diagnose syphilis, sample size, and prevalence of syphilis. In addition, prevalence of syphilis by marital status, presence/absence of previous history of multiple sexual partner, and history of previous STI were extracted whenever reported. Two independent reviewers (HG and DG) were involved in study selection, quality assessment, and data extraction. Disagreement between the reviewers was solved by discussion.

Statistical methods and analysis
Data were entered into Microsoft Excel and then exported to STATA 14 statistical software for further analysis. The $I^2$ values of 25, 50, and 75% was considered low, medium, and high heterogeneity, respectively [14]. In pooled prevalence analysis and 95% confidence intervals (CIs), the random effects model (DerSimonian-Laird method) [15] was used. The overall and subgroup analysis of random effects model with 95% CIs were calculated and demonstrated using a forest plot. Nevertheless, for studies with small (near 0) or large (near 1) prevalence, the inverse variance method was not stable and hence, we used Freeman Tukey arcsine methodology to address stabilizing variances as evidenced elsewhere [12, 16, 17]. Syphilis-HIV co-infection and the effect of selected predictor variables including marital status, previous history of multiple sexual partner, and previous history of STI on syphilis prevalence was analyzed using separate categories of meta-analysis.

The existence of publication bias was determined using funnel plot and Egger’s regression test. In Egger’s test, $p < 0.05$ was considered statistically significant [18]. The effect of each study on the overall pooled prevalence was determined by using sensitivity analysis. Sensitivity test eliminates each study step by step in the analysis to indicate the pooled effect sizes and related heterogeneity attributed by each individual study.

Result
Study selection
A comprehensive combined literature search generated a total of 936 possible articles, of which 20 were chosen sensibly for detailed full-text assessment and 13 studies were found to be appropriate for consideration in the meta-synthesis (Fig. 1).

![Fig. 1 PRISMA flow chart for the studies screened, reviewed and included](image-url)
Characteristics of included studies

Among the 13 studies [19–31] included in the meta-analysis, a total of 17,656 pregnant women were screened for syphilis in three different regions of Ethiopia. Regional distribution of studies revealed that eight [19–24, 26, 29] from Amhara, two [25, 28] from Southern Nations Nationalities and Peoples (SNNP) and three [27, 30, 31] from Oromia region. In spite of that, there was no any study from other regions of Ethiopia fulfilling the inclusion criteria. Except a single prospective cohort study [27] which was conducted in Assela, Oromia region; all included records were cross-sectional studies with study participants ranging from 181 [31] to 4346 [30] and conducted from 2005 to 2019. Different diagnostic tests were employed for screening syphilis; six studies [19–22, 25, 31] used treponema pallidum hemagglutination assay (TPHA), three studies [24, 28, 29] used venereal diseases research laboratory (VDRL), two studies [26, 30] employed rapid plasma reagin (RPR) test, and one study [27] used syphilis rapid immunoassay test. As per the established literature, the TPHA, syphilis rapid immunoassay and ICS test constitute the treponemal test while the VDRL and RPR are non-treponemal diagnostic tests [32].

Out of 17,656 pregnant women screened for syphilis, 416 were found positive. Out of 416 seropositive women, 172 were from SNNP, 183 were from Amhara, and 61 were from Oromia (Table 1).

Prevalence of syphilis among pregnant women

The pooled prevalence of syphilis among pregnant women in Ethiopia from the random effects model was 2.32% (95% CI, 1.68–2.97; $I^2 = 98.9$%; Eggers test, $p = 0.14$). Subgroup analysis based on the diagnostic test employed indicated that 2.53% (95% CI, 1.92–3.14%) and 1.90% (95% CI, 0.40–3.40%) syphilis prevalence using treponemal and non-treponemal test, respectively (Fig. 2). As graphically demonstrated in the symmetrical funnel plot, there was no evidence of publication bias within included studies (Fig. 3). Moreover, this was also assured by Egger’s test ($p = 0.14$). In addition, the sensitivity analysis clearly indicated that the influence of individual studies on the summary effect estimate was not significant. Consequently, the pooled effect size estimate of syphilis among pregnant women in Ethiopia was steady and reliable when analyzed by omitting one study at a time (Table 2).

Syphilis prevalence in different regions of Ethiopia

The prevalence of syphilis based on the geographical regions of Ethiopia was 4.06% (95% CI, 2.86–5.26) in SNNP, 2.16% (95% CI, 1.57–2.75) in Amhara, and 1.46% (95% CI, 0.69–2.23) in Oromia region (Table 3).

Pooled prevalence of syphilis-HIV co-infection among pregnant women

The overall pooled prevalence of syphilis-HIV co-infection among pregnant women in this meta-analysis was 0.80% (95% CI, 0.60–1.01; $I^2 = 95.2$%) (Fig. 4).

Table 1 Summary characteristics of studies included in the meta-analysis

| Author and year | Study area/region | Study period | Sample size | Test used | Prevalence of syphilis | Quality assessment |
|-----------------|-------------------|--------------|-------------|-----------|-----------------------|-------------------|
| Mulu et al., 2007 [26] | Gondar/Amhara | March to June, 2005 | 480 | RPR | 1% | High quality |
| Abate Assefa, 2014 [19] | Gondar/Amhara | January 2009 to December 2011 | 2385 | TPHA | 2.9% | High quality |
| Endris et al., 2015 [21] | Gondar/Amhara | February to June 2011 | 385 | TPHA | 2.9% | High quality |
| Melku et al., 2015 [22] | Gondar/Amhara | March to May 2012 | 300 | TPHA | 3.7% | High quality |
| Schonfeld et al., 2017 [27] | Assela/Oromia | May 2014 to September 2015 | 580 | SRIA* | 2.2% | High quality |
| Ageru et al., 2018 [28] | Wolaita Sodo/SNNP | 30 June 2012 to 30, June 2016 | 4022 | VDRL | 3.7% | High quality |
| Amsalu et al., 2018 [25] | Yirgalem/SNNP | October 2015 to August 2016 | 494 | TPHA | 5.1% | High quality |
| Zinabie et al., 2018 [29] | Debre Berhan/Amhara | September 2015 to August 2017 | 385 | VDRL | 1.8% | High quality |
| Asfaw et al., 2019 [31] | Jimma/Oromia | January to June, 2016 | 181 | TPHA | 1.1% | High quality |
| Biadgo et al., 2019 [20] | Gondar/Amhara | January 2011 to April 2015 | 3504 | TPHA | 1.9% | High quality |
| Mekonnen et al., 2019 [30] | Shashemene/Oromia | January 2014 to December 2016 | 4346 | RPR | 1.1% | High quality |
| Tareke et al., 2019 [23] | Bahir Dar/Amhara | November 2013 to June 2014 | 384 | ICS test | 2.6% | High quality |
| Yitbarek et al., 2019 [24] | Sede Muja District/Amhara | November 2018 to January 2019 | 210 | VDRL | 1.9% | High quality |

ICS immuno-chromatography strips, RPR rapid plasma reagin, SNNP Southern Nations Nationalities and Peoples of Ethiopia, TPHA treponema pallidum hemagglutination assay, VDRL venereal diseases research laboratory, SRIA* syphilis rapid immunoassay test
Factors associated with syphilis sero-positivity

The association of syphilis sero-positivity with marital status, previous history of multiple sexual partner and previous history of STI was quantified using OR. The odds of having syphilis are almost three times higher in pregnant women who had history of multiple sexual partner and

| Study                      | ES (95% CI)     | Weight |
|----------------------------|-----------------|--------|
| Treponemal                 |                 |        |
| Abate Assefa 2014          | 2.90 (2.67, 3.13)| 9.50   |
| Endris et al 2015          | 2.90 (1.47, 4.33)| 6.51   |
| Meliku et al, 2015         | 3.70 (1.37, 6.03)| 4.26   |
| Schonfeld et al, 2017      | 2.20 (1.47, 2.93)| 6.57   |
| Amsalu et al, 2018         | 5.10 (3.18, 7.02)| 5.19   |
| Biado et al, 2019          | 1.90 (1.80, 2.00)| 9.59   |
| Tareke et al, 2019         | 2.60 (1.31, 3.88)| 6.93   |
| Asfaw et al, 2019          | 1.10 (-0.08, 2.28)| 7.28   |
| Subtotal (I-squared = 90.8%, p = 0.000) | 2.53 (1.92, 3.14)| 57.83 |

| Study                      | ES (95% CI)     | Weight |
|----------------------------|-----------------|--------|
| Nontreponemal              |                 |        |
| Mulu et al, 2007           | 1.00 (0.60, 1.40)| 9.26   |
| Zinable et al, 2018        | 1.80 (0.90, 2.70)| 8.10   |
| Ageru et al, 2018          | 3.70 (3.53, 3.87)| 9.55   |
| Yitbarek et al, 2019       | 1.90 (0.16, 3.64)| 5.65   |
| Mekonnen et al, 2019       | 1.10 (0.105, 1.15)| 9.61   |
| Subtotal (I-squared = 99.5%, p = 0.000) | 1.90 (0.40, 3.40)| 42.17 |

| Overall (I-squared = 98.9%, p = 0.000) | 2.32 (1.68, 2.97)| 100.00 |

*NOTE: Weights are from random effects analysis*. 

Fig. 2 Pooled prevalence estimate (ES) of syphilis among pregnant women in Ethiopia

Fig. 3 Funnel plot, evaluating the existence of publication bias for pooled syphilis prevalence
partners than those who had no such partner, OR 2.98 (95% CI, 1.15–7.70). Pregnant women who had previous history of STI were more than four times more likely to develop syphilis than women who had no such history, OR 4.88 (95% CI, 1.35–17.62), whereas married women had 63% lower risk of having syphilis compared to those who are not cohabiting (single, widowed, divorced), OR 0.37 (95% CI, 0.12–0.91) (Table 4).

Discussion

Syphilis is one of the easily preventable and treatable sexually transmitted infections but continued to exert a high burden worldwide especially in sub-Saharan Africa where resources are scarce. Therefore, prevention and control of syphilis among pregnant women using appropriate intervention measures is crucial for the emergence of syphilis free generation. For that, determining the actual burden of syphilis infection and antenatal care (ANC) follow-up screening plays a great role in early diagnosing and treatment of syphilis, and prevention of its vertical transmission. This study aimed to determine the updated pooled prevalence and factors associated to syphilis positivity, and the pooled syphilis-HIV co-infection among pregnant women in Ethiopia.

This meta-analysis showed that the overall pooled prevalence of syphilis among pregnant women in Ethiopia was 2.32% (95% CI, 1.68–2.97). The pooled prevalence was two times higher than the recent nationwide HIV/syphilis sentinel reports among ANC attendees in Ethiopia (1.1%) [33]. The variation in syphilis

Table 2  Sensitivity analysis

| Excluded studies          | Pooled prevalence (95% CI) |
|---------------------------|-----------------------------|
| Mulu et al., 2007 [26]    | 2.46 (1.77, 3.14)           |
| Abate Assefa., 2014 [19]  | 2.26 (1.59, 2.93)           |
| Endris et al, 2015 [21]   | 2.28 (1.62, 2.95)           |
| Melku et al, 2015 [22]    | 2.26 (1.60, 2.92)           |
| Schonfeld et al., 2017 [27]| 2.33 (1.66, 3.01)           |
| Amsalu et al, 2018 [25]   | 2.17 (1.51, 2.83)           |
| Zinabie et al, 2018 [29]  | 2.37 (1.69, 3.04)           |
| Ageru et al, 2018 [28]    | 2.09 (1.59, 2.59)           |
| Biadgo et al, 2019 [20]   | 2.41 (1.50, 3.33)           |
| Tareke et al, 2019 [23]   | 2.30 (1.63, 2.97)           |
| Yitbarek et al, 2019 [24] | 2.35 (1.68, 3.01)           |
| Mekonnen et al, 2019 [30] | 2.45 (1.79, 3.11)           |
| Asfaw et al., 2019 [31]   | 2.42 (1.75, 3.09)           |
| Combined                  | 2.32 (1.68, 2.97)           |

Table 3  Summary estimate for syphilis prevalence in different regions of Ethiopia

| Geographical regions | Included studies | Prevalence (95% CI) | I² |
|----------------------|------------------|---------------------|----|
| Amhara               | 8                | 2.16% (95% CI, 1.57–2.75) | 92.2% |
| Oromia               | 3                | 1.46% (95% CI, 0.69–2.23) | 77.2% |
| SNNP                 | 2                | 4.06% (95% CI, 2.86–5.26) | 50.6% |

**SNNP**: Southern Nations Nationalities and Peoples of Ethiopia

Fig. 4  Pooled prevalence estimate (ES) of syphilis-HIV co-infection among pregnant women in Ethiopia
prevalence between this meta-analysis and the sentinel survey might be due to the rough estimate nature of the sentinel report that might underestimate the actual burden of the diseases. Contrastingly, previous meta-analysis reported higher prevalence of syphilis among pregnant women in Ethiopia, 3.67% [9] relative to this study. This might be because of the previous meta-analysis includes studies with higher syphilis prevalence conducted in the pre-antiretroviral therapy availability. However, given the known synergy between HIV and syphilis [11], we have excluded studies conducted prior to HIV antiretroviral availability in this meta-analysis. Besides, the diseases dynamics may change overtime and impact the observed prevalence [34]. Partly, it may be due to the difference in the number of included studies (5 studies in earlier meta-analysis and 13 articles in this study), indicating the sample size may impact the prevalence.

On the other hand, the pooled prevalence in this meta-analysis was comparable with the report from a countrywide surveillance of HIV/syphilis prevalence among pregnant women attending ANC in Tanzania (2.5%) [35]. The similarity in syphilis prevalence in Ethiopia and Tanzania might be partly due to the WHO’s increased focus and prioritization of antenatal syphilis (in conjunction with HIV and hepatitis B virus) screening for better intervention measures. Besides, combination rapid HIV/syphilis tests are now used to a greater degree in ANC which may also result in increased syphilis testing/diagnoses in different countries of the world to indicate the actual picture of the diseases in various settings.

As shown in subgroup analysis, this study indicated relatively higher prevalence of syphilis using treponemal diagnostic test modality 2.53% (95% CI, 1.92–3.14%) compared to the non-treponemal test 1.90% (95% CI, 0.40–3.40%). This could be due to reactivity to a treponemal test implies infection but it does not determine whether the infection is recent or remote or whether it has been treated or not [32]. Thus, it suggests that treponemal tests stay positive for decades after treatment and may not always indicate active infection. On the other hand, non-treponemal tests have a high false-positive rate and are difficult to interpret on their own [32]. Therefore, considering the difficulty of syphilis diagnosis, the results have to be interpreted with care.

Regional analysis showed a higher and lower prevalence of syphilis in SNNP (4.06%) and in Oromia (1.46%) respectively. Higher syphilis prevalence in SNNP might be attributed to the risky socio-cultural practices such as polygamy is more practiced in SNNP [36] relative to other regions of Ethiopia. Partly, it might be due to the difference in the number of studies included in each category.

In addition, pooled estimate of syphilis-HIV co-infection was also assessed. Consequently, the overall pooled prevalence was 0.80% (95% CI, 0.60–1.01%). A comparable result (0.73%) was obtained from Republic of Congo [37]. While a study in Tanzania reported lower (0.3%) prevalence [35], another study from Rwanda indicated higher (1.2%) prevalence [38] of syphilis-HIV co-infection relative to the findings of this study. The difference in prevalence of syphilis-HIV co-infection might be attributed to the variation in level of implementation and integration of STI prevention and control measures in different countries.

Besides, the association between pregnant women with and without previous history of multiple sexual partner and syphilis sero-positivity was measured in this study. Accordingly, pregnant women with previous history of multiple sexual partners were 2.98 times more likely to get syphilis infection compared to women without such partner. This was consistent with previously established evidences [39–42] and could be attributed to the fact that people with multiple sexual partner has higher risk of getting STI including syphilis. This study also showed that pregnant women with previous history of syphilis were 4.88 times at higher risk of developing syphilis relative to pregnant women without such history. This is in line with the findings from Malawi [43] and China [44]; this could be partly due to lack of behavioral change and other prevention interventions that resulted in maintaining risky behaviors among women who had history of previous STI.

On the other hand, this meta-analysis also demonstrated that married women had 63% lower risk of developing syphilis compared to those who are not cohabiting (single, widowed, divorced). This was comparable with...
earlier studies conducted in rural Tanzania [45] and three sub-Saharan countries [46]. This might be due to the tendency of non-cohabiting women to practice high-risk sexual behaviors like having multiple sexual partners. Partly, this could be because of women in ANC are sexually active age groups, suggesting that if they are non-cohabiting they may have high-risk sexual behaviors.

Limitations
Given the difficulty of syphilis diagnosis, most of the included studies used treponemal diagnostic test alone which may impact the prevalence report. Some regions in the country were not represented in this study due to lack of established original studies in the area. Furthermore, all included studies were facility based. Thus, interpretation of findings has to be with due consideration of these limitations.

Conclusions
This review provides evidence of relatively high prevalence of syphilis among pregnant women. This study also assessed the burden of syphilis–HIV co-infection and determinants of syphilis sero-reactivity in Ethiopia. Therefore, it is recommended to further ramping up of current intervention measures, like routine screening of all ANC women and integration of syphilis testing and treatment to the already established HIV prevention program in the country. Further, nationwide studies involving all regions are needed to assess the magnitude and determinant factors for syphilis among pregnant women in Ethiopia.

Abbreviations
ANC: Antenatal care; CI: Confidence interval; HIV: Human immunodeficiency virus; ICS test: Immuno-chromatography test; OR: Odds ratio; RPR: Rapid plasma reagin; SNNP: Southern Nations Nationalities and Peoples of Ethiopia; STI: Sexually transmitted infection; TPHA: Treponema pallidum hemagglutination assay; VDRL: Venereal diseases research laboratory.

Supplementary Information
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Additional file 1: PRISMA-P 2015 Checklist.

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Authors’ contributions
HG and DG. Conceived and designed the study, reviewed literatures, extracted and analyzed data, interpreted results, and drafted the manuscript. Both authors have read and approved the final manuscript.

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Availability of data and materials
Any data related to this manuscript will be accessible by requesting the corresponding author.

Declarations

Ethics
Not applicable.

Consent for publication
Not applicable.

Competing interests
The authors declare no that they have competing interests.

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