








# BMJ Open Effect of interventions based on educational technologies on the prevention of sexually transmitted infections in incarcerated women: protocol of a systematic review and meta-analysis

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## ABSTRACT

**Introduction** Prisons are places with high vulnerability and high risk for the development of sexually transmitted infections. World Health Agencies recommend establishing intervention measures, such as information and education, on the prevention of diseases. Thus, technologies as tools for health education have been used to reduce sexually transmitted infections. However, no systematic review has investigated the effectiveness of these interventions. Therefore, this review's objective is to examine the effect of educational technologies used for preventing sexually transmitted infections in incarcerated women.

**Methods and analysis** Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines will be strictly followed. The following electronic databases will be searched: Scopus; Cumulative Index of Nursing and Allied Health, Education Resources Information Center, Embase, PsycINFO, PubMed/Medline, Web of Science and Google Scholar. Randomised clinical trials of interventions that used educational technologies to prevent sexually transmitted infections in incarcerated women will be searched in the databases from the beginning of 2020 until December by two researchers independently. A narrative synthesis will be constructed for all included studies, and if there are sufficient data, a meta-analysis will be performed using the Review Manager software (V.5.3). Continuous results will be presented as the weighted mean difference or the standardised mean difference with 95% CIs. Under the heterogeneity of the included studies, a random-effects or fixed-effects model will be used. The studies' heterogeneity will be assessed by the  $I^2$  method. The sensitivity analysis will be carried out to examine the magnitude of each study's influence on the general results. A significance level of  $p \leq 0.05$  will be adopted.

## Strengths and limitations of this study

- This systematic review and meta-analysis will combine results from different studies that have comparable effect sizes.
- The protocol of systematic reviews presents a robust methodology that can serve as a basis for other researchers.
- The study screening, data extraction and quality assessment will be performed by two independent reviewers and a third researcher as an arbitrator.
- Different types of technologies may be identified and cause considerable heterogeneity, influencing clear conclusions.
- The analyses are carried out based on group data and not on the individual data of the participants.

**Ethics and disclosure** Ethical approval is not required because no primary data will be collected. The results will be published in journals reviewed by peers.

**PROSPERO registration number** CRD42020163820.

## INTRODUCTION

There are about 714 000 incarcerated women globally, and these numbers may be higher due to the lack of record or under-reporting of data present in some countries. This number corresponds to 6.9% of the prison population worldwide, and the countries with the highest number of incarcerated women are the USA (200 000), followed by China (107 131), Russia (48 478), Brazil (approximately 44 700) and Thailand (41 119). Furthermore, in the year 2000, this population increased by about 53%, the world population by 21%, and the male prison population by 20%.<sup>1</sup>

Due to their structure, prisons are places with high vulnerability and high risk for the development of sexually transmitted infections (STIs).<sup>2</sup> Such infections can be caused by more than 30 different aetiological agents, and their main route of transmission is sexual contact. Some of these infections have high prevalence rates and are more severe in women because when not detected in time, they can cause, for example, congenital syphilis, pelvic inflammatory disease, infertility and cervical cancer. Besides, there are stigma, discrimination and prejudice resulting from STI's cultural, psychological and biological aspects.<sup>3</sup> Among the researches on STI in incarcerated women, there stands out the association with HIV, syphilis, genital herpes, viral hepatitis, gonorrhoea, chlamydia and human papilloma virus.<sup>4-11</sup>

Moreover, it is crucial to consider that the high number of incarcerated people, coupled with the high turnover present in prison systems and the considerable number of prisoners who will return to the community, makes STI a severe public health problem.<sup>12</sup> In the face of this problem, the WHO proposes comprehensive intervention measures on STI as a strategy for these spaces.<sup>2</sup> Equally, the Bangkok Rules, which discuss the treatment provided to incarcerated women and non-custodial measures for women offenders, report the need to provide information and education on how to prevent STI in prison spaces. These rules were defined to have a global scope concerning these women's particularities. They were developed based on other resolutions adopted by the United Nations, in line with current international law.<sup>13</sup>

Thus, technologies as tools for health education can improve teaching quality through a pedagogical approach considering creativity and interactivity between educator and learner.<sup>14</sup> Furthermore, they can represent one more tool to be considered in preventing STIs in prisons. The educational technologies used for this purpose can be of different types, such as printed materials (cards, pictures, folders e-booklets), games, videos and simulators.<sup>15</sup> In this way, the technologies described here go beyond the universe of digital equipment.

Different technologies can be used in the teaching process of STI, including prisons.<sup>16-18</sup> However, no systematic review was found as the International Prospective Registry of Systematic Reviews (PROSPERO) and Cochrane Database of Systematic Reviews did not have systematic review/meta-analysis records on the subject. Thus, this systematic review with meta-analysis aims to examine the effect of interventions based on educational technologies in the prevention of STIs in incarcerated women.

## METHODS

The protocol was developed according to the Preferred Reporting Items for Systematic Review and Meta-Analysis Protocols<sup>19</sup> and the Cochrane Handbook guidelines for Systematic Reviews of Interventions.<sup>20</sup> The protocol was registered with the PROSPERO.

## Study eligibility

### Characteristics of the studies

The literature search will include studies according to the review's pre-established criteria, objectives, participants, interventions, comparisons and outcomes (PICO). Randomised clinical trials (RCTs) or cluster-RCT primary studies of interventions that used educational technologies in the prevention of STIs in incarcerated women will be selected according to the recommendations of the Cochrane Effective Practice and Organization of Care (EPOC),<sup>21</sup> without language restriction, published until 31 December 2020.

### Participants

Studies with incarcerated women, regardless of age, race, nationality and education level, developed in the prison environment, who have received some intervention using educational technologies related to STI, will be considered. Studies in which it is not possible to identify an educational technology will be excluded.

### Interventions

Studies evaluating the effect of educational technologies as an independent or combined intervention (association with expository or dialogued expository lecture, for example) will be considered interventions. This systematic review will adopt the concept of educational technology as 'the ethical study and practice to facilitate learning and improve performance, creating, using, and managing appropriate technological processes and resources'.<sup>22</sup>

### Comparators

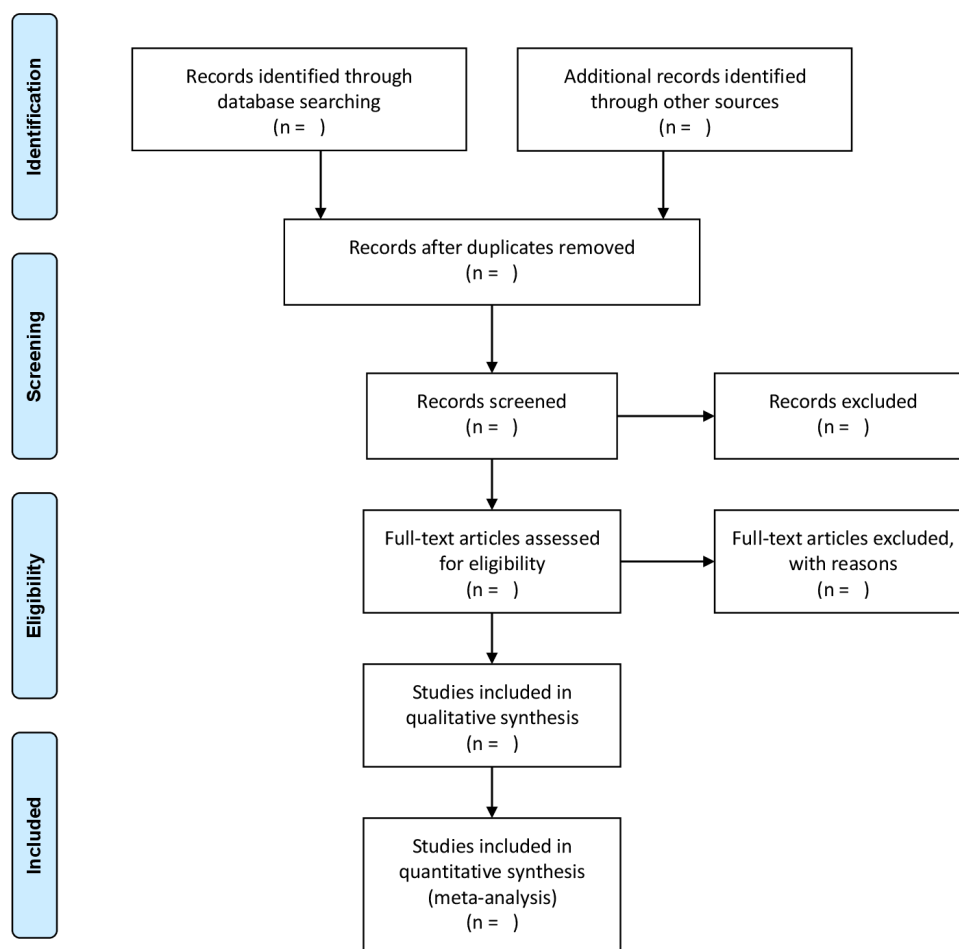
Studies whose control group received any educational intervention without a preventive focus on STI or no educational intervention will be considered comparators.

### Outcome measures

Studies reporting the effect of educational technologies related to sexual risk (reduced number of episodes of unprotected sex or sexual partners or correct use of a condom) in incarcerated women will be considered the outcome measures. According to the concept of educational technologies described above, educational technologies will include digital and non-digital technologies, which involve, for example, printed materials, games, videos and simulators.<sup>15</sup> Secondary outcomes will include knowledge about STI and incidence of STI.

### Information sources

The systematic search will comprise eight electronic databases: Scopus; Cumulative Index of Nursing and Allied Health; Education Resources Information Center; Embase; PsycINFO; PubMed/Medline; Web of Science and Google Scholar. Additionally, a manual search of the list of references of eligible articles will also be carried out to identify any potentially relevant study. The EndNote bibliographical software will be used to manage all references to ensure a comprehensive survey.



**Figure 1** Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) flow chart. Source: PRISMA flow diagram.<sup>24</sup>

The PICO<sup>23</sup> strategy was used to formulate the research question: ‘What is the effect of interventions based on educational technologies on STI prevention in incarcerated women?’ Initially, controlled Medical Subject Headings terms, and their synonyms were selected. The terms were combined using the Boolean operators “AND” and “OR”. The search strategy was developed at PubMed and subsequently will be adapted for other databases (online supplemental appendix 1).

### Selection of studies

Two researchers will independently select scientific articles through the title and abstract’s initial reading, based on the eligibility criteria mentioned above. Subsequently, the articles will be fully read. All differences arising at any step in the selection and analysis process will be solved through consensus between the researchers. In case of no agreement, a third researcher will be contacted to establish a final decision. The reasons for excluding some articles will be presented in a Preferred Reporting Items for Systematic Reviews and Meta-Analyses flow chart (figure 1).<sup>24</sup>

Data extraction will be performed by two researchers with the aid of a specific form<sup>25</sup> and will also consider the data extraction form for primary studies proposed by the

EPOC.<sup>26</sup> The items that will be collected are described below. As the review will assess the effect of educational interventions on sexual risk, the details of the interventions will be selected to establish the studies’ reproducibility.<sup>27</sup> Essential data for the risk of bias assessment (allocation and blinding) will also be extracted. The data will be inserted in the RevMan software (V.5.3.5),<sup>28</sup> which allows for managing and analysing data during a systematic review.

The following items will be extracted from the selected articles: authors; year of publication; country of study; objective; research project; settings; characteristics of the participants (age and ethnicity); health; sampling method; sample size; number and percentage of participants; research method/data collection tool; intervention (type of technology); control (alternative intervention/placebo/conventional care); total duration of the intervention (weeks/months); stages of measurement—baseline, follow-up 1, 2, 3 (weeks/months after the baseline); outcome/discoveries (including mean values, SD and CIs); conclusion of the authors; quality of the study (evaluation of the reviewers/comments on the study).<sup>25 26</sup>

## Risk of bias in individual studies

Two researchers will assess the risk of bias based on the criteria proposed by the EPOC,<sup>29</sup> which includes nine criteria whose outcome may be the following options: low risk, high risk and unclear risk of bias. The criteria are the generation of random sequence; allocation concealment; blinding of participants and personnel; blinding of outcome assessment; incomplete outcome data; selective outcome reporting; and other risks of bias.

The publication bias will be evaluated and classified according to the number of studies included in the meta-analysis: number of studies bigger or equal to 10—funnel chart in RevMan (V.5.3).<sup>28</sup> The interpretation will occur based on the Cochrane Handbook for Systematic Reviews of Interventions.<sup>20</sup>

## Data synthesis

The RevMan (V.5.3) software will be used for the meta-analysis.<sup>28</sup> The meta-analysis will assess the effect of interventions based on educational technologies in the prevention of STI. It will be composed only of studies with a low risk of bias, according to the EPOC.<sup>29</sup> In case of no article with a low risk of bias, the meta-analysis will not be developed. Continuous results will be presented as the weighted mean difference (MD) or the standardised MD with 95% CIs. Under the heterogeneity of the included studies, a random-effects or fixed-effects model will be used. The heterogeneity between the studies will be measured by the Cochran Q and I<sup>2</sup> tests.<sup>30</sup> A significance level of  $p \leq 0.05$  will be adopted.

The size of each study's primary effect will be calculated from the first stage of measurement after implementing the intervention. For studies with more than one intervention group, the primary effect size will be measured for the group presenting the intervention's main characteristics or a smaller number of methodological flaws. In turn, for studies with more than one control group, the primary effect will be calculated for the group presenting the highest expected difference concerning the intervention group.<sup>30</sup>

## Subgroup analysis

If there are enough studies, we will conduct a subgroup analysis based on the different educational technologies used to prevent STIs in incarcerated women.

## Sensitivity analysis

A sensitivity analysis will be conducted to examine the magnitude of each study's influence on the general results of the comparisons.

## Quality of evidence assessment

The quality of evidence will be assessed through the Grading of Recommendations Assessment, Development and Evaluation approach. Each outcome can be classified according to the following levels of evidence: very low, low, moderate and high. These results represent the level of confidence in the estimated effect. The quality of evidence includes methodological limitations

(risk of bias), inconsistency, indirect evidence, inaccuracy and publication bias. Two independent researchers will conduct this quality assessment.<sup>31</sup>

## Qualitative data synthesis

Additionally, a qualitative data synthesis will be presented, even if the meta-analysis is not performed, including a description of aspects such as year of publication, country of origin, sample size, type of intervention and outcomes. A chart will also be presented with a general summary of the educational technologies evaluated.

## Ethics and dissemination

This study is a protocol whose goal is to perform a systematic review and meta-analysis of studies previously published. Thus, there was no human recruitment, and the analyses will not occur concerning the individual participants. Therefore, there is no need for approval from a research ethics committee. The results of this study will be published in journals reviewed by peers.

## Patient and public involvement

No patient involved.

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**Contributors** IdSC—definition of PICO question, writing the protocol and the present manuscript and coordination of the protocol development. RCMGM—outcome definition and review of protocol and manuscript. PdOCM—outcome definition and review of protocol and manuscript. CFS—outcome definition and review of protocol and manuscript. LPL—definition of PICO question, outcome definition and review of protocol and manuscript. TGG—outcome definition and review of protocol and manuscript. GIRO—definition of PICO question, outcome definition and review of protocol and manuscript. FMPL—definition of PICO question, outcome definition and review of protocol and manuscript, and coordination of the protocol development.

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## APPENDIX

### Search Strategy PubMed

("Women"[MeSH Terms] OR "Women"[All Fields] OR "Girl"[All Fields] OR "Woman"[All Fields] OR "Girls"[All Fields] OR "Female"[MeSH Terms] OR "Females"[All Fields] OR "Female"[All Fields] OR "Women's Health"[MeSH Terms] OR "Women's Health"[All Fields] OR "Womens Health"[All Fields] OR "Woman's Health"[All Fields]) AND ("Prisons"[MeSH Terms] OR "Prisons"[All Fields] OR "Prison"[All Fields] OR "Penitentiaries"[All Fields] OR "Penitentiary"[All Fields] OR "Prisoners"[MeSH Terms] OR "Prisoner"[All Fields] OR "Prisoners"[All Fields] OR "Incarceration"[All Fields] OR "Incarcerated"[All Fields]) AND ("Educational Technology"[MeSH Terms] OR "Educational Technology"[All Fields] OR "Educational Technologies"[All Fields] OR "Instructional Technology"[All Fields] OR "Instructional Technologies"[All Fields] OR "Health Education"[MeSH Terms] OR "Health Education"[All Fields] OR "Health Promotion"[MeSH Terms] OR "Health Promotion"[All Fields] OR "Promotion of Health"[All Fields] OR "Health Promotions"[All Fields] OR "Promotional Items"[All Fields] OR "Promotional Item"[All Fields] OR "Wellness Programs"[All Fields] OR "Wellness Program"[All Fields] OR "Health Campaigns"[All Fields] OR "Health Campaign"[All Fields]) AND ("Sexually Transmitted Diseases"[MeSH Terms] OR "Sexually Transmitted Diseases"[All Fields] OR "Sexually Transmitted Disease"[All Fields] OR "STIs"[All Fields] OR "STI"[All Fields] OR "Venereal Diseases"[All Fields] OR "Venereal Disease"[All Fields] OR "Sexually Transmitted Infections"[All Fields] OR "Sexually Transmitted Infection"[All Fields] OR "STDs"[All Fields] OR "Sex Education"[MeSH Terms] OR "Sex Education"[All Fields] OR "Gonorrhea"[MeSH Terms] OR "Gonorrhea"[All Fields] OR "Neisseria gonorrhoeae Infection"[All Fields] OR "Granuloma Inguinale"[MeSH Terms] OR "Granuloma Inguinale"[All Fields] OR "Granuloma Venereum"[All Fields] OR "Donovanosis"[All Fields] OR "Syphilis"[MeSH Terms] OR "Syphilis"[All Fields] OR "Great Pox"[All Fields] OR "Chancroid"[MeSH Terms] OR "Chancroid"[All Fields] OR "Chlamydia Infections"[MeSH Terms] OR "Chlamydia Infections"[All Fields] OR "Chlamydia Infection"[All Fields] OR "Chlamydia"[All Fields] OR "Lymphogranuloma Venereum"[MeSH Terms] OR "Lymphogranuloma Venereum"[All Fields] OR "Lymphogranuloma Inguinale"[All Fields] OR "Condylomata Acuminata"[MeSH Terms] OR "Condylomata Acuminata"[All Fields] OR "Venereal Warts"[All Fields] OR "Venereal Wart"[All Fields] OR "Genital Warts"[All Fields] OR "Genital Wart"[All Fields] OR "buschke lowenstein tumor"[MeSH Terms] OR "buschke lowenstein tumor"[All Fields] OR "buschke lowenstein tumor"[All Fields] OR "Giant Condyloma Acuminatum"[All Fields] OR "Giant Condyloma of Buschke and Lowenstein"[All Fields] OR "buschke lowenstein tumor"[All Fields] OR "buschke lowenstein tumor"[All Fields] OR "Giant Condyloma of Buschke and Lowenstein"[All Fields] OR "Herpes Genitalis"[MeSH Terms] OR "Herpes Genitalis"[All Fields] OR "Genital Herpes"[All Fields] OR "Herpes Simplex Virus

Genital Infection"[All Fields] OR "Genital Herpes Simplex"[All Fields] OR "HIV Infections"[MeSH Terms] OR "HIV Infections"[All Fields] OR "HIV Infection"[All Fields] OR "HTLV III LAV Infections"[All Fields] OR "HTLV III LAV Infection"[All Fields] OR "HTLV III Infections"[All Fields] OR "HTLV III Infection"[All Fields] OR "HIV Coinfection"[All Fields] OR "HIV Coinfections"[All Fields] OR "Acquired Immunodeficiency Syndrome"[MeSH Terms] OR "Acquired Immunodeficiency Syndrome"[All Fields] OR "Acquired Immune Deficiency Syndrome"[All Fields] OR "Acquired Immuno Deficiency Syndrome"[All Fields] OR "Acquired Immunodeficiency Syndromes"[All Fields] OR "AIDS"[All Fields] OR "HIV"[MeSH Terms] OR "HIV"[All Fields] OR "Human Immunodeficiency Virus"[All Fields] OR "Human Immunodeficiency Viruses"[All Fields] OR "Human T Cell Lymphotropic Virus Type III"[All Fields] OR "Human T Cell Leukemia Virus Type III"[All Fields] OR "LAV-HTLV-III"[All Fields] OR "Lymphadenopathy Associated Virus"[All Fields] OR "Human T Lymphotropic Virus Type III"[All Fields] OR "Acquired Immune Deficiency Syndrome Virus"[All Fields] OR "Acquired Immunodeficiency Syndrome Virus"[All Fields] OR "HTLV-III"[All Fields] OR "Papillomavirus Infections"[MeSH Terms] OR "Papillomavirus Infections"[All Fields] OR "Papillomavirus Infection"[All Fields] OR "Human Papillomavirus Infection"[All Fields] OR "Human Papillomavirus Infections"[All Fields] OR "HPV Infection"[All Fields] OR "HPV Infections"[All Fields] OR "HPV"[All Fields] OR "Papillomaviridae"[MeSH Terms] OR "Papillomaviridae"[All Fields] OR "Human Papilloma Virus"[All Fields] OR "Human Papilloma Viruses"[All Fields] OR "Human Papillomavirus Viruses"[All Fields] OR "Human Papillomavirus Virus"[All Fields] OR "hepatitis, viral, human"[MeSH Terms] OR "Human Viral Hepatitides"[All Fields] OR "Human Viral Hepatitis"[All Fields] OR "Hepatitis B"[MeSH Terms] OR "Hepatitis B"[All Fields] OR "Hepatitis C"[MeSH Terms] OR "Hepatitis C"[All Fields] OR "Hepatitis D"[MeSH Terms] OR "Hepatitis D"[All Fields] OR "Delta Hepatitis"[All Fields] OR "Delta Infection"[All Fields] OR "Delta Superinfections"[All Fields] OR "Delta Superinfection"[All Fields] OR "Amazon Black Fever"[All Fields] OR "Labrea Disease"[All Fields])