

Developing and testing of an interactive internet-based intervention to reduce sexual harm of sexualised drug use ('chemsex') among men who have sex with men in Hong Kong

Submission date 15/03/2021	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 17/03/2021	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 09/01/2023	Condition category Infections and Infestations	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Sexualised drug use, known as 'chemsex' or 'chemfun', is the practice of intentionally using illicit drugs before or during sexual activities to enhance sexual arousal and pleasure. This practice, however, poses harmful risks to sexual health. Chemsex is common among men who have sex with men (MSM). Chemsex has been found to increase the likelihood of engaging in other risky sexual behaviours that might increase the spread of sexually transmitted infections (STIs) and HIV.

We propose this trial because chemsex, which is an emerging issue among MSM, substantially increases the risk of HIV and STIs. In spite of this, there has been very limited research evaluating the effectiveness of interventions or health promotion programmes that specifically aim to reduce the intention to engage in chemsex and the actual chemsex behaviours among MSM.

Who can participate?

Men who have sex with men aged over 18 years.

What does the study involve?

Participants will be allocated to one of two groups, with an equal chance of being in either group (like tossing a coin). Participants and researchers will not have a choice in the treatment given. Participants in the first group will receive a web-based treatment containing interactive content that aims to:

1. Lower the desire to use drugs at sex by enhancing participants' knowledge of chemsex, including negative health impacts, risks, social stigmas, and legal consequences
2. Foster a positive attitude towards consistent condom use
3. Foster a positive attitude toward regular HIV/STI testing
4. Increase the self-efficacy of refusing chemsex

5. Improve participants' perceived self-efficacy in making informed decisions on HIV/STI prevention in both sober and drug-influenced sex
6. Set expectations that consistent condom use and regular HIV/STI testing are normative

The second group will receive only web-based information without sexual health components.

Participants in both groups will be evaluated before starting the web-based treatment and three months after starting the web-based treatment.

What are the possible benefits and risks of participating?

One possible benefit is that participants can enhance their sexual health knowledge. There is no significant risk in the study.

Where is the study run from?

School of Nursing, University of Hong Kong (Hong Kong)

When is the study starting and how long is it expected to run for?

From June 2020 to February 2022

Who is funding the study?

Council for the AIDS Trust Fund, Hong Kong (Hong Kong)

Who is the main contact?

Dr Edmond Pui Hang Choi, h0714919@connect.hku.hk

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

ClinicalTrials.gov (NCT)

Nil known

Protocol serial number

Nil known

Study information

Scientific Title

Developing and testing of an interactive internet-based intervention to reduce sexual harm of sexualised drug use ('chemsex') among men who have sex with men in Hong Kong

Study objectives

Participants receiving the internet-based intervention will be more likely to exhibit better self-efficacy in refusing risky sexual behaviours and chemsex, lower intention to engage in chemsex and chemsex behaviours, higher condom use consistency, and more HIV and STI testing, compared to MSM in the control group.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 28/10/2020, Institutional Review Board of the University of Hong Kong/Hospital Authority Hong Kong West Cluster (Room 901, 9/F, Administration Block, Queen Mary Hospital, 102 Pokfulam Road, Hong Kong; +852 2255 4086; hkwirb@ha.org.hk), ref: HKU/HA HKW IRB: UW 20-650

Study design

Randomized controlled trial

Primary study design

Interventional

Study type(s)

Prevention

Health condition(s) or problem(s) studied

Safe sex to prevent sexually transmitted infections (STI)

Interventions

Participants will be randomly assigned to either the intervention group or control group via computer-generated block randomisation (with blocks of size 4) on a 1:1 randomisation ratio; no stratification will be applied. Men who have sex with men will be recruited and randomly allocated into either the intervention (n = 125) or control group (n = 125). Subjects in the intervention group will receive the web-based intervention containing interactive content that aims to:

1. Lower the desire to use drugs at sex by enhancing participants' knowledge of chemsex, including negative health impacts, risks, social stigmas, and legal consequences

2. Foster a positive attitude towards consistent condom use
3. Foster a positive attitude toward regular HIV/STI testing
4. Increase the self-efficacy of refusing chemsex
5. Improve participants' perceived self-efficacy in making informed decisions on HIV/STI prevention in both sober and drug-influenced sex
6. Set expectations that consistent condom use and regular HIV/STI testing are normative

Participants in the control group will receive a web-based intervention without any sexual health information and without any interactive components. The contents for the control group will only include educational materials on general health information.

Intervention Type

Behavioural

Primary outcome(s)

Self-efficacy in refusing risky sexual behaviours and chemsex as measured using the Chinese version of Drug Avoidance Self-Efficacy Scale (DASES), the Self-Efficacy for Sexual Safety scale, and the traditional Chinese version of the Condom Self-Efficacy Scale (CSES) at baseline and 3 months

Key secondary outcome(s)

1. Intention to have chemsex measured using participant interview at baseline and 3 months
2. Actual engagement in chemsex measured using participant interview at baseline and 3 months
3. Practice of condomless sex both sober and drug-influenced measured using participant interview at baseline and 3 months
4. The number of HIV/STI tests in the previous three months measured using participant interview at baseline and 3 months

Completion date

16/02/2022

Eligibility

Key inclusion criteria

1. Cisgender men who have sex with men
2. Aged ≥ 18 years
3. Internet access
4. Able to read and understand Chinese

Participant type(s)

Other

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

Male

Total final enrolment

316

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

15/06/2021

Date of final enrolment

05/11/2021

Locations**Countries of recruitment**

Hong Kong

Study participating centre

School of Nursing, The University of Hong Kong

4/F, William M.W. Mong Block

21 Sassoon Road

Pokfulam

Hong Kong

Hong Kong

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Sponsor information**Organisation**

University of Hong Kong

ROR

<https://ror.org/02zhqgq86>

Funder(s)**Funder type**

Government

Funder Name

Council for the AIDS Trust Fund, Hong Kong

Results and Publications

Individual participant data (IPD) sharing plan

The data sharing plans for the current study are unknown and will be made available at a later date

IPD sharing plan summary

Data sharing statement to be made available at a later date

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	primary and secondary results	05/01/2023	09/01/2023	Yes	No
Protocol article		13/04/2021	15/04/2021	Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes