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Evaluation of the efficacy of a mobile application developed for cannabis users to help them reduce or quit their consumption

Submission date
03/10/2024

Registration date
25/11/2024

Last Edited
05/09/2025

Recruitment status
No longer recruiting

Overall study status
Completed

Condition category
Mental and Behavioural Disorders

[ ] Prospectively registered

[ ] Protocol

[ ] Statistical analysis plan

[ ] Results

[ ] Individual participant data

[X] Record updated in last year

Plain English summary of protocol

Background and study aim
Digital interventions offer opportunities for cannabis users willing to stop or reduce their consumption, but who may be reluctant to seek physical help or are far from a specialised health centre. A personalised 5-week e-coaching program has been designed and developed based on an app in association with other services such as helpline, peer forums and chat services. The main aim of this study is to assess the effectiveness of the cannabis app in helping to stop or reduce cannabis use.

Who can participate?
People aged 18 years or over who are willing to stop or reduce their current cannabis consumption

What does the study involve?
Participants are recruited mainly through the drogues-info-service.fr (DIS) website managed by Santé publique France and randomly allocated to an intervention group or a control group. The intervention group are invited to download the app (available on Google Play and Apple Store). The control group will receive a leaflet with basic information about the harmful effects of cannabis and how to find help. Data will be collected using online questionnaires sent to participants at the start of the study, after the intervention and 3 months later.
To evaluate the effectiveness of the intervention, the number of joints smoked in the last 7 days are compared between the two groups at 3 months after the end of the intervention. Secondly, the number of joints smoked in the last 7 days just after the intervention and the number of days with cannabis consumption in the last 7 days at 3 months will also be compared. The level of satisfaction and perceived utility of the app will also be analysed.

What are the possible benefits and risks of participating?
Participants of the intervention group can continue to use the app beyond the duration of the experiment. Participants in the control group can download the app after answering the last

follow-up questionnaire. Participants who have fully completed the three study questionnaires will receive a gift card worth €20 and access to a random draw allowing five of them to win an additional gift card worth €300.

Where is the study run from?
Santé publique France, the French national public health agency, manages the study. As the experiment involves only the use of a personal smartphone, there is no special or dedicated place to participate.

When is the study starting and how long is it expected to run for?
March 2022 to January 2026

Who is funding the study?
Santé publique France

Who is the main contact?
Emmanuel Lahaie, emmanuel.lahaie@santepubliquefrance.fr

Contact information

Type(s)
Public, Scientific, Principal investigator

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
Efficacy of a digital intervention to reduce or stop cannabis use: study protocol for a randomized controlled trial (ESCAL)

Acronym
ESCAL-protocol

Study objectives
This mobile application increase the chance of reducing or quiting the cannabis use of cannabis users willing to reduce or quit their consumption

Ethics approval required
Ethics approval not required

Ethics approval(s)
In view of the information collected and the type of intervention, this study has been qualified as an evaluation in the field of health outside research involving humans, in accordance with the recommendations of the referal methodology MR-004 of the CNIL (independent administrative authority created by the National Data Protection Act in 1978)

As a research of public interest, the data controller has undertaken to collect only data that is strictly necessary and relevant to the objectives of the research. The protocol has been registered in a public directory held by the Health Data Hub (health-data-hub.fr, number 17990932)

Study design
Interventional two-arm randomized controlled trial

Primary study design
Interventional

Study type(s)
Prevention

Health condition(s) or problem(s) studied
Problematic cannabis use

Interventions
Cannabis users are invited to complete an online information and consent questionnaire and for those eligible a first questionnaire about their use (T0). Those who have completed the T0 questionnaire are included and randomised into two groups. Randomization follows the order of arrival of participants. Allocation is done every other time in each group.

Participants in the intervention group receive:
1. An invitation (email) to download the app, named "Jeanne" and to use it during the 5-week program of the app
2. Invitations to complete two follow-up questionnaires at the end of the program (T1) and 3 months after (T2)

Participants in the control (non-intervention) group receive :
1. A PDF leaflet including basic information about the risks of cannabis use
2. Invitations to complete two follow-up questionnaires at T1 and 3 months after (T2)

Intervention Type
Behavioural

Primary outcome(s)
The number of joints smoked in the last 7 days, measured at 3 months after the end of the intervention (T2)

Key secondary outcome(s))
1. The number of joints smoked in the last 7 days, measured just after the intervention (T1)
2. The number of days with cannabis consumption in the last 7 days, measured at 3 months (T2)
3. The level of satisfaction and perceived utility of the app, measured at T1

Completion date
15/01/2026

Eligibility

Key inclusion criteria
1. Age >= 18 years old
2. Have used cannabis in the last 7 days
3. Willing to reduce or quit cannabis use in the next 15 days
4. Have a smartphone connected to the internet
5. French speaking

Participant type(s)
Population

Healthy volunteers allowed
No

Age group
Adult

Lower age limit
18 years

Upper age limit
99 years

Sex
All

Total final enrolment
629

Key exclusion criteria

1. Not currently following treatment for cannabis addiction
2. Not pregnant
3. Not recently hospitalized for mental disorders
4. Not having used digital or online help to stop or reduce cannabis in the last few months

Date of first enrolment

17/10/2024

Date of final enrolment

31/08/2025

Locations

Countries of recruitment

France

French Guiana

French Polynesia

French Southern Territories

Saint Martin (French part)

Study participating centre
Santé publique France

12, rue du val d'osne
Saint-Maurice
France
94415

Sponsor information

Organisation

Santé Publique France

ROR

https://ror.org/00dfw9p58

Funder(s)

Funder type

Government

Funder Name
Santé Publique France

Results and Publications

Individual participant data (IPD) sharing plan
The datasets generated during and/or analysed during the current study will be available upon request from Emmanuel Lahaie (emmanuel.lahaie@santepubliquefrance.fr). Anonymized data can be shared with external teams after analysis of the project by Santé publique France and the signing of a confidentiality commitment

IPD sharing plan summary
Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Other files	Data collection protocol		11/10/2024	No	No
Participant information sheet			11/10/2024	No	Yes
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes
Study website	Study website	11/11/2025	11/11/2025	No	Yes