

# Blocking online alcohol exposure with a internet browser plugin

<b>Submission date</b> 06/02/2026	<b>Recruitment status</b> Recruiting	<input checked="" type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 10/02/2026	<b>Overall study status</b> Ongoing	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
<b>Last Edited</b> 06/05/2026	<b>Condition category</b> Mental and Behavioural Disorders	<input type="checkbox"/> Individual participant data
		<input checked="" type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

Alcohol imagery is pervasive online. Exposure to alcohol imagery triggers alcohol use and harmful consequences. Through recent advancements in the computational abilities of devices and advances in machine learning, it is now possible to develop a plugin which can detect and block alcohol in images. People with an alcohol use disorder are constantly and sub-consciously seduced to drink through exposure to online alcohol imagery and alcohol ads, which contributes to relapse and treatment failure. Allowing people with an alcohol use disorder to control their online exposure to alcohol imagery could enhance treatment success.

This study aims to test a browser plugin prototype that will block alcohol-related images in a sample of participants treated for alcohol use disorder (AUD). The plugin works similarly to the parental control plugins for sex and violence imagery. Specifically, the study aims to 1) test the feasibility and acceptability of the plugin, and 2) assess the impact of the plugin on perception of alcohol imagery exposure, alcohol craving, and alcohol use.

The project will provide crucial information from end users for the adaptation of a worldwide unique browser plugin to prevent alcohol exposure online. Assessing the acceptability and feasibility of the plugin will lay the foundation for a future large randomized trial.

### Who can participate?

Adult patients with AUD.

### What does the study involve?

The study will be a pilot study with follow-up at 1 and 3 months with an online questionnaire. Participants will be randomly assigned (1:1) to receive access to a browser plugin blocking alcohol imagery or to a control group (access to the plugin provided 3 months later). Study measures will include: hours of online presence, perception of online exposure to alcohol imagery, alcohol use, and alcohol craving. In the intervention group, acceptability of the plugin will be assessed. Participants will also be asked whether they would be willing to use it in the future. Feasibility will be measured by whether the participants kept the plugin active during the intervention period. As this study will be a pilot trial, focus will be on descriptive statistics and effect size estimations using confidence intervals, rather than statistical hypothesis testing.

Descriptive analyses will be conducted on measures of acceptability and feasibility and the confidence intervals will be computed to describe the range of effects on perception of exposure to online alcohol imagery, craving, and alcohol use.

A subset of participants randomized into the browser plugin condition will be invited to a one-to-one audio-recorded semi-structured interview about their experiences with the plugin at the end of the follow-up period. Audio content will be transcribed verbatim, coded and analyzed using inductive research methods.

What are the possible benefits and risks of participating?

Benefits and risks not provided at time of registration

Where is the study run from?

Lausanne University Hospital, Switzerland.

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

April 2026 to February 2027.

Who is funding the study?

Swiss National Science Foundation, Switzerland.

Who is the main contact?

Prof Nicolas Bertholet, nicolas.bertholet@chuv.ch

## Contact information

### Type(s)

Principal investigator, Public, Scientific

### Contact name

Prof Nicolas Bertholet

### ORCID ID

<https://orcid.org/0000-0001-5064-6377>

### Contact details

Lausanne University Hospital

Addiction medicine

Rue du Bugnon 23 A

Lausanne

Switzerland

1011

+41213148400

Nicolas.Bertholet@chuv.ch

## Additional identifiers

## Study information

Scientific Title

Blocking online alcohol exposure with a browser plugin: a proof-of-concept mixed methods, pilot randomized controlled trial

## **Acronym**

PLUGIN study

## **Study objectives**

To test a browser plugin prototype that will block alcohol-related images in a sample of participants treated for alcohol use disorder (AUD).

## **Ethics approval required**

Ethics approval required

## **Ethics approval(s)**

approved 27/01/2026, Commission cantonale d'éthique de la recherche sur l'être humain (CER-VD) (Av. de Chailly 23, Lausanne, 1012, Switzerland; +41213161830; scientifique.cer@vd.ch), ref: 2025-02636

## **Primary study design**

Interventional

## **Allocation**

Randomized controlled trial

## **Masking**

Open (masking not used)

## **Control**

Placebo

## **Assignment**

Sequential

## **Purpose**

Prevention

## **Study type(s)**

## **Health condition(s) or problem(s) studied**

Alcohol use disorder (AUD)

## **Interventions**

The study intervention will be to receive access to a browser plugin blocking alcohol imagery online. The browser plugin prototype is a Chrome extension implemented using a deep learning software framework designed to train deep learning models for the browser environment. The plugin will work by intercepting and scanning every image on a webpage. It will blur the image if it is identified as depicting alcohol. In this pilot phase, it has been chosen to focus on one internet browser. Chrome has been chosen to develop the first prototype as it is the browser with the largest market share in Switzerland (48%).

Intervention: participants in the intervention group will receive access to a plugin blocking (i.e. blurring) alcohol imagery online. The plugin has an “enabled” and a “disabled” button, and thus can be activated or deactivated as the users want. In the intervention group, participants will be provided with the plugin and will be able to use it as much or as little as they want. The study is a pilot trial in which assessing the acceptability and feasibility of this approach is key.

Control: participants in the control group will not be given access to the plugin (no intervention control condition). They will be able to install it at the end of the study.

## **Intervention Type**

Other

## **Primary outcome(s)**

1. Perception of online exposure to alcohol imagery measured using the self reported number of alcohol images seen per hour of online presence at 1 and 3 months
2. Feasibility and acceptability: plugin activation, intervention group only, measured using self-report and usage statistics at 1 and 3 months
3. feasibility and acceptability: Chrome usage, intervention group only, measured using self-reported compliance with using the Chrome browser for online activities at 1 and 3 months
4. Acceptability, intervention group only, measured using items adapted from Van DerLaan and from the Client Satisfaction Questionnaire for Internet interventions (CSQ-I) at 1 and 3 months

## **Key secondary outcome(s)**

1. alcohol craving measured using the Mini Alcohol Craving Experience Questionnaire at 1 and 3 months
2. Alcohol use measured using self reported quantity and frequency questions on alcohol use quantity, frequency, and heavy drinking days at 1 and 3 months

## **Completion date**

28/02/2027

# **Eligibility**

## **Key inclusion criteria**

1. Age 18 or over
2. Current diagnostic of AUD
3. No alcohol use in the past 5 days
4. Using a personal computer to access the internet
5. Willingness to complete the follow-up assessments
6. Willingness to install the plugin and to share its usage data
7. Willingness to use Chrome as their main browser (as the prototype has been developed as a Chrome extension)
8. Able to understand and to sign a written informed consent (written in French) prior to the study

## **Healthy volunteers allowed**

No

**Age group**

Mixed

**Lower age limit**

18 years

**Upper age limit**

100 years

**Sex**

All

**Total final enrolment**

0

**Key exclusion criteria**

People admitted under court order or mandated treatment

**Date of first enrolment**

05/05/2026

**Date of final enrolment**

01/11/2026

**Locations****Countries of recruitment**

Switzerland

**Sponsor information****Organisation**

University Hospital of Lausanne

**ROR**

<https://ror.org/05a353079>

**Funder(s)****Funder type****Funder Name**

Schweizerischer Nationalfonds zur Förderung der Wissenschaftlichen Forschung

**Alternative Name(s)**

Schweizerischer Nationalfonds, Swiss National Science Foundation, Fonds National Suisse de la Recherche Scientifique, Fondo Nazionale Svizzero per la Ricerca Scientifica, Fonds National Suisse, Fondo Nazionale Svizzero, Schweizerische Nationalfonds, The Swiss National Science Foundation (SNSF), SNF, SNSF, FNS

**Funding Body Type**

Private sector organisation

**Funding Body Subtype**

Trusts, charities, foundations (both public and private)

**Location**

Switzerland

## **Results and Publications**

**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not expected to be made available