

BMJ Open Efficacy of pragmatic same-day ring prophylaxis for adult individuals exposed to SARS-CoV-2 in Switzerland (COPEP): protocol of an open-label cluster randomised trial

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ABSTRACT

Introduction Lopinavir/ritonavir (LPV/r) has been proposed as repurposed drugs for pre-exposure and postexposure prophylaxis as well as therapy of COVID-19. Coronavirus postexposure prophylaxis (COPEP) trial aims at assessing their efficacy as postexposure ring-prophylaxis among adults exposed to SARS-CoV-2.

Methods and analysis COPEP is a two-arm open-label cluster-randomised trial conducted in three cantons of Switzerland. Asymptomatic contacts (≥ 16 years) of individuals diagnosed with COVID-19 will be randomised (2:1) to either LPV/r (400 mg/100 mg two times per day) for 5 days, or a standard of care arm (no treatment). Asymptomatic individuals may be either SARS-CoV-2 positive or negative. Contacts living in the single household will form a cluster and will be randomised into the same arm. All participants will be followed-up for 21 days and undergo daily monitoring for COVID-19 symptoms. The primary endpoint is 21-day incidence of laboratory-confirmed COVID-19 with ≥ 1 compatible symptom, analysed in an intention-to-treat (ITT) analysis. The secondary endpoints include the 21-day incidence of COVID-19 as well as SARS-CoV-2 infection in a modified ITT analysis, excluding participants who had a positive SARS-CoV-2 RT-PCR from oropharyngeal swab and/or a positive SARS-CoV-2 IgG serology at baseline. Assuming a 21-day incidence for COVID-19 of 20% among contacts without postexposure chemoprophylaxis, to detect a relative risk reduction of 60% (ie, translating in an absolute reduction from 20% to 8%), with a power of 80%, an alpha of 5%. Accounting for design effect of cluster design of circa 1.1, we plan to enrol 200 participants to the LPV/r arm and 100 to the standard of care arm, 300 participants in total.

Ethics and dissemination Ethics approval has been granted by the Commission Cantonale d'Ethique de la Recherche, Ethikkommission Nordwest- und Zentralschweiz and Comitato Etico Cantonale (ref 2020-

Strengths and limitations of this study

- Among the first clinical trials to study two prophylactic candidates for COVID-19 in the general population as well as healthcare workers.
- The trial will test pragmatic ring prophylactic strategies that can be prescribed the same day with minimal clinical and laboratory assessment.
- Recruitment may be slowed down by the evolution of the epidemic.
- This is an unblinded trial.

00864) and Swissmedic (2020DR3056). Results from this trial will be disseminated via journal articles and presentations at national and international conferences.

Trial registration number Clinicaltrials.gov Registry (NCT04364022); Swiss National Clinical Trial Portal Registry (SNCTP 000003732).

Registered report identifier CCER 2020-0864.

INTRODUCTION

Since the beginning of the SARS-CoV-2 outbreak in December 2019, the medical and scientific community has scrambled to identify effective pharmacological candidates for its prevention and treatment. COVID-19 (which is the clinical manifestation of SARS-CoV-2 infection) has caused unprecedented pressure on modern health systems across the world. The sudden and uncontrolled influx of severe COVID-19 cases has caused critical shortage of hospital-based resources and forced many countries into lockdown in early 2020.



Containing the COVID-19 pandemic may necessitate a multipronged strategy, including immunisation, treatment and prophylaxis. The latter would facilitate the containment of future outbreaks of the disease in non-immunised populations. This current pandemic wave will likely be followed by subsequent outbreaks and authorities will need to consider less strict isolation measures when appraising multiple economic impacts (including on health-related outcome) over a prolonged period of time.

Several drugs have been identified as possible prophylactic candidates. These include the protease inhibitors lopinavir/ritonavir (LPV/r) used for the treatment of HIV.^{1,2} Protease inhibitors work by preventing viral replication and LPV/r has been shown to bind to active site of the SARS-CoV protease in vitro, with studies confirming that the spatial structure of the binding site was conserved between SARS-CoV and SARS-CoV-2.^{3–5} LPV/r is licensed in most countries, are part of the WHO essential list, and are available globally and at low cost.⁶

However, to date, there are no published randomised studies assessing the efficacy of LPV/r as prophylaxis for COVID-19. Coronavirus postexposure prophylaxis (COPEP) is a Swiss two-arm open-label cluster randomised trial that will assess the efficacy, safety and acceptability of same-day LPV/r-based prophylaxis, compared with standard of care alone (no treatment) for asymptomatic individuals exposed to SARS-CoV-2.

METHODS

COPEP is an open-label two-arm (2:1) cluster randomised superiority trial to assess efficacy, safety and acceptability of same-day LPV/r-prophylaxis vs standard of care (no treatment) for asymptomatic adult individuals exposed to SARS-CoV-2. Asymptomatic individuals may be either SARS-CoV-2 positive or negative. The study will be performed at three sites in Switzerland, Geneva, Basel and Lugano, and seeks to recruit 300 participants

over 8–9 months. The main objective of COPEP is to assess prevention of laboratory-confirmed COVID-19 with ≥ 1 compatible symptom, with secondary objectives, including preventing of SARS-CoV-2 infection and attenuation of COVID-19 severity.

Participants

This trial enrolls individuals (≥ 16 years) who have come into close contact with a confirmed case of SARS-CoV-2 infection. Individuals are eligible to be enrolled if the contact occurred within the last 7 days but no more than 72 hours after the index case was diagnosed. A close contact is defined, as per the Swiss Federal Office of Public Health, as a person who spent >15 min in <2 m distance or shared closed space with a confirmed case for a prolonged period (eg, more than 2 hours) in the period extending from 48 hours before onset of symptoms (or before the date of testing in the absence of symptoms), or a person who had direct contact with the body fluids or laboratory specimens of a case without recommended personal protective equipment (PPE) or in case of failure of PPE.

Recruitment of participants to this study will be performed through two routes; (1) the Public Health Authorities in Geneva, Basel and Lugano, who, as the standard of care, contact all individuals tested positive for SARS-CoV-2 to provide them with instructions on isolating procedures and self-monitoring and (2) health-care workers will be recruited via social media platforms and internal hospital-based platforms. All close contacts of confirmed SARS-CoV-2 positive index case will be screened for eligibility (table 1).

Intervention

Participants will be randomised in household clusters, 2:1, to receive either LPV/r, 400 mg/100 mg (2 \times 200 mg/50 mg pills) two times per day for 5 days; or standard of care (no treatment). The first dose of LPV/r will

Table 1 Inclusion and exclusion criteria

Inclusion criteria	Exclusion criteria
1. Documented close contact with a PCR-confirmed SARS-CoV-2 positive individual	1. Fever (temperature $>38.0^{\circ}\text{C}$) and/or respiratory symptoms (cough, dyspnoea) and/or new anosmia/ageusia
2. ≥ 16 years of age	2. Individuals with previous confirmed SARS-CoV-2 infection within the last 6 months
3. Informed consent documented by signature (including parent's or legal guardian's signature if the participant is 16 and 18 years old)	3. Known impairment of liver function
	4. Known hypersensitivity to any of the study medications
	5. Use of any medications that are contraindicated with LPV/r using the website www.hiv-druginteractions.org/checker
	6. Individuals on boosted protease inhibitor as part of an antiretroviral therapy
	7. Inability to be followed-up for the trial period

LPV/r, lopinavir/ritonavir.

be taken during the baseline visit, as directly observed therapy.

Outcomes

Primary endpoint

The primary endpoint is the 21-day incidence of laboratory-confirmed COVID-19 with ≥ 1 compatible symptom in individuals exposed to SARS-CoV-2 who are asymptomatic (either SARS-CoV-2 positive or negative) at baseline. The primary analysis includes all individuals enrolled, irrespective of their baseline oropharyngeal swab results or the baseline SARS-CoV-2 serology (intention-to-treat (ITT) analysis).

Secondary endpoints

- ▶ 21-day incidence of COVID-19 in individuals exposed to SARS-CoV-2 who are asymptomatic, and have negative SARS-CoV-2 PCR and serology at baseline (modified ITT).
- ▶ 21-day incidence of SARS-CoV-2 infection in individuals exposed to SARS-CoV-2 who are asymptomatic and have negative SARS-CoV-2 PCR and serology at baseline (modified ITT).
- ▶ Severity of clinical COVID-19 on a 7-point ordinal scale.
- ▶ Incidence of serious adverse events.

Explorative endpoints

- ▶ Acceptability of a prophylaxis for COVID-19.
- ▶ Reported adherence to LPV/r for participants on the LPV/r arm and HCQ and LPV/r drug levels on day 5 among all individuals.

Measurements

The following clinical definitions will be used. COVID-19 is defined as ≥ 1 symptom compatible with COVID-19 (cough, dyspnoea, anosmia, ageusia, elevated temperature ($>38.0^{\circ}\text{C}$)) and either

1. a positive PCR for SARS-CoV-2 in oropharyngeal swab and/or
2. a seroconversion of IgG only or IgG and IgA for SARS-CoV-2 at day 21 in individuals with negative serology at baseline. In case of seroconversion of IgA only, seroconversion of IgG using more sensitive spike-based recombination immunofluorescence assay (S-rIFA) will be assessed.

New SARS-CoV-2 infection is defined as

1. a positive PCR for SARS-CoV-2 (oropharyngeal swab) among those with a negative PCR at baseline and/or
2. a seroconversion of IgG only or IgG and IgA for SARS-CoV-2 at day 21 in individuals with negative serology at baseline. In case of seroconversion of IgA only, seroconversion of IgG using more sensitive S-rIFA will be used.

Seroconversion for SARS-CoV-2 is defined as negative results for IgG in ELISA at baseline and either

1. positive result for IgG in ELISA and confirmation by S-rIFA at day 21 or

2. doubtful result for IgG in ELISA at day 21 and confirmation by S-rIFA. In case of negative result for IgG in ELISA at day 21 seroconversion can alternatively be defined as follows: Negative result of IgA in ELISA at baseline and (a) positive or doubtful result for IgA in ELISA at day 21 and (b) positive result for IgG in S-rIFA at day 21.

Procedures

All participants will be followed-up for 21 days (figure 1 and table 2). Participants will undergo an oropharyngeal swab for SARS-CoV-2 PCR and a SARS-CoV-2 serology at baseline and on day 21. In addition, participants will be asked to provide on day 5, a capillary puncture on dried blood spot (DBS) to assess blood levels of LPV/r. Participants will be given the option of 'self-test', or of a home-visit for this DBS procedure.

During follow-up, individuals will be asked to complete a daily online COVID-19 symptoms questionnaire. The online questionnaire generates alerts when individuals report a symptom associated with COVID-19 and if participants do not complete the questionnaire for 2 consecutive days. The online questionnaire also serves as a reminder for those on the LPV/r arm, to take their daily medication up to day 5. Paper questionnaires will be made available for those without access to internet. All questionnaires are available in several languages.

The team will contact the participant in case an alert is triggered and do a first clinical assessment by phone. Participants who report COVID-related symptoms (eg, dyspnoea, cough, fever ($>38.0^{\circ}\text{C}$), anosmia) will be asked to come onsite for an extra visit and undergo clinical assessment and an oropharyngeal swab to confirm/exclude SARS-CoV-2 infection. If found positive, participants will be provided with appropriate care, as per local protocol. A follow-up visit 14 days after a positive diagnosis will be conducted, in order to establish the level of clinical manifestation of disease.

On day 21, participants will further complete a questionnaire on adverse events and acceptability of the treatment. Participants on the LPV/r arm are asked to bring the phials which contained the medication to day 21 visit, so that the study team can record the number of returned pills, as per good clinical practice standard.

Sample size

The sample size for the primary endpoint assumes that without treatment 20% of close contacts will develop COVID-19, based on the clinical observations made by the team. To detect a relative risk reduction of 60% (ie, translating in an absolute reduction from 20% to 8%), with a power of 80%, an alpha of 5% and accounting for design effect of cluster design of circa 1.1 (based on $DE=(n-1)r+1$, where $n=3$ denotes the average cluster size, and $r=0.05$ is the assumed intraclass correlation coefficient). A sample size of 200 participants on LPV/r and 100 on standard of care arm, or 300 in total will thus be needed.

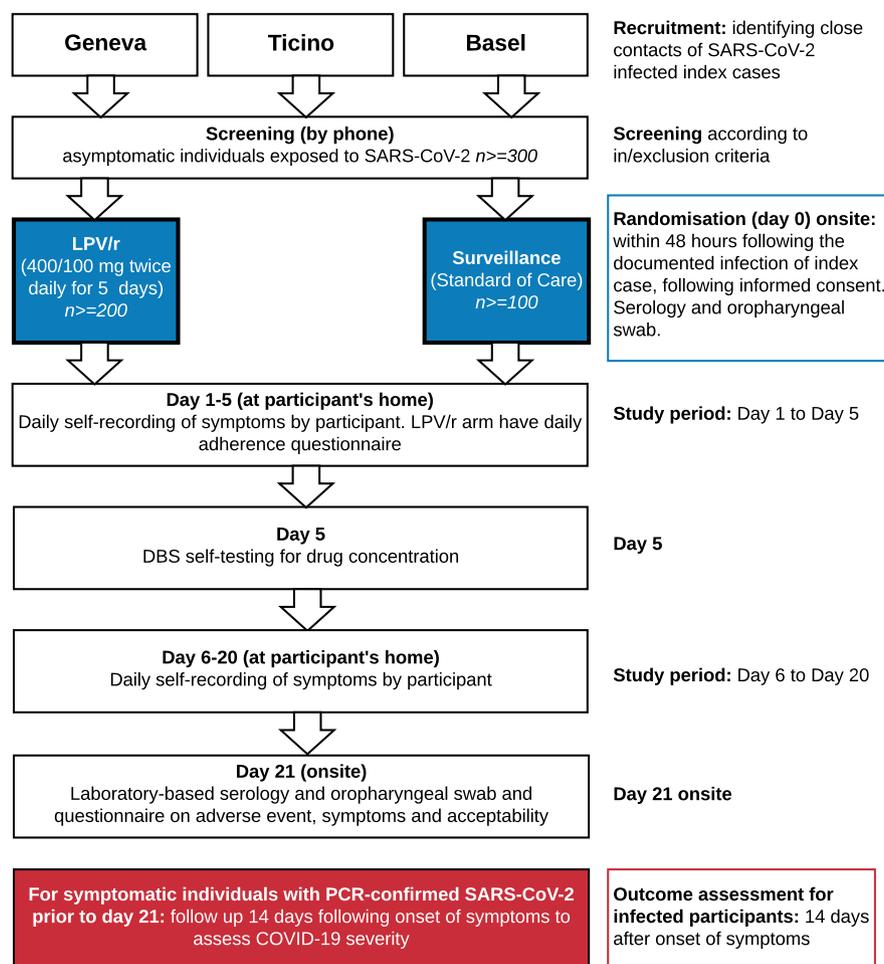


Figure 1 Consolidated Standards of Reporting Trials flow diagram. DBS, dried blood spot; LPV/r, lopinavir/ritonavir.

For the first of the secondary endpoint (occurrence of new SARS-CoV-2 infection) we assume 40% of close contact without prophylaxis will become infected and 16% with prophylaxis; but we also expect a baseline prevalence of positive PCR of 30% (these participants will be excluded from this analysis in a modified ITT). With an effective sample size of 140 individuals on the LPV/r arm (200×0.7) and 70 on the standard of care arm (100×0.7) the power will be 95% for this endpoint.

Randomisation and blinding

Randomisation will be done in variable-sized blocks (sizes 6 or 9) in random sequence, stratified by site (Geneva, Basel and Lugano). Randomisation will be done by cluster, that is where close contacts reside in the same household (eg, the family or friends of the index case that live together). The randomisation procedure will consist of sealed envelopes, one for each household cluster, prepared by individuals who are not associated with the recruitment for the trial. Treatment will be identical within households because of the risk of cross-contamination and to simplify treatment administration to individuals in the same household. This is an open-label trial.

Statistical analysis

For the analysis of the primary endpoint, 21-day incidence of COVID-19, we will perform an ITT, including all individuals who were randomised (including those who will retrospectively be found to be PCR-confirmed SARS-CoV-2 positive at baseline as well as individuals retrospectively found SARS-CoV-2 seropositive by serology).

The intervention arm, LPV/r, will be compared with the surveillance arm, using separate indicator variables for the active treatment arms. Because the hypotheses about treatment efficacy are unrelated and independent, and because we will focus on estimation and CIs rather than on statistical tests, no adjustment will be done for multiplicity. Since individual observations will be clustered within households (randomisation units), themselves nested within index cases, we will use mixed complementary log-log regression models for the main analysis (complementary log-log regression is similar to logistic regression, but yields relative hazards, rather than ORs; relative hazards are more readily interpretable in the context of disease incidence). The outcome variable will be the occurrence of COVID-19 by day 21.

Table 2 Standard Protocol Items: Recommendations for Interventional Trials table

	Screening	Baseline: day 0	Days 1–21 (daily self-assessment)		Day 21
	Phone	Onsite	Participant's home	Day 5*	Onsite
Informed consent procedure	X†	X‡			
Eligibility check; inclusion and exclusion criteria	X§	X¶			
Oropharyngeal swab		X	Prompted by COVID-19 compatible symptoms**		X
Laboratory-based serology		X			X
Randomisation		X			
Administration of prophylaxis††		X	X		
Daily self-assessment			X		
Questionnaire on adherence of intervention‡‡§§			X‡‡§§		
Questionnaire on adverse events				X	X
Dried blood spot collection for blood drug concentration				X	
Questionnaire on acceptability					X

Participants who report COVID-19 symptoms on or prior to day 21; will have onsite visit as soon as possible. An oropharyngeal swab will be performed, and if SARS-CoV-2 is PCR-confirmed, participants will undergo a follow-up visit 14 days after symptoms onset.**

*Participants will be given the option of either a self-test dried blood spot, or home visit by a team member.

†Study information will be provided to participants by phone.

‡Informed consent will be signed by participant and medical investigator onsite.

§Eligibility criteria will be checked with participant by phone.

¶Eligibility criteria will be confirmed.

**Participants with a positive SARS-CoV-2 swab during follow-up will be provided with appropriate care, as per local protocol.

††Only for participants randomised to lopinavir/ritonavir (LPV/r), as directly observed therapy for first dose.

‡‡Only for participants randomised to LPV/r.

§§Only between baseline and day 5.

A random intercept will be defined by each household, nested within the index case. The main fixed effect will be treatment. The main statistical model will be adjusted for potential confounding variables, guided by the most up to date evidence. We foresee that for the incidence of COVID-19, adjustment variables are: age, presence of comorbidities (specifically cardiac, liver or pulmonary disease), treatment of the index case, occupational versus non-occupational exposure and/or positive serology.

In a modified intent to treat analysis, we will evaluate the secondary endpoints; (1) 21-day incidence of COVID-19 in individuals exposed to SARS-CoV-2 who are asymptomatic, PCR-confirmed SARS-CoV-2 negative and have negative SARS-CoV-2 serology at baseline (modified ITT), (2) 21-day incidence of SARS-CoV-2 infection in individuals exposed to SARS-CoV-2 who are asymptomatic, PCR-confirmed SARS-CoV-2 negative and have negative SARS-CoV-2 serology at baseline (modified ITT). The output will be an adjusted relative hazard of the treatment effect versus surveillance, with a 95% CI.

For the secondary outcome 'severity of disease', rated on a 7-point scale, we will use a Mann-Whitney test to compare this ordinal outcome variable between intervention arm and surveillance. For the analysis of individual-reported adverse events, the population will include all randomised participants, and the comparisons will be

by cross-tabulations and χ^2 tests, since the probability of experiencing adverse events will likely not be affected by clustering. For the analysis of acceptability/compliance, only the intervention arm will be evaluated. We will use R software V.1.2.5019 and Stata software V.16 for the analysis.

Data collection and management

Any data collected during the course of this study will be protected under secure password, or kept under lock and key in the research office. No records bearing patient identification will be provided to anyone outside of the institution except regulatory agencies. Patients will not be identifiable as individuals in any publication or presentation of this study. Data will be recorded in RedCap and analysed as described above. The trial will follow all standard procedures, including reporting of adverse events. The protocol was written in accordance with Standard Protocol Items: Recommendations for Interventional Trials checklist (see online supplemental appendix 1)

Patient and public involvement

No patient or patient advisor was involved with study design, recruitment or conduct.

ETHICS AND DISSEMINATION

The study (protocol v3 09.06.2020) has been approved by the following boards: Commission Cantonale d'éthique de la Recherche, Geneva, Switzerland (2020-00864), Ethikkommission Nordwest- und Zentralschweiz, Comitato Etico Cantonale and Swissmedic (Swiss Agency for Therapeutic Products).

All participants will be asked for written informed consent with a dedicated member of the research team. This trial will be conducted in accordance with good clinical practice and the Helsinki Declaration. The WHO Trial Registration Data Set is in online supplemental appendix 2.

The COPEP-trial will establish whether LPV/r is effective as postexposure chemoprophylaxis against clinical COVID-19 and/or SARS-CoV-2 infection compared with surveillance among asymptomatic individuals with recent contact to a person infected with SARS-CoV-2. Furthermore, this trial will aim to investigate whether LPV/r reduces the severity of clinical COVID-19 over surveillance alone, and whether these prophylaxes are safe and acceptable for postexposure prophylaxis of SARS-CoV-2 and COVID-19.

If effective, such a pragmatic approach with prescription the same day and without waiting for laboratory results, will provide a feasible and relatively low-cost strategy to contain local COVID-19 outbreaks in future. All results will be disseminated in peer-reviewed journals and national and international conferences.

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Contributors AC and NDJ conceived the overall trial, including the overall design and therapeutic candidates and are the PI and Co-PI. AC, NDJ, MSm, TP, MSt and AM designed the trial, including decision on intervention dosage and administration, study endpoints and procedures. MSm and TP developed all statistical aspects of the trial design. MSm will perform the trial data analysis, with the support of TP. DL and SF contributed to the formulation of recruitment via the cantonal health authorities, together with MBack and CS. AM and DOA developed the patient and laboratory pathways. BM, LK and LD helped with all laboratory aspects of the trial design. MSm wrote the first draft of this manuscript. GJN, SR, AM, MSt, NDJ and AC are the study physicians and together with FJ, TA, HS, JS, MBattegay, IG and FC who manage the respective patients with COVID-19 or infectious diseases unit contributed to the patient pathway. All authors contributed to the redrafting of the manuscript and in the process of approving the final draft.

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Reporting checklist for protocol of a clinical trial.

Based on the SPIRIT guidelines.

Instructions to authors

Complete this checklist by entering the page numbers from your manuscript where readers will find each of the items listed below.

Your article may not currently address all the items on the checklist. Please modify your text to include the missing information. If you are certain that an item does not apply, please write "n/a" and provide a short explanation.

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		Reporting Item	Page Number
Administrative information			
Title	#1	Descriptive title identifying the study design, population, interventions, and, if applicable, trial acronym	1,2
Trial registration	#2a	Trial identifier and registry name. If not yet registered, name of intended registry	2, 4
Trial registration: data set	#2b	All items from the World Health Organization Trial Registration Data Set	15
Protocol version	#3	Date and version identifier	15
Funding	#4	Sources and types of financial, material, and other support	16

Roles and responsibilities: contributorship	#5a	Names, affiliations, and roles of protocol contributors	1,16
Roles and responsibilities: sponsor contact information	#5b	Name and contact information for the trial sponsor	ICF
Roles and responsibilities: sponsor and funder	#5c	Role of study sponsor and funders, if any, in study design; collection, management, analysis, and interpretation of data; writing of the report; and the decision to submit the report for publication, including whether they will have ultimate authority over any of these activities	16
Roles and responsibilities: committees	#5d	Composition, roles, and responsibilities of the coordinating centre, steering committee, endpoint adjudication committee, data management team, and other individuals or groups overseeing the trial, if applicable (see Item 21a for data monitoring committee)	NA

Introduction

Background and rationale	#6a	Description of research question and justification for undertaking the trial, including summary of relevant studies (published and unpublished) examining benefits and harms for each intervention	6
Background and rationale: choice of comparators	#6b	Explanation for choice of comparators	6
Objectives	#7	Specific objectives or hypotheses	6
Trial design	#8	Description of trial design including type of trial (eg, parallel group, crossover, factorial, single group), allocation ratio, and framework (eg, superiority, equivalence, non-inferiority, exploratory)	6

Methods: Participants,

Interventions, and outcomes

Study setting	#9	Description of study settings (eg, community clinic, academic hospital) and list of countries where data will be collected. Reference to where list of study sites can be obtained	7
Eligibility criteria	#10	Inclusion and exclusion criteria for participants. If applicable, eligibility criteria for study centres and individuals who will perform the interventions (eg, surgeons, psychotherapists)	7-8
Interventions: description	#11a	Interventions for each group with sufficient detail to allow replication, including how and when they will be administered	8
Interventions: modifications	#11b	Criteria for discontinuing or modifying allocated interventions for a given trial participant (eg, drug dose change in response to harms, participant request, or improving / worsening disease)	NA
Interventions: adherence	#11c	Strategies to improve adherence to intervention protocols, and any procedures for monitoring adherence (eg, drug tablet return; laboratory tests)	10-11
Interventions: concomitant care	#11d	Relevant concomitant care and interventions that are permitted or prohibited during the trial	7-8
Outcomes	#12	Primary, secondary, and other outcomes, including the specific measurement variable (eg, systolic blood pressure), analysis metric (eg, change from baseline, final value, time to event), method of aggregation (eg, median, proportion), and time point for each outcome. Explanation of the clinical relevance of chosen efficacy and harm outcomes is strongly recommended	8-9
Participant timeline	#13	Time schedule of enrolment, interventions (including any run-ins and washouts), assessments, and visits for participants. A schematic diagram is highly recommended (see Figure)	10-11
Sample size	#14	Estimated number of participants needed to achieve study objectives and how it was determined, including	11

		clinical and statistical assumptions supporting any sample size calculations	
Recruitment	#15	Strategies for achieving adequate participant enrolment to reach target sample size	10-11
Methods:			
Assignment of interventions (for controlled trials)			
Allocation: sequence generation	#16a	Method of generating the allocation sequence (eg, computer-generated random numbers), and list of any factors for stratification. To reduce predictability of a random sequence, details of any planned restriction (eg, blocking) should be provided in a separate document that is unavailable to those who enrol participants or assign interventions	12
Allocation concealment mechanism	#16b	Mechanism of implementing the allocation sequence (eg, central telephone; sequentially numbered, opaque, sealed envelopes), describing any steps to conceal the sequence until interventions are assigned	12
Allocation: implementation	#16c	Who will generate the allocation sequence, who will enrol participants, and who will assign participants to interventions	12
Blinding (masking)	#17a	Who will be blinded after assignment to interventions (eg, trial participants, care providers, outcome assessors, data analysts), and how	NA
Blinding (masking): emergency unblinding	#17b	If blinded, circumstances under which unblinding is permissible, and procedure for revealing a participant's allocated intervention during the trial	NA
Methods: Data collection, management, and analysis			
Data collection plan	#18a	Plans for assessment and collection of outcome, baseline, and other trial data, including any related processes to promote data quality (eg, duplicate	13-14

		measurements, training of assessors) and a description of study instruments (eg, questionnaires, laboratory tests) along with their reliability and validity, if known. Reference to where data collection forms can be found, if not in the protocol	
Data collection plan: retention	#18b	Plans to promote participant retention and complete follow-up, including list of any outcome data to be collected for participants who discontinue or deviate from intervention protocols	13-14
Data management	#19	Plans for data entry, coding, security, and storage, including any related processes to promote data quality (eg, double data entry; range checks for data values). Reference to where details of data management procedures can be found, if not in the protocol	13-14
Statistics: outcomes	#20a	Statistical methods for analysing primary and secondary outcomes. Reference to where other details of the statistical analysis plan can be found, if not in the protocol	12-13
Statistics: additional analyses	#20b	Methods for any additional analyses (eg, subgroup and adjusted analyses)	12-13
Statistics: analysis population and missing data	#20c	Definition of analysis population relating to protocol non-adherence (eg, as randomised analysis), and any statistical methods to handle missing data (eg, multiple imputation)	12-13
Methods:			
Monitoring			
Data monitoring: formal committee	#21a	Composition of data monitoring committee (DMC); summary of its role and reporting structure; statement of whether it is independent from the sponsor and competing interests; and reference to where further details about its charter can be found, if not in the protocol. Alternatively, an explanation of why a DMC is not needed	NA

Data monitoring: interim analysis	#21b	Description of any interim analyses and stopping guidelines, including who will have access to these interim results and make the final decision to terminate the trial	NA
Harms	#22	Plans for collecting, assessing, reporting, and managing solicited and spontaneously reported adverse events and other unintended effects of trial interventions or trial conduct	13-z14
Auditing	#23	Frequency and procedures for auditing trial conduct, if any, and whether the process will be independent from investigators and the sponsor	NA
Ethics and dissemination			
Research ethics approval	#24	Plans for seeking research ethics committee / institutional review board (REC / IRB) approval	15
Protocol amendments	#25	Plans for communicating important protocol modifications (eg, changes to eligibility criteria, outcomes, analyses) to relevant parties (eg, investigators, REC / IRBs, trial participants, trial registries, journals, regulators)	NA
Consent or assent	#26a	Who will obtain informed consent or assent from potential trial participants or authorised surrogates, and how (see Item 32)	15
Consent or assent: ancillary studies	#26b	Additional consent provisions for collection and use of participant data and biological specimens in ancillary studies, if applicable	NA
Confidentiality	#27	How personal information about potential and enrolled participants will be collected, shared, and maintained in order to protect confidentiality before, during, and after the trial	13-14
Declaration of interests	#28	Financial and other competing interests for principal investigators for the overall trial and each study site	16

Data access	#29	Statement of who will have access to the final trial dataset, and disclosure of contractual agreements that limit such access for investigators	13-14
Ancillary and post trial care	#30	Provisions, if any, for ancillary and post-trial care, and for compensation to those who suffer harm from trial participation	NA
Dissemination policy: trial results	#31a	Plans for investigators and sponsor to communicate trial results to participants, healthcare professionals, the public, and other relevant groups (eg, via publication, reporting in results databases, or other data sharing arrangements), including any publication restrictions	15
Dissemination policy: authorship	#31b	Authorship eligibility guidelines and any intended use of professional writers	NA
Dissemination policy: reproducible research	#31c	Plans, if any, for granting public access to the full protocol, participant-level dataset, and statistical code	NA

Appendices

Informed consent materials	#32	Model consent form and other related documentation given to participants and authorised surrogates	Supplement
Biological specimens	#33	Plans for collection, laboratory evaluation, and storage of biological specimens for genetic or molecular analysis in the current trial and for future use in ancillary studies, if applicable	NA

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Supplementary material

BMJ Open

Supplement 2. The World Health Organization Trial Registration Data Set for the COPEP trial

1. Primary Registry and Trial Identifying Number

Swiss National Clinical Trial Portal (SNCTP 000003732)

2. Date of Registration in Primary Registry

17.04.2020

3. Secondary Identifying Numbers

Clinicaltrials.gov (NCT04364022);

4. Source of Monetary or Material Support

Fondation privée des HUG

5. Primary Sponsor

Fondation privée des HUG

6. Secondary Sponsor(s)

SNF grant submitted

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9. Public Title

Médecin adjointe agréée
Director of the HIV Unite

10. Scientific Title

Chef de Clinique Scientifique
Epidemiologist

11. Countries of Recruitment

Switzerland

12. Health Conditions(s) or Problem(s) Studied

COVID-19

13. Intervention(s)

Supplementary material

BMJ Open

Participants will be randomized in household clusters, 2:1 to receive either LPV/r, 400mg/100mg (2x 200mg/50mg pills) twice daily for 5 days; or surveillance. The first dose of LPV/r will be taken during the baseline visit, as directly observed therapy.

14. Key Inclusion and Exclusion Criteria

Inclusion Criteria

1. Documented close contact with a PCR-confirmed SARS-CoV-2 positive individual;
2. ≥ 16 years of age;
3. Informed consent documented by signature (including parent's or legal guardian's signature if the participant is 16 and 18 y.o.);

Exclusion Criteria

1. Fever (temperature $>38.0^{\circ}$) and/or respiratory symptoms (cough, dyspnoea) and/or new anosmia/ageusia;
2. Individuals with previous confirmed SARS-CoV-2 infection within the last six months;
3. Known impairment of liver function;
4. Known hypersensitivity to any of the study medications;
5. Use of any medications that are contraindicated with LPV/r using the website www.hiv-druginteractions.org/checker
6. Individuals on boosted protease inhibitor as part of an antiretroviral therapy
7. Inability to be followed-up for the trial period

15. Study type:

Study type: interventional

Method of allocation: randomized

Masking: open

16. Date of First Enrollment

23/04/20

17. Sample Size

300

18. Recuitement Status

Recruiting

19. Primary outcomes:

Outcome: incidence of COVID-19

Measurement: ≥ 1 symptom compatible with COVID-19 and either

i) a positive PCR for SARS-CoV-2 in oropharyngeal swab and/or

ii) a seroconversion of IgG only or IgG and IgA for SARS-CoV-2 at day 21 in individuals with negative serology at baseline. In case of seroconversion of IgA only, seroconversion of IgG using more sensitive spike-based recombination immunofluorescence assay (S-rIFA) will be assessed.

Timepoint: 21 days

Supplementary material

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20. Key Secondary Outcomes

Outcome: incidence of COVID-19

Measurement: modified ITT (negative SARS-CoV-2 PCR and serology at baseline)

Timepoint: 21 days

Outcome: incidence of SARS-CoV-2

Measurement: i) a positive PCR for SARS-CoV-2 (oropharyngeal swab) amongst those with a negative PCR at baseline and/or

ii) a seroconversion of IgG only or IgG and IgA for SARS-CoV-2 at day 21 in individuals with negative serology at baseline. In case of seroconversion of IgA only, seroconversion of IgG using more sensitive S-rIFA will be used.

Timepoint: 21 days

Outcome: severity of COVID-19

Measurement: 7-point ordinal scale

Timepoint: 14 days post onset of disease, end of hospitalization where applicable

21. Ethics review

The study has been approved by the following boards: Commission cantonale d'éthique de la recherche, Geneva, Switzerland (2020-00864), Ethikkommission Nordwest- und Zentralschweiz, Comitato Etico Cantonale Swissmedic (Swiss Agency for Therapeutic Products).

All participants will be asked for written informed consent. This trial will be conducted in accordance to Good Clinical Practice and the Helsinki Declaration.

22. Completion data

N/A

23. Summary Results:

N/A

24. IPD sharing statement:

Plan to share IPD: the IPD for this trial will not be made available

Plan description: study protocol