Participant ID:	HVTN 405/HPTN 1901	Visit Code:

## Form: Screening Outcome

informed consent date	
s the participant eligible to enroll in the study?	Yes
if "No", go to "Eligibility status".	No
Group	Group 1 - not previously hospitalized, without specific clinical spectrums or outcomes Group 2 - previously hospitalized, without specific clinical spectrums or outcomes Group 3 - with specific clinical spectrums or outcomes
If "Group 3", specify:	Recovered after intubation
	Prolonged viral shedding
	Myocarditis/pericarditis
	Rapid recovery from COVID-19
	Second positive SARS-CoV-2 RT-PCR test result after a negative result Other
If "Other", specify: (max. 200 characters)	
Eligibility status	Eligible and enrolled  Ineligible  Incomplete Screening
If "Ineligible", select reason(s) why participant is ineligible.	Inclusion Criterion 1 - Age 18 or older.  Inclusion Criterion 2 - Reports having had a positive test for SARS-CoV-2.  Inclusion Criterion 3 - Reports resolution of COVID-19 within 1-8 weeks of enrollment OR, if asymptomatic infection, reports positive SARS-CoV-2 test within 2-10 weeks of enrollment.  Inclusion Criterion 4 - Access to a participating HVTN or HPTN CRS and willingness to be followed for the planned duration of the study.  Inclusion Criterion 5 - Ability and willingness to provide informed consent.  Inclusion Criterion 6 - Assessment of understanding: volunteer demonstrates understanding of this study.

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		Inclusion Criterion 7 - Volunteers
		who were assigned female sex at birth: negative urine or serum beta human chorionic gonadotropin (β-HCG) pregnancy test within 4 days of
		enrollment visit (ie, prior to enrollment blood draw or nasal collections).  Exclusion Criterion 1 - Reports
		current COVID-19. Exclusion Criterion 2 - Pregnant.
		Exclusion Criterion 3 - Receipt of SARS-CoV-2 specific antibodies (eg, convalescent plasma or sera, monoclonal antibodies, hyperimmune globulin).  Exclusion Criterion 4 -
		SARS-CoV-2 vaccine(s) received in a prior vaccine trial.  Exclusion Criterion 5 - Any
		medical, psychiatric, occupational, or other condition that, in the judgment of the investigator, would interfere
		with, or serve as a contraindication to, protocol adherence or a volunteer's ability to give informed consent.
		Volunteer inappropriate for enrollment in the judgement of the investigator.