Participant ID:	CoVPN 5001	Visit Code:

## Form: Screening Outcome

Is the participant eligible to enroll in the study?	Yes
If "No", go to "Eligibility status".	No
Enrollment date	
Group at Enrollment	Group 1 - asymptomatic
	Group 2 - mild symptoms, not
	hospitalized — Group 3 - hospitalized
	Group 3 - Hospitalized
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 - Test result
	indicating presence of SARS-CoV-2 virus.
	Inclusion Criterion 3 - Ability and willingness to provide informed
	consent.
	Inclusion Criterion 4 - Willingness to have clinical
	research staff come to place of
	residence or hospital if needed.
	Inclusion Criterion 5 -
	Willingness to be followed for the planned duration of the
	study.
	Inclusion Criterion 6 -
	Assessment of understanding:  volunteer demonstrates
	understanding of this study.
	Inclusion Criterion 7 -
	Agreement to allow access to
	medical records.
	Inclusion Criterion for Group 1 - No current symptoms.
	Inclusion Criterion for Group 1 -
	No symptoms consistent with
	COVID-19 within 2 weeks prior
	to positive test.  Inclusion Criterion for Group 1 -
	Positive SARS-CoV-2 RNA test or
	antigen test within six days prior
	to enrollment (target time) up to
	10 days prior to enrollment
	(upper allowable window). Inclusion Criterion for Group 2 -
	Onset of mild symptoms
	consistent with COVID-19 within
	six days prior to enrollment
	(target time) up to 14 days prior
	to enrollment (upper allowable window).
	willuow).

Participant ID:	CoVPN 5001	Visit Code:
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		Inclusion Criterion for Group 2 - Positive SARS-CoV-2 RNA test or antigen test within six days prior to enrollment (target time) up to 10 days prior to enrollment (upper allowable window). Inclusion Criterion for Group 3 - Participant hospitalized for COVID-19 within 3 days prior to enrollment. Exclusion Criterion 1 - 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 investigator's judgement

This module contains Form "Screening Outcome" (pages 28-29) from the full document "Prospective Study of Acute Immune Responses to SARS COV-2 Infection"