Subject Enrolment And Informed Consent v2

Subject ID _____

The format for Subject ID is: [IDP][Study number]_AAA_0000

"[IDP][Study Num]" is the Program Code and Study Code for this study
"AAA" is the applicable approved Site Code (only upper case letters). Refer to the Site dropdown menu below for the appropriate code.
"0000" is the Subject ID Number (only digits 0-9)
Underscores must be used (no dashes or other symbols)

If the Subject ID you entered does not match this format, please exit this form and start again.

NOTE: The subject ID appears to be incorrect.

Only underscores may be used (no dashes or other symbols). There should be exactly 14 characters.

Please check the format and correct as necessary.

NOTE: The subject ID appears to be incorrect.

The Program Code should be [IDP] ("[IDP]00"). Only upper case letters are accepted.

Please check the format and correct as necessary

NOTE: The subject ID appears to be incorrect.

The Study Code should be 00 ("[IDP]00")

Please check the format and correct as necessary

NOTE: The subject ID appears to be incorrect.

The Site Code is invalid. Only upper case letters are accepted.

Please check the format and correct as necessary

Was Informed Consent obtained?

- ◯ Yes
- ◯ No

If No, indicate reason: _____

How was consent obtained?

- ◯ Verbally (by telephone)
- ◯ E-consent
- ◯ In-person
- ◯ Other (specify)

Specify other consent method: _____

Who provided consent?

- ☐ Participant
- ☐ Substitute decision maker / parent / study partner

Date consent was signed: _____

NOTE: The date cannot be after today's date or before study start date.

Version date of Informed Consent Form:
- ◯ PAR02_EPO_204_Participant_12MAR2021
- ◯ PAR02_EPO_204_Participant_JULY2020
- ◯ PAR02_EPO_204_Participant_AUG2021
- ◯ PAR02_EPO_204_Participant_DEC2021

Person Obtaining Consent: _____

Assent

Was assent obtained?
- ◯ Yes
- ◯ No
- ◯ Not applicable

Assent Version:
- ◯ Version 1.0
- ◯ Version 2.0

Consent for Future, Unspecified Research

As per TCPS2 updates (new application of article 3.13), a 'separate consent' is required for storage of data and human biological materials for future, unspecified research. Prospective participants must be provided with an option to consent to the study and future research separately.

	Yes	No	Not applicable(i.e., it is included in main study consent)
Was consent collected for future, unspecified research?	◯	◯	◯

Date separate consent for future, unspecific research was signed: _____

NOTE: The date cannot be after today's date or before study start date.

Additional Attestation

Attestation received? ◯ Yes ◯ No

Health Card Number:

Please follow the link below to enter the subject's Ontario Health Card Number in the secure Brain-CODE Subject Registry.

This information is NOT being stored in the study database.

Go to Secure Brain-CODE Subject Registry

Has the subject's Health Card Number been entered into the Subject Registry?	◯ Yes ◯ No
If No, indicate reason:	◯ Refusal ◯ Unknown ◯ Prefer not to answer ◯ Not Applicable ◯ Other (specify)
Other reason:	_____