

INFORMED PATIENT AGREEMENT - COLLECTION OF BIOLOGICAL SAMPLES

1. The patient's data	Name and first name:
	Domicile/ Residence:
	ID card Number/Passport Number:
	Phone Number: Email address:
2. The patient's legal representative* * It is used in the case of minors and adults without discernment (for art. 8 paragraphs (3) - (5) of the methodological norms).	Name and first name:
	Domicile/Residence:
	As:

3. Medical Act (description) 4. Date of collection 5. Consent for the collection of samples	Collection of biological samples _____
	The patient agrees to the collection, the storage and usage of the biological products YES NO
	I also declare that I have been informed about the degree of testing error, understanding that it is assumed by the manufacturer, without the involvement of LABORATOR CUZA VODA SRL. YES NO **Please read the information on the back

6. Requested Investigations 7. Execution time	<input type="checkbox"/> Detection of SARS – CoV - 2 - Real - Time RT – PCR
	<input type="checkbox"/> 12 h <input type="checkbox"/> Romanian <input type="checkbox"/> 24 - 48 h <input type="checkbox"/> English
8. Communication	I understand that the result of the laboratory analyzes will be available online at www.centrul-provita.ro based on the user code and the access password that will be provided to me together with the analysis request, after the expiration of the execution term that was communicated.

**** INFORMATION ON THE DEGREE OF TESTING ERROR**

The test uses viral RNA amplification (E gene, RdRp gene and N gene) of SARS-CoV-2 virus. The result is viral status at harvest, so one or more undetectable results do not rule out the possibility of Covid-19 infection.

One or more negative results, especially from samples taken from the upper respiratory tract, do not exclude the possibility of SARS-CoV2 infection.

A false-negative result can be explained by:

- sample collected too early or too late during the infection;

-appearance of mutations in the virus or inhibitors in the PCR reaction. In patients with a firm suspicion of SARS-CoV-2 infection, additional samples may be taken and tested from the lower respiratory tract.

The identification of another pathogen does not exclude the infection with the new coronavirus, the role of coinfection in pathology not being fully known.

To certify the disposal of the virus, it is recommended to repeat the sampling until the results are negated in two consecutive tests by gene amplification reactions.

My signature represents the fact that I have been informed and I agree with the information in this document.

Date: _____ **Signature (Patient / Legal Representative):** _____