

PATIENT FIRST NAME	PATIENT LAST NAME	PATIENT D.O.B.
SHIPPING ADDRESS	CITY S	STATE ZIP CODE
PATIENT EMAIL	PATIENT PHONE #1	PATIENT PHONE #2
OCCUPATION	EMPLOYER	EDUCATION LEVEL
I, the undersigned patient of the consent.	or representative or legal gu	ardian of the minor named above, give
I am attempting to learn meRNA vaccine or having contracte		e burden in my body from having had the
• • • • • • • • • • • • • • • • • • • •	test "Basic S1 Immune Sub	nefit from the test described by this uset Panel" may help my referring dy.









IMPORTANT NOTICE:

The U.S. Food and Drug Administration (FDA) has NOT approved or cleared this test. However, FDA approval or clearance is not required to use the test for this consent.

This basic S1 Spike Protein with Immune Subset blood test (S1) has been made available for clinical research by Dr. Alan Bain, D.O. The pricing and specifics of this (S1 Spike Protein) test have been negotiated solely for the ongoing study of the Homeopathic Spike Detox (HSD) protocol. Any use of this (Spike Protein S1) test outside of (HSD) or personal research, without going through Dr. Bain, D.O., violates his agreement and may incur a \$25,000 fine per violation. This S1 Spike Protein with Immune Subset panel does not cover all tests offered by IncelIDx. Currently, only tests for classical, intermediate, and nonclassical monocytes with S1 Spike Protein are available; tests for CD4, CD8, the CD4:CD8 ratio, or CBC are not included.

The test determines if I am still encumbered with spike protein which may still challenge my immune system. The test measures S1 spike proteins in monocyte and analyzes if spike protein is still attached. The test was developed by IncellDx, Ca, and its performance characteristics were determined by Radiance Diagnostics. Radiance will conduct the test at Radiance's laboratory, which is separate from Dr. Alan Bain, D.O.'s office. Within 5-7 business days after Radiance received the blood specimen, Radiance will prepare a report indicating the results of the test, which Radiance will deliver to the ordering Physician, Dr. Alan Bain, D.O. and forwarded to your referring physician.

The U.S. Food and Drug Administration (FDA) has not approved or cleared this test. However, FDA approval or clearance is not required to use the test for this consent.

Radiance is a CLIA-Certified high-complexity laboratory. Dr. Alan Bain, D.O. will remit payment for the test to Radiance.

PLEASE INITIAL NEXT TO EACH STATEMENT

The cost of the test may not be covered by my insurance. I understand and I am financially
responsible for the full cost of the test, and I voluntarily consent to pay Dr. Alan Bain, D.O. for the test
n full, in advance and in the amount of \$360.00 USD (the test cost). This test is given at a specialized
discount to protect any conflicts of interest. The test fee does not include any telehealth or office visits
with Dr. Alan Bain, D.O.
If I am not working with a provider I agree to an informed concept concultation with Dr. Alan

_____If I am not working with a provider, I agree to an informed consent consultation with Dr. Alan Bain, D.O. concerning the Test in addition to the above information. I understand the test results review and consultation fee with Dr. Alan Bain is not included in the test kit fee. If you do not have insurance coverage, choose to waive my right to bill insurance OR if I am located outside of Dr. Bain's practicing









area, an Alternative Homeopathic visit fee of \$227 will be due at the time of my visit. I will refrain from discussing the results of my lab, the test price or information obtained during this process publicly, privately and on any social media and chat group. I understand that COVID Long Hauler Cytokine panel, S1 and Immune Subset panel tests are Laboratory Developed Tests (LDT). This test panel was developed by incellDx and their performance characteristics were determined by Radiance Diagnostics, which is certified under the Clinical Laboratories Improvement Amendments of 1988 (CLIA) as qualified to perform high complexity testing. The U. S. Food and Drug Administration (FDA) has not reviewed, approved, or cleared these tests I understand that the laboratory tests performed at Radiance Diagnostics are done at my request. In consideration for receiving the opportunity to participate in laboratory testing, which is provided by The Radiance Diagnostics, I hear by release, waive, discharge, covenant not to sue, and agree to hold harmless for any and all purposes company and their healthcare staff, members, shareholders, officers, servants, agents, volunteers, or employees (herein referred to as "indemnities") from any and all liabilities, claims, demands, injuries, (including death), or damages, including court costs and attorney's fees and expenses, that may be sustained by me while participating in testing. I understand I am responsible for any fees associated with having my blood sample drawn. Phlebotomist fees are not included in the home test kit fee. I am fully responsible for scheduling and locating a phlebotomist to draw my sample. I AGREE to send my sample back to the lab no later than WEDNESDAY to ensure proper delivery and processing time. I am fully aware that, the company is not providing medical care or medical diagnosis with Testing. I further understand that it is my responsibility to consult my own medical doctor for interpretation, analysis, evaluation, and explanation of my test results. I understand that neither Radiance Diagnostics nor its ordering physician will analyze, evaluate, critique, or otherwise interpret the results of said tests. I agree that Radiance Diagnostics, its officers, shareholders, directors, employed physicians or its other agent or employee shall not be liable for any claims including, but not limited to, any claim arising out of or related to, inaccurate, uninterrupted, misinterpreted or results not received and do hereby expressly forever release and discharge all claims, demands, injuries, damage, actions or causes of action. I hereby waive my rights regarding protected health information under HIPAA, to the extent necessary to complete the Testing and to allow Company to provide the results of Testing to the organization which has arranged for the testing or me. Protected health information will not be reused or disclosed by Company to any person or entity other than above, except as required by law.

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l u	understand that there is no refu	nd once the kit has been ordered through Dr. Alan Bain,
PLEASE NO	OTE:	
	consible for finding a phleboton suggest https://www.anylabtes	nist to draw your blood and any fees associated with that tnow.com
TEST ORI	DERING: [BS1IMST] SAR	S-COV2-S1-IMMUNE SUBSET PANEL
	ently COVID 19 PCR Positive? t is the date you first tested pos	
Current Sym	nptoms (Circle all that apply)	
F	atigue	Loss of smell
S	hortness of breath	Sensitivity
Т	emperature	Brain Fog
L	oss of Taste	Cough
N	leuropathy	Insomnia
С	chills	Joint Pain
С	chest Pain	Rash
G	Sastrointestinal Issues	Myalgias
T.	achycardia	N/A









NAME OF REFERRING PROVIDER, PHONE AND EMAIL:

Terms, Cancellation Policy:

I understand that COVID Long Hauler Cytokine panel, S1 and Immune Subset panel tests are Laboratory Developed Tests (LDT). The U. S. Food and Drug Administration (FDA) has not reviewed, approved, or cleared these tests.

I understand that the laboratory tests performed are done at my request. In consideration for receiving the opportunity to participate in laboratory testing, I hear by release, waive, discharge, covenant not to sue, and agree to hold harmless for any and all purposes company and their healthcare staff, members, shareholders, officers, servants, agents, volunteers, or employees (herein referred to as "indemnities") from any and all liabilities, claims, demands, injuries, (including death), or damages, including court costs and attorney's fees and expenses, that may be sustained by me while participating in testing.

I am fully aware that, the lab company is not providing medical care or medical diagnosis with Testing. I further understand that it is my responsibility to consult my own medical doctor for interpretation, analysis, evaluation, and explanation of my test results. I understand that neither the lab nor its ordering physician will analyze, evaluate, critique, or otherwise interpret the results of said tests. I agree that the lab, its officers, shareholders, directors, employed physicians or its other agent or employee shall not be liable for any claims including, but not limited to, any claim arising out of or related to, inaccurate, uninterrupted, misinterpreted or results not received and do hereby expressly forever release and discharge all claims, demands, injuries, damage, actions or causes of action.

I hereby waive my rights regarding protected health information under HIPAA, to the extent necessary to complete the Testing and to allow Company to provide the results of Testing to the organization which has arranged for the testing or me. Protected health information will not be reused or disclosed by Company to any person or entity other than above, except as required by law.

Cancellation and Refund Policy: I understand, I may cancel the order before my consultation with Dr. Alan Bain. Refunds cannot be given once the test kit has been ordered and shipped.









S1 PRIVATE STUDY QUESTIONNAIRE:

The Covid-19 vaccination program was rolled out early in December 2020 with 50% of the total vaccinations being conducted by the end of June 2021. Do you remember the approximate date of your FIRST vaccination(s)?

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PLEASE LIST ANY DISEASE DIA VACCINATIONS.	GNOSIS OR CHRONIC CONDITION TH	AT OCCURRED AFTER YOUR COVID-19
HAVE ANY OF THE ABOVE GO CONDITION WORSENED AND		VID-19 VACCINATION? IF YES, WHAT
PLEASE LIST ANY DISEASE DIA VACCINATION.	GNOSIS OR CHRIONIC CONDITIONS T	THAT HAVE OCCURRED PIOR TO YOUR
PLEASE LIST ANY SYMPTOMS STARTED AND IF THEY STILL P	YOU BELIEVE MAY HAVE BEEN CAUSI ERSIST.	ED BY THE VACCINE, WHEN THEY
19 VACCINE NEGATIVELY EFFE	ECTED MY HEALTH.	THE FOLLOWIN STATEMENT. THE COVIE
WHICH COMPANY DEVELOPE	D THE COVID-19 VACCINATION YOU	RECEIVED?
PLEASE LIST ANY COVID-19 VA	ACCINATION YOU HAVE RECEIVED WI	TH APPROX. DATE OF FIRST DOSE.



HOW MANY TIMES HAVE YOU BEEN INFECTED WITH OR BELIEVE YOU'VE BEEN INFECTED WITH COVID-19, BY YEAR. PLEASE NOTE HOW MANY TIMES NEXT TO EACH YEAR. 2019 2020 _____ 2021 _____ 2022 _____ 2023 +_____ HOW MANY COVID CLINICS HAVE YOU SEEN? HOW MANY SPECIALIST OR TRADITIONAL DOCTORS HAVE YOU SEEN? SINCE COVID-19, HAVE YOU RECEIVED NEW DIAGNOSIS? PLEASE LIST ANY SUPPLEMENTS AND MEDICATIONS YOU ARE CURRENTLY TAKING. PLEASE LIST ANY TREATMENTS YOU HAVE TRIED BUT HAVE FAILED. *WE WILL BE IN CONTACT WITH YOU FOR PAYMENT WITHIN 48 HOURS* SIGNATURE PRINTED NAME DATE

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