

Supplementary Materials

Table S1. Clinical sample information

Demographic, clinical, and selected lab test characteristics of COVID-19 patients. CRP=C-reactive protein, HFNC=high flow nasal canula, RA=room air, FM=face mask, MV=mechanical ventilation, NC=nasal cannula

No.	Gender	Age	Onset of symptoms	%Sat	O2 support	Severity of disease	CRP (mg/dL)	Ferritin (ng/mL)	Absolute Lymph count (10 ³ /uL)	D-dimer (ng/mL)
P1	Male	56	<10 days	85	RA	Severe	19.00	NA	1.10	
P2	Male	62	>10 days	91	RA	Moderate	0.34	NA	1.00	628
P3	Male	65	<10 days	95	RA	Mild	8.90		0.66	
P4	Male	57	<10 days	94	RA	Moderate	2.80		0.73	
P5	Male	66	<10 days	< 90	HFNC	Severe	10.70		0.96	622
P6	Male	60	<10 days	< 90	HFNC	Severe	2.30		1.08	1790
P7	Male	41	<10 days	< 90	HFNC	Severe	14.20	2120	0.78	1509
P8	Female	58	<10 days	96	RA	Mild	0.3	NA	2.6	
P9	Female	64	>10 days	94	RA	Moderate	3.9	NA	1.4	286
P10	Male	52	<10 days	90	NC	Severe	24.8	1332	1.25	
P11	Male	47	<10 days	94	RA	Mild			1.70	
P12	Female	43	<10 days	97	RA	Mild	6.00		1.50	483
P13	Male	74	<10 days	< 90	HFNC	Severe	23.60	830	0.65	1733
P14	Male	48	<10 days	94	RA	Mild	7		1.05	784
P15	Male	48	<10 days	< 90	HFNC	Severe	20	465	0.77	998
P16	Male	57	<10 days	93	RA	Moderate	NA		1.6	
P17	Male	58	<10 days	< 90	HFNC	Severe	4.8		2.1	413
P18	Female	74	<10 days	93	RA	Moderate	2.7		1.48	1258
P19	Male	23	<10 days	98	RA	Moderate	NA			
P20	Male	69	<10 days	< 90	MV	Critical	24.00	136	1.38	11595
P21	Female	99	<10 days	95	100% FM	Critical	19.00		1.00	
P22	Female	61	<10 days	88	RA	Severe	10.90	1685	0.55	451
P24	Male	60	<10 days	96	RA	Mild	3.70	586	0.48	586
P25	Female	40	<10 days	96	RA	Mild	5.7	70	1.06	154
P26	Female	55	<10 days	91	NC	Severe	14.78	244	1.78	
P27	Female	23	<10 days	96	RA	Mild			1.7	
P28	Female	31	>10 days	96	RA	Mild			1.30	
P29	Male	77	>10 days	< 90	MV	Severe	15.00		0.55	4995
P30	Male	48	<10 days	94	RA	Mild	9.40		0.95	
P31	Female	89	<10 days	83	RA	Severe	2.98			
P32	Female	77	<10 days	94	RA	Mild	9.70		0.67	
P33	Female	58	<10 days	96	NC	Severe	10.80	137	1.94	298
P34	Male	58	<10 days	93	NC	Severe	24.60	768	0.60	1166
P35	Male	64	<10 days	93	RA	Mild	6.00	469	1.50	10035
P36	Female	30	<10 days	94	RA	Mild	0.16		1.04	193
P37	Female	80	<10 days	96	RA	Moderate	7.10	306	0.70	580
P38	Female	59	<10 days	97	RA	Mild	0.20		1.10	626

P39	Male	51	<10 days	96	RA	Mild	1.60		1.67	183
P40	Female	76	<10 days	94	RA	Mild	3.2	101	1.48	
P41	Female	87	<10 days	94	NC	Moderate	1.73	295	0.5	
P42	Female	93	<10 days	92	RA	Severe	3.75		2.9	2333
P43	Male	87	<10 days	90	RA	Severe	18.7		0.7	
P44	Male	57	<10 days	94	RA	Moderate	9		0.7	
P45	Male	76	<10 days	90	RA	Severe	4.20		1.30	
P46	Female	90	<10 days	93	NC	Severe	18.70		1.90	313
P47	Female	84	<10 days	90	100% FM	Critical	14.30	670	0.60	
P48	Male	87	<10 days	< 90	HFNC	Severe	32.00	821	0.59	1384
P49	Female	76	<10 days	93	NC	Severe	0.50	85	2.80	
P50	Male	62	<10 days	98	RA	Mild	2.90	528	1.04	299
P51	Male	71	<10 days	93	RA	Moderate	17.00		1.20	2016
P53	Female	67	<10 days	90	NC	Severe	4.00	143	0.75	

Table S2. Gene specific q-PCR primers used in the study

Gene name	Primers
<i>Loop-II</i>	Forward TCAACTGCAACTCCAAAGCC Reverse GGGGACGAGAAGGGATTGA
<i>Loop -I</i>	Forward CGTACATTACTGCCAGCCAC Reverse GGCTTTGGAGTTGCAGTTGA
<i>COX-III</i>	Forward ATGACCCACCAATCACATGC Reverse ATCACATGGCTAGGCCGGAG
<i>ND-2</i>	Forward CACAGAAGCTGCCATCAAGTA Reverse CCGGAGAGTATATTGTTGAAGAG
<i>E</i>	Forward TCGTTTCGGAAGAGACAGGT Reverse ACTAGCCATCCTTACTGCG
<i>N</i>	Forward AGAATGGAGAACGCAGTGGGG Reverse GTCTTGGTTCACCGCTCTCA
<i>3b</i>	Forward TGATGCCAACTATTTTCTTTG Reverse TATTGTAAGGTATACAATAG
<i>β-Actin</i>	Forward CATTGCTGACAGGATGCAGAAGG Reverse TGCTGGAAGGTGGACAGTGAGG

Table S3. Result of a Machine Learning model to predict disease outcomes from mtDNA alone. A predictive model was built with a leave-one-out approach (n=34). The confusion matrix compared predicted and actual outcomes.

		Predicted			
		Healthy	Mild	Moderate	Severe
Actual	Healthy	24	0	0	0
	Mild	0	23	1	5
	Moderate	0	0	7	0
	Severe	0	5	2	4

Description of proteomic-identified proteins found at high levels in COVID-19 patients: their function and relation to diseases

The p49 protein band comprises a number of proteins. These include:

Haptoglobin—a glycoprotein primarily secreted by the liver that binds to hemoglobin, where the haptoglobin–hemoglobin is removed by the reticuloendothelial system [1]. This protein plays a critical role in tissue repair, and regeneration and prevention of oxidative damage, and it is proposed as a biomarker of various pathologies [1]. Haptoglobin is linked to inflammation and many diseases [2], such as diabetic nephropathy [3], Crohn's disease [4], Parkinson's disease [5], and others.

Complement component C8-beta chain—one of three subunits C8 α ($M_r \sim 64,000$), C8 β ($M_r \sim 63,500$), and C8 γ ($M_r \sim 20,400$) of the complement component 8 (C8) protein. C8 is one component of the membrane attack complex (MAC), which mediates cell lysis and initiates membrane penetration of the complex. C8 α and C8 β are involved in the formation of the MAC and affect cell membrane integrity [6]. While C8 γ belongs to the lipocalin family, it is not essential for the MAC-mediated cytolytic activity that kills gram-negative bacteria [7]. The MAC plays a key role in the innate and adaptive immune response by forming pores in the plasma membrane of target cells. A rare defect of the complement classical pathway is associated with susceptibility to severe recurrent infections.

Alpha-1-antichymotrypsin (AACT, SERPINA3)—a serine protease inhibitor that is mainly synthesized in the liver, secreted into the blood, and then induced during inflammation. Its main physiological target is neutrophil cathepsin G, an enzyme that plays a key role in killing of bacteria by the neutrophils. AACT may serve as a diagnostic or prognostic biomarker or therapeutic target in treating tumors [8].

Serum albumin—a 65kDa globular, water-soluble protein that is synthesized by liver hepatocytes and is rapidly excreted into the bloodstream. Albumin is the most abundant protein found in mammal blood and has a very diverse role. It is essential for maintaining plasma oncotic pressure, and it functions as a plasma carrier protein for steroids, hemin, fatty acids, and thyroid hormones. As an anionic protein, albumin binds calcium in blood serum. Its low levels (hypoalbuminemia) can result from a serious underlying medical condition, such as kidney, liver, or heart failure [9].

The p17 protein band comprises the following proteins:

C-reactive protein (CRP)—a pentameric protein synthesized by the liver. CRP has both proinflammatory and anti-inflammatory properties. It is the prototypical acute phase serum protein, which can rise rapidly in response to inflammation and infection [10]. Markedly elevated levels of CRP are most often associated with infections [11] such as severe dengue infection [12]. CRP can activate the classic complement pathway by binding to phosphocholine expressed on the surface of bacterial cells such as pneumococcus. Thus, it is considered to be a relatively sensitive biomarker in infectious and non-infectious inflammation [13]. CRP has been found to trigger cell death in ischemic cells [14]. People infected with COVID-19 in Wuhan, China, had elevated levels of this protein [15, 16], and a correlation between serum CRP and COVID-19 severity has been proposed [17].

Apolipoprotein A-1 (ApoA1)—a main protein constituent of high-density lipoprotein (HDL) particles that has several well-documented cardioprotective functions. ApoA1 binding to lipids is induced by interaction between a highly hydrophobic C-terminal domain with the surface lipids of a particle. As ApoA1 is directly involved in lipid metabolism, this property is important for considering this apolipoprotein as one of the key molecular players in the pathogenesis of cardiometabolic diseases such as atherosclerosis. Beneficial functions of APOA1 in atherosclerosis, thrombosis, diabetes, cancer, and neurological disorders is increasing exponentially [18].

Alpha-1-anti-trypsin (AAT)—a serine protease inhibitor produced in the liver. Its primary function is to protect the lungs from neutrophil elastase which damages delicate lung tissue [19]. AAT deficiency is the most common hereditary disorder in adults [20]. It is associated with an increased risk of developing pulmonary emphysema and liver disease, and also skin problems. The increase observed may reflect the body's response to protect the lungs.

Immunoglobulin Joining chain (J chain)—a 137-residue polypeptide that essential in the formation and stabilization of polymeric Ig structures. It regulates multimerization of secretory IgM and IgA [21].

The p14 protein band includes:

Transthyretin (TTR or TBPA)—a 15.9 kDa that forms a tetrameric protein that transports the thyroid hormone thyroxine and the retinol-binding protein bound to retinol (vitamin A) [22]; TTR is mainly synthesized by the liver (13) and the choroid plexus of brain (14), which are the sources of this protein in plasma and cerebrospinal fluid (CSF), respectively. In humans, 90% of plasma TTR is secreted from liver after binding to the RBP-retinol complex before secretion into the plasma. The protein may also be involved in other intracellular processes including proteolysis, nerve regeneration, autophagy, and glucose homeostasis. TTR misfolding and aggregation forming amyloid transthyretin (ATTR) is known to be associated with amyloid deposition that predominantly affects the peripheral nerves or heart. Patients with ATTR amyloidosis represent a population particularly vulnerable to COVID-19 [23]. TTR is associated with several diseases [24], and functions in the preservation and regulation of memory function and behavior, as well as in protection against several types of neurodegeneration in AD models [25].

Serum amyloid A4 (SAA4)—a constitutive apolipoprotein of high-density lipoprotein. Different isoforms of SAA are expressed constitutively at different levels or in response to inflammatory stimuli. These proteins are produced predominantly by the liver. SAAs composed of two sub-groups, comprising acute-phase SAA1 and SAA2 are associated with HDL and inflammation [26], and SAA4 comprises >90% of the total serum SAA proteins [27]. SAA is also an acute phase marker that responds rapidly, increasing within hours after an inflammatory stimulus, and the magnitude of increase may be greater than that of C-reactive protein (CRP).

Apolipoprotein C-IV—also known as apolipoprotein C4 (APOC4), a lipid-binding protein belonging to the apolipoprotein gene family. It has a role in lipoprotein metabolism and is a critical cofactor for the activation of lipoprotein lipase (LPL), a plasma enzyme that hydrolyzes triglycerides (TGs) on TG-rich lipoproteins [28]. It is associated with diseases such as hyperlipoproteinemia and cardiovascular disease (CVD) [29].

Immunoglobulin kappa constant (IGKC)—immunoglobulin (Ig) isotypes are divided into classes and subclasses depending on the heavy chain they possess; IgM (μ); IgD (δ); IgG (γ) which encompasses subclasses IgG1, IgG2, IgG3, and IgG4; IgA (α) which encompasses subclasses IgA1 and IgA2; and IgE (ϵ) [30]. Each isotype is paired with either a kappa (κ) or lambda (λ) light chain to create a tetrameric immunoglobulin complex capable of triggering unique effector functions. Diseases associated with IGKC include immunoglobulin kappa light chain deficiency and amyloidosis. Among its related pathways are the immune response Fc epsilon RI pathway and immune response nuclear factor of activated T cells (NFAT).

CD5 antigen-like (CD5L)—also called apoptosis inhibitor of macrophage, is secreted primarily from macrophages in lymphoid tissues during an inflammatory response, and it regulates mechanisms in inflammatory responses, such as infection or atherosclerosis [31]. It is expressed in the spleen, lymph nodes, thymus, and bone marrow [32]. This protein acts as an inhibitor of apoptosis and as a key regulator of lipid synthesis, inducing lipolysis in the progression of obesity, and it participates in obesity-associated autoimmunity. Due to its multifaceted roles that range from pattern recognition to autophagy, cell polarization, and the regulation of lipid metabolism, CD5L plays a key role in highly prevalent diseases that develop due to either acute or chronic inflammation, including several infectious, metabolic, and autoimmune conditions [33].

Retinol-binding protein 4 (RBP4)—a transporter protein for retinol from the liver that is stored in the peripheral tissues [34]. In plasma, the RBP-retinol complex interacts with transthyretin that transport retinol from the plasma and cerebrospinal fluid to the liver. After associating with transthyretin (TTR), the retinol/RBP4/TTR complex is released into the bloodstream and delivers retinol to the tissues via binding to specific membrane receptor tissues [34]. We found that both **RBP4** and TTR were

increased in Covid-19 patients (Table 2). This was expected, as after the formation of the complex, retinol/RBP4/TTR is secreted into the circulation [34]. However, it is also reported that RBP4 levels were decreased in hospitalized patients with critical illness compared to nonpatients [35].

Cytosolic retinoic acid receptor responder 2 (CRABP-II)—a soluble protein and member of the family of intracellular lipid-binding proteins transporting retinoic acid (RA) from the cytosol to the nucleus where it directly associates with the RA receptor, thereby enhancing its transcriptional activity [36]. Its functions include antimicrobial activity against bacteria and fungi that [37]. Viral RNA recognition mechanism through RIG-I receptors can quickly consume a large amount of the body's retinoid reserve, which causes the retinol levels to fall below the normal serum levels. This causes retinoid insufficiency and impaired retinoid signaling, which leads to an interruption in Type I interferon synthesis and excessive inflammation. Interestingly, three of the identified proteins **CRABP-II**, **RBP4**, and **transthyretin** are related to retinoid signaling, and thereby the regulation of Type I interferon synthesis and excessive inflammation.

Fig. S1. The sequences with their LC-MS/MS identified peptides of the proteins identified in the protein bands p49, p17 and p14

Protein band-p49

1. Haptoglobin – P00738

MSALGAVIALLLWGQLFAVDSGNDVTDIADDGCPKPPEIAHGYVEHSVRYQCKNYYKLRTEGDGVY
TLNDKKQWINKAVGDKLPECEADDGCPKPPEIAHGYVEHSVRYQCKNYYKLRTEGDGVYTLNNEK
QWINKAVGDKLPECEAVCGKPKNPANPVQRILGGHLDAGSFPWQAKMVSHHNLTTGATLINEQW
LLTTAKNLFLNHSNATAKDIAPTLTLYVGKKQLVEIEKVVLHPNYSQVDIGLIKQKQVSVNERVMP
ICLPSKDYAEVGRVGYVSGWGRNANFKFTDHLKYVMLPVADQDQCIRHYEGSTVPEKKTPKSPVGV
QPILNEHTFCAGMSKYQEDTCYGDAGSAFAVHDLEEDTWYATGILSFDKSCAVAEGVYVKVTSIQ
DWVQKTIAEN

2. Complement component C8 beta chain – P07358

MKNSRTWAWRAPVELFLCAALGCLSLPGSRGERPHSFGSNVKNKSAKSRQMRSDVTLMPIDCE
LSSWSSWTTCDPCQKKRYRYAYLLQPSQFHGEPNCFSDKEVEDCVTNRPCRSQVRCEGFVCAQTGR
CVNRLLCNGDNDGQSDDEANCRRIYKCKCHEMDQYWIGSLASGINLFTNSFEGPVLDRHYAG
GCSPHYILNTRFRKPYNVESYTPQTQGYEFILKEYESYSDFERNVTEKMASKSGFSFGFKIPGIFELGI
SSQSDRGKHYIRRTKRFSHTKSVFLHARSDLEVAHYKLKPRSLMLHYEFLQRVKRLPLEYSYGEYRD
LFRDFGTHYTEAVLGGIYEYTLVMNKEAMERGDYTLNNVHACAKNDFKIGGAIEEVYVSLGVSVG
KCRGILNEIKDRNKRDTMVEDLVVLVRGGASEHITLAYQELPTADLMQEWGDAVQYNPAIKVKVE
PLYELVTATDFAYSSTVRQNMKQALEEFQKEVSSCHCAPCQNGVPVLKGSRCDCICPVGSQGLACE
VSRYRKNTPIDGKWNCWSNWSSCSGRRKTRQRQCNNPPPQNGGSPCSGPASETLDCS

3. Serum albumin OS-69.3kdw – P02768

MKWVTFISLLFLFSSAYSRGVFRDAHKSEVAHRFKDLGEENFKALVLIFAQYLQQCPFEDHVKLV
NEVTEFAKTCVADESAENCDKSLHTLFGDKLCTVATLRETYGEMADCCAKQEPERNECFQHKDDN
PNLPRLVRPEVDVMCTAFHDNEETFLKKYLYEIARRHPYFYAPELLFAKRYKAAFTCECCQAADKAA
CLLPKLDELRLDEGKASSAKQRLKASLQKFGERAFKAWAVARLSQRFPKAEFAEVSCLVDTLTKVH
TECCHGDLLECADDRADLAKYICENQDSISSKLKECCEKPLEKSHCIAEVENDEMPADLPSLAADFV
ESKDVCKNYAEAKDVFGLMFLYEYARRHPDYSVLLLLRLAKTYETTLEKCCAAADPHECYAKVFDE
FKPLVEEPQNLIKQNCLEFEQLGEYKFQNALLVRYTKKVPQVSTPTLVEVSRNLGKVGSKCKKHPEA
KRMPCAEDYLSVVLNQLCVLHEKTPVSDRVTKCTESLVNRRPCFSALEVDETYVPKEFNAETFTFH
ADICTLSEKERQIKQTALVELVKHKPKATKEQLKAVMDDFAAFVEKCKADDKETCFAEEGKKLV
AASQAALGL

4. Alpha-1-antichymo trypsin - P01011

MERMLPLLALGLLAAGFCPAVLCHPNPLDEENLTQENQDRGTHVDLGLASANVDFAFSLYKQLVL
KAPDKNVIFSPLSISTALAFSLGAHNNTLTLEILKGLKFNLTETSEAEIHQSFOHLLRTLNLQSSDELQLS
MGNAMFVKEQLSLLDRFTEDAKRLYGSEAFATDFQDSAAAKKLINDYVKNTRGKITDLIKDLDSQT
MMVLVNYIFFKAKWEMPFDPDQTHQSRFYLSKKKWVMVPMMSLHHLTIPYFRDEELSCTVVELKYT

GNASALFILPDQDKMEEVEAMLLPETLKRWRDSLEFREIGELYLPKFSISR DYNLNDILLQLGIEEAFTS
KADLSGITGARNLAVSQVVHKAVLDVFEEGTEASAATAVKITLLSALVETR TIVRFNRPFLMIIVPTDT
QNIFFMSKVTNPKQA

5. CD5 antigen - O43866

MALLFSLILAICTRPGFLASPSGVRLVGGLHRCEGRVEVEQKGQWGTV CDDGWDIKDVAVLCRELGC
GAASGTPSGILYEPPAEKEQKVLIQSVSCTGTEDTLAQCEQEEVYDCSHDEDAGASCENPESSFSVPVE
GVRLADGPGHCKGRVEVKHQNQWYTVCTGWSLRAAKVVCRLGCGRAVLTQKRCNKHAYGRK
PIWLSQMSCSGREATLQDCPSGPWGKNTCNHDEDTWVECEDPFDLRLVGGDNLCSGRLEVLHKG V
WGSVCDDNWGEKEDQVVCKQLGCGKSLSPSFRDRKCYGPGVGRIWLDNVRCSGEEQSLEQCQHRF
WGFHDCTHQEDVAVICS

Protein band-p17

1. C-creative protein – P02741

MEKLLCFLVLTSLSHAFGQTDMSRKAFVFPKESDTSYVSLKAPLTKPLKAFTVCLHFYTELSSSTRGYSI
FSYATKRQDNEILIFWSKDIGYSFTVGGSEILFEVPEVTVAPVHICTSWESASGIVEFWVDGKPRVRKS
LKKGYTVGAESIILGQEQDSFGGNFEQSLSVGDIGNVMWDFVLSPDEINTIYLGPPFSPNVLNWR
ALKYEVQGEVFTKPQLWP

2. Apolipoprotein A-I OS-30.8kdw – P55056

MKAAVLTAVLFLTGSQARHFWQQDEPPQSPWDRVKDLATVYVDVLKDSGRDYVSQFEGSALGKQ
LNLKLLDNWDSVTSTFSKLREQLGPTQEFWDNLEKETEGLRQEMSKDLEEVKAKVQPYLDDFQKK
WQEEMELYRQKVEPLRAELQEGARQKLHELQEKLSPGGEEMRDRARAHVDALRTHLAPYSDELQR
LAARLEALKENGARLA EYHAKATEHLSTLSEKAKPALEDLRQGLLPVLESFKVSFLSALEEYTKKL
NTQ

3. Alpha-1-antitrypsin OS – PO1009

Isoform 1

MPSSVSWGILLLAGLCCLVPVSLAEDPQGDAQAQKTDTSHHDDQHPTFNKITPNLAEFASFSLYRQLAH
QSNSTNIFFSPVSIATAFAMLSLGTKADTHDEILEGLNFNLTEIPEAQIHEGFQELLRTL NQPDSQLQLTT
GNGLFLSEGLKLVDKFLLEDVKKLYHSEAFTVNFGDTEEAKKQINDYVEKGTQGKIVDLVKELDRDTV
FALVNYIFFKGKWERPFVVKDTEEDDFHVDQVTTVKVPMMKRLGMFNIQHCKKLSSWVLLMKYLG
NATAIFFLPDEGKLQHLENELTHDIITKFLNEDRRSASLHLPKLSITGTYDLKSVLGQLGITKVF SNGA
DLSGVTEEAPLKLSKAVHKAVLTIDEKGTEAAGAMFLEAIPMSIPPEVKFNKPFVFLMIEQNTKSPLFM
GKVVNPTQK

Isoform 2

MPSSVSWGILLLAGLCCLVPVSLAEDPQGDAQAQKTDTSHHDDQHPTFNKITPNLAEFASFSLYRQLAH
QSNSTNIFFSPVSIATAFAMLSLGTKADTHDEILEGLNFNLTEIPEAQIHEGFQELLRTL NQPDSQLQLTT
GNGLFLSEGLKLVDKFLLEDVKKLYHSEAFTVNFGDTEEAKKQINDYVEKGTQGKIVDLVKELDRDTV
FALVNYIFFKGKWERPFVVKDTEEDDFHVDQVTTVKVPMMKRLGMFNIQHCKKLSSWVLLMKYLG
NATAIFFLPDEGKLQHLENELTHDIITKFLNEDRRSASLHLPKLSITGTYDLKSVLGQLGITKVF SNGA
DLSGVTEEAPLKLSKVRSP

4. Complement component C8 gamma chain, P07360

MLPPGTATLLTLLAAGSLGQKPQRPRRASPSTIQPKANFDAQQFAGTWLLVAVGSACRFLQEQQH
RAEATTLHVAPQGTAMAVSTFRKLDGICWQVRQLYGD TGVLGRFLLQARDARGAVHVVAETDYQ
SFAVLYLERAGQLSVKLYARSLPVSDSVLSGFQVRVQEAHLTEDQIFYFPKYGFCEAADQFHVLD EVR
R

5. Immunoglobulin J chain, P01591

MKNHLLFWGVLAVFIKAVHVKAQEDERIVLVDNCKCARITSRIIRSSDPNEDIVERNIRIIVPLNNRE
NISDPTSPLRTRFVYHLSDLCKKCDPTEVELDNQIVTATQSNICDEDSATETCYTYDRNKC YTAVVPL
VYGETKMOVETALTPDACYPD

6. Retinoic acid receptor responder protein – P02753

MRRLLIPLALWLGA VGVGVAELTEAQRRLGLQVALEEFHKHPPVQWAFQETSVESAVDTPFPAGIFVR
LEFKLQQTSCRKRDWKKPECKVRPNGRKRKCLACIKLGSEDKVLGRLVHCPIETQVLR EAEHQETQ
CLRVRAGEDPHSFYFPGQFAFSKALPRS

7. Retinol-binding protein 4 – Q99969

MKWVWALLLLAALGSGRAERDCRVSSFRVKENFDKARFSGTWYAMAKKDPEGLFLQDNIVAEFSV
DETQMSATAKGRVRLNNWVDCADMVGTFTDTPAKFKMKYWGVASFLLQKGNDDHWIVDTD
YDTYAVQYSCRLLNLDGTCADSYSFVFSRDPNGLPPEAQKIVRQRQEELCLARQYRLIVHNGYCDGR
SERNLL

Protein band-p14

1. Transthyretin-1 – P02766

MASHRLLLLCLAGLVFVSEAGPTGTGESKCPLMVKVLDAVRGSPAINVAHVFRKAADDTWEPFAS
GKTSESGELHGLTTEEEFVEGIYKVEIDTKSYWKALGISPFHEHAEEVFTANDSGPRRYTIAALLSPYS
YSTTAVVTNPKE

2. Serum amyloid – P35542

MRLFTGIVFCSLVMGVTSESWRSFFKEALQGVGDMGRAYWDIMISNHQNSNRPLYARGNYDAAQR
GPGGVWAAKLISRSRVYLQGLIDCYLFGNSSTVLEDSKSNEKAEWGRSGKDPDRFRPDGLPKKY

3. Apolipoprotein C – P55056

MSLLRNRLQALPALCLCVLVLACIGACQPEAQEGTSLPPPCLKMSRWSLVRGRMKELLETVVNRTD
GWQWFWSPTFRGFMQTYDDHLRDLGLPLTKAWFLESKDSLLKKTHSLCPRLVCGDKDQG

4. Immunoglobulin kappa constant - P01834

RTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYS
LSSTLTLSKADYEEKHKVYACEVTHQGLSSPVTKSFNRGEC

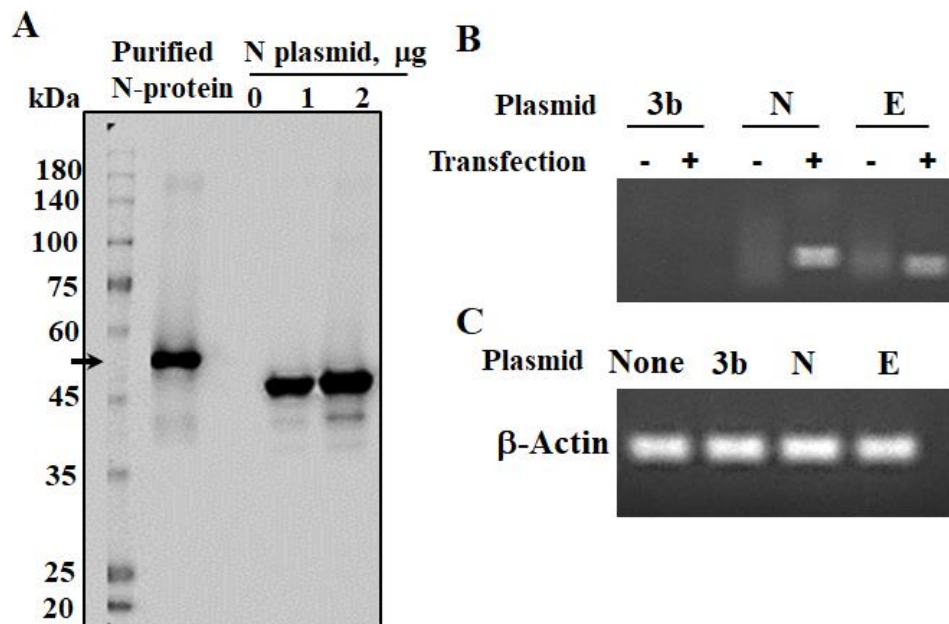


Fig. S2: Cells expression of SARS-CoV-2 proteins

(A) SDS-PAGE and Western blot of purified SARS-CoV-2 N-protein C-terminal DYKDDDDK tagged (indicated by an arrow) (Cat. No. S014660-05; Synbio Technologies; NJ, USA) and lysates of SHSY-5Y cells 48h post transfection with 1 or 2 μ g of N-Protein encoding plasmid. (B,C) SHSY-5Y cells were transfected with empty plasmid or encoding for N, E or 3b proteins. 48h post transfection cells were harvested, and lysates were subjected to PCR using specific primers for 3b, N, E (B) or for β -Actin shown as a loading control (C).

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