

Supplementary material to

**Identification and analysis of unstructured, linear B-cell epitopes in SARS-CoV-2 virion proteins
for vaccine development**

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Supplementary Data S1. Conservation of B-cell epitopes in S, E, M and N protein across coronavirus species. Independent sequence alignments between uBCELS/BCEH of SARS-CoV-2, and equivalent sequences in SARS-CoV and three bat coronavirus samples: bat-SL-CoVZXC21, bat-SL-CoVZC45 and RaTG13. For clarity, SARS-CoV and SARS-CoV-2 headers are in bold letters. Residues in uBCEs are in capital letters while the rest of the loop is in lowercase. Identical residues to SARS-CoV-2 sequences are highlighted in green. Deletions are indicated with dashes.

S protein

uBCEL-S1

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SARS-CoV          20_ddvqaPNYTQHTS_32
Bat-SL-CoVZXC21   17_-dltGRTPLINENY_28
Bat-SL-CoVZC45   17_vnltGRTPLINENY_29
RaTG13            15_vnlttRTQLPPAY_27
SARS-CoV-2      15_vnlttRTQLPPAY_27
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uBCEL-S2

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SARS-CoV          72_in-----HTFGNpvi_p_82
Bat-SL-CoVZXC21   68_lttN-NAAATKRFDNpild_84
Bat-SL-CoVZC45   69_lttN-NAAATKRFDNpild_85
RaTG13            68_ihvSGTNGIKRFDNpvlp_85
SARS-CoV-2      68_ihvSGTNGTKRFDNpvlp_85
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uBCEL-S3

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SARS-CoV          389_vkGDDVRQIAPggtgviadynyklpddf_416
Bat-SL-CoVZXC21   397_lrFSEVRQVAPggtgviadynyklpddf_424
Bat-SL-CoVZC45   398_lrFSEVRQVAPggtgviadynyklpddf_425
RaTG13            402_itGDEVQRQIAPggtgkiadynyklpddf_429
SARS-CoV-2      402_lrGDEVQRQIAPggtgkiadynyklpddf_429
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uBCEL-S4

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SARS-CoV          427_NIDATStgnyn_437
Bat-SL-CoVZXC21   435_KQDT-----gh_440
Bat-SL-CoVZC45   436_KQDV-----gn_441
RaTG13            440_HIDAKEggnfn_450
SARS-CoV-2      440_NLDSKVggnyn_450
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uBCEL-S5

SARS-CoV 442_ylrhGKLRPFERDISNVPFSPDGKPCtp-palncywp_477
 Bat-SL-CoVZXC21 445_shrsTKLKPFERDLSSDENG-----vrt_467
 Bat-SL-CoVZC45 446_shrsTKLKPFERDLSSDENG-----vrt_468
 RaTG13 455_lfrkANLKPFERDISTEIQAGSKPCngqtglncyyp_491
SARS-CoV-2 455_lfrkSNLKPFERDISTEIQAGSTPCngvegfnicyfp_491

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uBCEL-S6

SARS-CoV 601_VNCTDVSTAIHADQLTpawriystgnnv_628
 Bat-SL-CoVZXC21 591_VNCTDVPTTIHADQLTpawriyaigtsv_618
 Bat-SL-CoVZC45 592_VNCTDVPTTIHADQLTpawriyatgtnv_619
 RaTG13 615_VNCTEVPVAIHADQLTptwrvystgsv_642
SARS-CoV-2 615_VNCTEVPVAIHADQLTptwrvystgsv_642

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uBCEL-S7

SARS-CoV 662_TVSL----LRSTsq_671
 Bat-SL-CoVZXC21 652_TASI----LRSTgq_661
 Bat-SL-CoVZC45 653_TASI----LRSTsq_662
 RaTG13 676_TQTNS----RSVas_685
SARS-CoV-2 676_TQTNSPRRARSVas_689

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uBCEL-S8

SARS-CoV 765_AQVKQMYKTPTLKYFggfnfs_785
 Bat-SL-CoVZXC21 755_AQVKQIYKTPPIKDFggfnfs_775
 Bat-SL-CoVZC45 756_AQVKQIYKTPPIKDFggfnfs_776
 RaTG13 779_AQVKQIYKTPPIKDFggfnfs_799
SARS-CoV-2 783_AQVKQIYKTPPIKDFggfnfs_803

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uBCEL-S9

SARS-CoV 1107_NCDVVIG_1113
 Bat-SL-CoVZXC21 1097_NCDVVIG_1103
 Bat-SL-CoVZC45 1098_NCDVVIG_1104
 RaTG13 1121_SCDVVIG_1127
SARS-CoV-2 1125_NCDVVIG_1131

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uBCEL-S10

SARS-CoV	1118	TVYDPLQPELDS	1129
Bat-SL-CoVZXC21	1108	TVYDPLQPELDS	1119
Bat-SL-CoVZC45	1109	TVYDPLQPELDS	1120
RaTG13	1132	TVYDPLQPELDS	1143
SARS-CoV-2	1136	TVYDPLQPELDS	1147

uBCEL-S11

SARS-CoV	1220	tsCCSCLKG	1228
Bat-SL-CoVZXC21	1210	tsCCSCLKG	1218
Bat-SL-CoVZC45	1211	tsCCSCLKG	1219
RaTG13	1234	tsCCSCLKG	1242
SARS-CoV-2	1238	tsCCSCLKG	1246

M protein

uBCEL-M1

SARS-CoV	208	DHAGSN	Dnia	llvq	221
Bat-SL-CoVZC45	209	DHSSSS	Dnia	llvq	222
Bat-SL-CoVZXC21	209	DHSSSS	Dnia	llvq	222
RaTG13	209	DHSSSS	Dnia	llvq	222
SARS-CoV-2	209	DHSSSS	Dnia	llvq	222

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N protein

uBCEL-N1

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SARS-CoV          19_ GGPTDSTDNQNGGRNGARPKQRRPQGLPNntaswfta 56
Bat-SL-CoVZXC21   18_ GGPSDSSDNSQNGERNGARPKQRRPQGLPNntaswfta 55
Bat-SL-CoVZC45    18_ GGPSDSSDNSKNGERNGARPKQRRPQGLPNntaswfta 55
RaTG13            18_ GGPSDSTGSNQNGERSGARPKQRRPQGLPNntaswfta 55
SARS-CoV-2        18_ GGPSDSTGSNQNGERSGARSKQRRPQGLPNntaswfta 55
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uBCEL-N2

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SARS-CoV          60_ HGKEELRFPRGQGVPIINTNS 79
Bat-SL-CoVZXC21   59_ HGKENLTFPRGQGVPIINTNS 78
Bat-SL-CoVZC45    59_ HGKENLTFPRGQGVPIINTNS 78
RaTG13            59_ HGKEDLKFPRGQGVPIINTNS 78
SARS-CoV-2        59_ HGKEDLKFPRGQGVPIINTNS 78
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uBCEL-N3

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SARS-CoV          136_ tegalntpkdhigtrnpnnaatVLQLPQGTTLPKG 171
Bat-SL-CoVZXC21   135_ tegalntpkdhigtrnpannaaiVLQLPQGTTLPKG 170
Bat-SL-CoVZC45    135_ tegalntpkdhigtrnpannaaiVLQLPQGTTLPKG 170
RaTG13            135_ tegalntpkdhigtrnpannaaiVLQLPQGTTLPKG 170
SARS-CoV-2        135_ tegalntpkdhigtrnpannaaiVLQLPQGTTLPKG 170
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uBCEL-N4

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SARS-CoV          174_ AEGSRGGSQASSRSSRSRGNSSRNSTPGSSRGNSPArmasg 214
Bat-SL-CoVZXC21   173_ AEGSRGGSQASSRSSRSRNSSRNSTPGSSRGTSPArmagn 213
Bat-SL-CoVZC45    173_ AEGSRGGSQASSRSSRSRNSSRNSTPGSSRGTSPArmagn 213
RaTG13            173_ AEGSRGGSQASSRSSRSRNSSRNSTPGSSRGTSPArmagn 213
SARS-CoV-2        173_ AEGSRGGSQASSRSSRSRNSSRNSTPGSSRGTSPArmagn 213
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uBCEL-N5

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SARS-CoV          236_ SGKGOQQGQTVT 248
Bat-SL-CoVZXC21   235_ SGKGOQQGQTVT 247
Bat-SL-CoVZC45    235_ SGKGOQQGQTVT 247
RaTG13            235_ SGKGOQQGQTVT 247
SARS-CoV-2        235_ SGKGOQQGQTVT 247
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uBCEL-N6

SARS-CoV	277	RRGPEQTQGNFG	288
Bat-SL-CoVZXC21	276	RRGPEQTQGNFG	287
Bat-SL-CoVZC45	276	RRGPEQTQGNFG	287
RaTG13	276	RRGPEQTQGNFG	287
SARS-CoV-2	276	RRGPEQTQGNFG	287

uBCEL-N7

SARS-CoV	340	LDDKDP	345
Bat-SL-CoVZXC21	339	LDDKDP	344
Bat-SL-CoVZC45	339	LDDKDP	344
RaTG13	339	LDDKDP	344
SARS-CoV-2	339	LDDKDP	344

uBCEL-N8

SARS-CoV	363	FPPTPEPKKDKKKKTDEAQLP	383
Bat-SL-CoVZXC21	363	FPPTPEPKKDKKKKADELQALP	383
Bat-SL-CoVZC45	363	FPPTPEPKKDKKKKADELQALP	383
RaTG13	363	FPPTPEPKKDKKKKADETQALP	383
SARS-CoV-2	363	FPPTPEPKKDKKKKADETQALP	383

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Supplementary Table S1. References for databases and methods utilized in this study.

Method/database	Application	Reference
AAPPred	Prediction of linear B-cell epitopes	[1]
ABCPred	Prediction of linear B-cell epitopes	[2]
Bepipred	Prediction of linear B-cell epitopes	[3]
Bepipred 2.0	Prediction of linear B-cell epitopes	[4]
BLAST	Sequence database searching	[5]
CD-HIT 4.8.1	Sequence clustering	[6]
Clustal Omega 1.2.4	Sequence alignment	[7]
COILS	Prediction of coiled-coil regions	[8]
DISOPRED 3.1	Prediction of disordered regions	[9]
Emini	Prediction of accessibility	[10]
IUPRED2	Prediction of unstructured regions	[11]
Janin	Prediction of exposed surface	[12]
Karplus	Prediction of flexibility	[13]
Kolaskar	Prediction of antigenicity	[14]
LBEEP	Prediction of linear B-cell epitopes	[15]

Mega 7	Environment for phylogeny analysis	[16]
NCBI	Sequence database for searching	[17]
Neighbor-Joining	Calculation of phylogeny	[18]
NetMHCpan EL 4.0	Prediction of HLA-1 epitopes	[19]
NetMHCIIpan 3.2	Prediction of HLA-2 epitopes	[20]
Parker	Prediction of hydrophilicity	[21]
Pellenquer	Prediction of turns	[22]
Pfam 32.0	Identification of protein domains	[23]
Phobius	Prediction of transmembrane helices	[24]
Poison correction	Calculation of phylogenetic branch lengths	[25]
Ponnuswamy	Prediction of polarity	[26,27]
PSI-PRED 4.0	Prediction of secondary structure	[27]
SDPPred	Identification of SDPs	[28]
SignalP 5.0	Prediction of signal peptide	[29]
SVMtrip	Prediction of linear B-cell epitopes	[30]

Supplementary Table S2. Linear B-cell epitopes identified in spike (S), envelope (E), membrane (M), and nucleocapsid (N) proteins of SARS-CoV and SARS-CoV-2 that have been validated *in vivo*.

Structural proteins and subunits			Epitope		Coronavirus	Experiment type	Experimental evidence	Reference		
			Location (residues)	Sequence						
S	S	S	357-365	RISNCVADY	SARS-CoV-2	<i>In silico</i>	Not tested	[31]		
			361-369	CVADYSVLY						
		1	1	369-391	YNSASFSTFKCYGVSP TK LNDLCF	SARS-CoV-2	<i>In vitro</i>	Not tested	[32]	
				424-437	KLPDDFTGCVIAWN	SARS-CoV-2	<i>In silico</i>	Not tested	[33]	
		447-458	NYNLYRLFRK							
		B	B	92-96	ASTEK	SARS-CoV-2	<i>In silico</i>	Not tested	[34]	
				110-115	LDSKTQ					
				352-357	AWNRKR					
					373-390	SFSTFKCYGVSP TKLNDL	SARS-CoV-2	<i>In silico</i>	Not tested	[35]
					434-467	GNYNKYRYLRHGKLRP FERDISNVPFSPDGKPC	SARS-CoV	<i>In vitro</i>	Sera contain neutralizing IgG antibody against peptides	[36]
					460-476	FSPDGK PCTPPALNCYW	SARS-CoV	<i>In vitro</i>	Monoclonal antibodies which neutralize the virus <i>in vitro</i>	[37]
					337-360	VYAWERKKISNCVADYS VLYNSTF	SARS-CoV	<i>In vitro</i>	Construction of mimotopes that reveal key amino acid residues involved in antibody binding	[38]
					380-399	SNVYADSFVVKGDDVRQ IAP				
					491-510	YQP YRVVLSFELLNAPA TV	SARS-CoV	<i>In vitro</i>	Epitope recognized by monoclonal antibodies with virus neutralizing activity	[39]
			437-459	NYKYRYLRHGKLRPFER DISNVP	SARS-CoV	<i>In silico</i> <i>in vitro</i> <i>in vivo</i>	Induction of cellular and humoral responses. Epitope recognized by polyclonal sera	[40]		

		336-352	SVYAWERKKISNCVADY	SARS-CoV	<i>In silico</i>	No tested	[41]
		424-435	NTRNIDATSTGN				
		442-458	YLKHGKLRPFERDISNV				
		459-470	PFSPDGKPCTPP				
		483-494	FYTTTGIGYQPY				
	424-435	NTRNIDATSTGN	SARS-CoV	<i>In silico</i> <i>in vitro</i> <i>in vivo</i>	Peptide recognized by antibodies in sera	[42]	
	536-552	GVLTPSSKRFQPFQFG	SARS-CoV	<i>In vitro</i> <i>in vivo</i>	Immunized mice produce antibodies with neutralizing activity	[43]	
	335-352	SVYAWERKKISNCVADY	SARS-CoV	<i>In vitro</i>	Neutralizing antibodies that recognized epitopes in the RBD	[44]	
	442-458	YLRHGKLRPFERDISNV					
	430-454	ATSTGNYNKYRYLRHG KLRPFERD	SARS-CoV	<i>In silico</i> <i>in vitro</i>	Antigen recognized by monoclonal antibody	[45]	
	447-458	LRPFERDISNV	SARS-CoV	<i>In silico</i> <i>in vitro</i>	Predicted epitope recognized by monoclonal antibody <i>in vitro</i>	[46]	
	343-367	KKISNCVADYSVLYNSTF FSTFKCY	SARS-CoV	<i>In vitro</i> <i>in vivo</i>	Antigen recognized by antibodies with virus neutralizing activity	[47]	
	373-390	KLNDLCFSNVYADSFVV K					
	411-428	KLPDDFMGCVLAWNTRN I					
	323-334	CPFGEVFNATKF	SARS-CoV	<i>In vitro</i>	Convalescent sera with high neutralizing activity against SARS-CoV recognized epitopes in the peptides	[48]	
467-480	CTPPALNCYWPLND						
471-503	ALNCYWPLNDYGFYTTT GIGYQPYRVVLSFEL	SARS-CoV-2	<i>In silico</i> <i>in vitro</i>	Antibodies against a peptide located in the receptor binding domain block binding and entrance of SARS-CoV into host cells <i>in vitro</i>	[49]		
287-317	DAVDCALDPLSETKCTLK SFTVEKGIYQTSN	SARS-CoV-2	<i>In silico</i>	Not tested	[50]		

		524-598	VCGPKKSTNLVKNKCVN FNFNGLTGTGVLTESNKK FLPFQQFGRDIADTTDAV RDPQTLLEILDITPCSFGGV SVI				
		815-823	RSFIEDLLF	SARS-CoV-2	<i>In silico</i>	Not tested	[31]
		1039-1047	RVDFCGKGY				
		1237-1245	MTSCCCLK				
		1264-1272	VLKGVKLHY				
		144-153	YYHKNNKSWM	SARS-CoV-2	<i>In silico</i>	Not tested	[34]
		277-283	KYNENGT				
		15-30	CVNLTTTRTQLPPAYTN	SARS-CoV-2	<i>In silico</i>	Not tested	[35]
		61-76	NVTWFHAIHVSGTNGT				
		175-201	KSGNFKHLREFVFKNKD GFLYVYKGY	SARS-CoV	<i>In vitro</i>	Sera contain neutralizing IgG against peptides	[36]
		174-195	EKSGNFKHLREFVFKNK DGFLY	SARS-CoV	<i>In silico</i> <i>in vitro</i>	Induction of cellular and humoral responses. Epitope recognized by polyclonal sera	[40]
		91-102	ATEKSNVVRGWV				
		291-308	KSFEIDKGIYQTSNFRVV	SARS-CoV	<i>In vitro</i> <i>in vivo</i>	Inoculation induces IgG and IgA in mice that exhibit potent neutralizing activity against SARS-CoV	[51]
		130-150	FELCDNPFFAVSKPMGTQ THT	SARS-CoV	<i>In vitro</i>	Antibodies block the binding of S protein to Vero E6 cells	[52]
		49-62	SDTLYLTQDLFLPF	SARS-CoV	<i>In vitro</i>	Convalescent sera with high neutralizing activity against SARS-CoV recognized epitopes in the peptides	[48]
		295-306	IDKGIYQTSNFR				
S		601-640	GTNTSNQVAVLYQDVNC TEVPVAIHADQLTPTWRV YSTGS	SARS-CoV-2	<i>In silico</i>	Not tested	[21]
		802-819	FSQILPDPSKPSKRSFIE				

2	888-909	FGAG LQIPFAMQMAYRFNG				
	814-826	KRSFIEDLLFNKV	SARS-CoV-2	<i>In silico</i>	Not tested	[53]
	560-571	LPFQQFGRDIAD	SARS-CoV-2	<i>In silico</i>	Not tested	[33]
	754-764	LQYGSFCTQLN				
	789-799	YKTPPIKDFGG				
	1139-1152	DPLQPELDSFKEEL				
	527-530	PKKS	SARS-CoV-2	<i>In silico</i>	Not tested	[34]
	601-606	GTNTSN				
	689-700	YQTQTNSPRRAR				
	771-779	EQDKNTQ				
	1154-1162	KYFKNHTSP				
	1141-1184	LQPELDSFKEELDKYFKN HTSPDVLGDISGINASV VNIQKEID	SARS-CoV-2	<i>In vitro</i>	Neutralizing antibodies recognize the S2 domain of S protein and confers protection against a wide range of SARS-CoVs	[54]
	1128-1159	DSFKEELDKYFKNHTSPD VDLGDISGINASVV				
	958-966	VLNDILSRL	SARS-CoV	<i>In vitro</i>	Sera contained neutralizing IgG antibody against peptides	[36]
	1156-1178	ASVVNIQKEIDRLNEVAK NLNES	SARS-CoV	<i>In silico</i> <i>in vitro</i> <i>in vivo</i>	Sera from vaccinated mice contain IgG that exhibit virus neutralization activity	[55]
556-568	SDFTDSVRDPKTS	SARS-CoV	<i>In silico</i> <i>in vitro</i>	Induction of cellular and humoral responses. Epitope recognized by polyclonal sera	[40]	
915-949	QIQESLTTTSTALGKLQD VVNQNAQALNTLVKQLS	SARS-CoV	<i>In silico</i>	Not tested	[41]	
1150-1184	ISGINASVVNIQKEIDRLN EVAKNLNESLIDLQE					
1065-1076	HEGKAYFPREGV	SARS-CoV	<i>In silico</i> <i>in vitro</i> <i>in vivo</i>	Peptide recognized by antibodies in sera	[56]	
520-537	IKNQCVMFNFNGLTGTGV	SARS-CoV	<i>In vitro</i>	Inoculation in mice induces IgG	[51]	

564-581	DPKTSEILDISPCSFGGV		<i>in vivo</i>	and IgA that exhibit a potent neutralizing activity against SARS-CoV	
540-554	PSSKRFQPFQQFGRD	SARS-CoV	<i>In silico in vitro</i>	Peptides at the N- the C-terminal regions of the N protein elicited strong antibody responses.	[57]
553-570	RDVSDFTDSVRDPKTSEI				
1236-1255	CKFDEDDSEPVLKGVKL HYT				
1091-	NFFSPQIITDNTFVSGNC DVVIGIINNTVYDPLQPEL DSFKEELDKYFKNHTSPD VDLGDISGINASVVNIQK EIDRLNEVAKNLNESLID LQELGKYEQYI	SARS-CoV	<i>In vitro</i>	Antibodies recognized HR2 domain and neutralize virus infection <i>in vitro</i>	[58]
1143-1157	SPDVDLGDISGINAS	SARS-CoV	<i>In vitro</i>	Antibodies recognized HR2 domain	[59]
539-559	TPSSKRFQPFQQFGRDVS DFT	SARS-CoV	<i>In vitro</i>	Four epitopes in S1 domain induce a humoral response in mice	[60]
548-567	FQQFGRDVSDFTDSVRDP KT				
583-606	VITPGTNASSEVAVLYQD VNCTDV				
607-630	STAIHADQLTPAWRIYST GNNVFQ				
927-937	GLGKLQDVVNQNGE	SARS-CoV	<i>In vitro</i>	Antibodies recognized SARS-CoV	[61]
942-951	ALNTLVKQLSSNC				
789-799	PDPLKPTKRSF	SARS-CoV	<i>In silico in vitro</i>	Epitope recognized by monoclonal antibody	[46]
540-559	PSSKRFQPFQQFGRDVSD FT	SARS-CoV	<i>In silico in vitro in vivo</i>	Antibodies without neutralizing activity but inhibit syncytia formation	[62]
731-753	CANLLLQYGSFCTQLNR ALSGIA				
786-811	QILPDPLKPTKRSFIEDLL FNKVTLA	SARS-CoV	<i>In vitro</i>	Convalescent sera recognize S2 domain	[63]
490-510	GYQPYRVVLSFELLNAP	SARS-CoV	<i>In vitro</i>	Antibodies block the binding of S	[52]

		AT			protein to Vero E6 cells	
	599-620	QDVNCTDVSTAIHADQL TPAWR	SARS-CoV	<i>In silico</i> <i>in vitro</i>	Sera from patients with SARS-CoV react with peptides	[64]
	947-965	QLSSNFGAISSVLNDILSR				
	607-627	STAIHADQLTPAWRIYST GN	SARS-CoV	<i>In vitro</i>	Peptide recognized by monoclonal antibodies which neutralize SARS-CoV	[65]
	599-620	QDVNCTDVSTAIHADQL TPAWR	SARS-CoV	<i>In silico</i> <i>in vitro</i>	Peptide recognized by antibodies in sera	[66]
	597-603	LYQDVNC	SARS-CoV	<i>In vitro</i>	Immunization with this peptide induces antibodies that enhance infection both <i>in vitro</i> and in non-human primate	[67]
	731-745	CANLLLQYGSFCTQL	SARS-CoV	<i>In vitro</i> <i>in vivo</i>	Antibodies recognize the cleavage site critical for SARS-CoV entry into host cells	[68]
	545-558	FQPFQQFGRDVSDF	SARS-CoV	<i>In vitro</i>	Convalescent sera with high neutralizing activity against SARS-CoV recognized epitopes in the peptides	[48]
	553-564	RDVSDFTDSVRD				
	651-662	PIGAGICASYHT				
	663-674	VSLLRSTSQKSI				
	695-708	AIPTNFSISITTEV				
	737-748	QYGSFCTQLNRA				
	879-890	PFAMQMAYRFNG				
	791-805	PLKPTKRSFIEDLLF	SARS-CoV	<i>In vitro</i>	Peptides recognized by convalescent sera	[69]
	1111-1130	VIGVINNTVYDPLQPELDSF	SARS-CoV	<i>In vitro</i>	Peptide recognized by neutralizing antibody that prevents viral	[70]
	60-76	SRVKNLNSSEGVPDLLV	SARS-CoV	<i>In silico</i> <i>in vitro</i>	Peptides elicit antibody responses	[57]

E	55-70	TVYVYSRVKLNLSSEG	SARS-CoV	<i>In vitro</i>	Convalescent sera with high neutralizing activity against SARS-CoV recognized epitopes in the peptides	[48]
M	1-24	MADNNGTITVEELKKLLE QWNLVI	SARS-CoV-2	<i>In silico</i>	No tested	[50]
	132-151	PLLESELVIGAVILRGHLR I				
	1-31	MADNNGTITVEELKQLLEQ WNLVIGFLFLAWI	SARS-CoV	<i>In silico</i> <i>in vitro</i>	Peptides react with convalescent sera and induce high antibody titers in rabbits	[44]
	132-161	LMESELVIGAVIIRGHLR MAGHSLGRCDIK				
	1-20	MADNNGTITVEELKQLLEQ WN	SARS-CoV	<i>In silico</i> <i>in vitro</i>	Peptide recognized by monoclonal antibody	[40]
	149-163	RMAGHSLGRCDIKDLP	SARS-CoV	<i>In silico</i> <i>in vitro</i> <i>in vivo</i>	Peptide recognized by monoclonal antibody	[42]
	173-185	PKEITVATSRTLS				
	204-221	KLNTDHAGSNDNIALLV Q				
	1-18	MADNNGTITVEELKQLLEQ	SARS-CoV	<i>In silico</i> <i>in vitro</i>	Peptides elicit an antibody response	[57]
	1-15	MADNNGTITVEELKQL	SARS-CoV	<i>In vitro</i>	Recognition by convalescent sera	[71]
	165-176	KEITVATSRTLS	SARS-CoV	<i>In vitro</i>	Convalescent sera with high neutralizing activity against SARS-CoV recognized epitopes in the peptides	[48]
5-16	GTITVEELKQLL					
137-158	LVIGAVIIRGHLRMAGHS LGR	SARS-CoV	<i>In silico</i> <i>in vitro</i>	Peptide recognized by antibodies in sera	[66]	
N	42-62	RPQGLPNNTASWFTALT QH GK	SARS-CoV-2	<i>In silico</i>	Not tested	[50]
	153-172	NNN TVLQLPQGTTLPKGF				
	355-401	NKHIDAYKTFPPTEPKKD KKKKTDEAQPLPQRQKK QPTVTLLP DM				

111-125	FYYLGTGPEASLPYG	SARS-CoV	<i>In vitro</i>	Peptide reacts with sera from mouse, rat, rabbit, civet, pig and horse	[72]
1-17	MSDNGPQSNQRSAPRIT	SARS-CoV	<i>In silico in vitro</i>	Peptides elicit antibody responses	[57]
13-30	APRITFGGPTDSTDNNQN				
25-43	TDNNQNGGRNGARPKQR RP				
38-55	PKQRRPQGLPNNIASWFT				
341-360	DDKDPQFKDNVILLNKHI DA				
356-375	KHIDAYKTFPPTPKKDK KK				
371-390	KDKKKKTDEAQPLPQRQ KKQ				
11-19	RSAPRITF	SARS-CoV	<i>In vitro</i>	Antibody binding in phage library screen	[73]
17-32	TFGGPTDSTDNNQNGG	SARS-CoV	<i>In vitro</i>	Peptides recognized by antibodies	[74]
135-150	ATEGALNTPKDHIGTR				
117-132	GPEASLPYGANKEGIV				
215-239	GGETALALLLDRLNQL SKVSGKG				
245-268	QTVTKKS EASKKPRQKRTATKQ				
274-283	AFGRRGPEQT	SARS-CoV	<i>In vitro</i>	Monoclonal antibodies recognized epitopes	[75]
286-295	NFGDQDLIRQ				
316-325	FGMSRIGMEV				
361-367	YKTFPPT				
1-30	MSDNGPQSNQRSAPRITF GGPTDSTDNNQN	SARS-CoV	<i>In vitro</i>	Peptide recognition by antibodies	[76]
81-95	PDDQIGYYRRATRRV				
86-100	GYRRATRRVRGGDG				
156-175	TVLQLPQGTTLPK				
300-320	HWPQIAQFAPSASAFFGM				

	SR				
336-350	GAIKLDDKDPQFKDN				
351-365	VILLNKHIDAYKTFP				
1-69	MSDNGPQSNQRSAPRITF GGPTDSTDNNQNGGRNG ARPKQRRPQGLPNNIASW FTALTQHGKEELRFPR	SARS-CoV	<i>In vitro</i>	Recognition by monoclonal antibodies	[77]
1-20	MSDNGPQSNQRSAPRITF GG	SARS-CoV	<i>In vitro</i>	Peptides react with convalescent sera. Antibody production after peptide immunization	[78]
150-170	RNPNNNTVLQLPQGTTLP K				
390-410	QPTVTLLPDMDDFSRQLQ N				
371-390	KDKKKKTDEAQPLPQRQ KKQ	SARS-CoV		Peptides react with convalescent sera	[79]
385-407	QRQKKQPTVTLLP DMDDFSRQ				
52-59	SWFTALTQ	SARS-CoV	<i>In silico in vitro</i>	Synthetic peptides react with convalescent sera and antisera from mice immunized with inactivate SARS-CoV	[80]
83-89	DQIGYYR				
156-166	TVLQLPQGT				
218-227	TALALLLLDR				
347-363	FKDNVILLNKHIDAYKT				
379-385	EAQPLPQ				
389-398	KQPTVTLLPA				
66-87	QLPQGTTLPKGFYAEGSR GGSQ	SARS-CoV	<i>In silico in vitro</i>	Peptide recognized by antibodies in sera	[66]
371-390	KDKKKKTDEAQPLPQRQ KKQ				

Abbreviations:

^a IEDB: Immune Epitope Database

^b PISA: Proteins, Interfaces, Structures and Assemblies

^c ViPR: Virus Pathogen Database and Analysis Resource

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