

- A. **GGGCCC**TCATGGCGGACAGCAACGGTACCATCACCGTGGAGGAACTGAAGAAACTGCTGGAGCA
GTGGAACCTGGTTATCGGCTTCCTGTTTCTGACCTGGATTGCTGCAATTCGCGTACGCGAACC
GTAACCGTTTTCTGTACATCATCAAGCTGATTTTCTGTGGCTGCTGTGGCCGGTGACCCTGGCGTGC
TTTGTGCTGGCGGCGGTTTACCGTATCAACTGGATTACCGGTGGCATCGCGATTGCGATGGCGTGCC
TGGTTGGTCTGATGTGGCTGAGCTATTTATCGCGAGCTTCCGTCTGTTTGC GCGTACCCGTAGCATG
TGGAGCTTCAACCCGGAGACCAACATCCTGCTGAACGTGCCGCTGCACGGCACCATTTCTGACCCGTC
CGCTGCTGGAGAGCGAACTGGTGATCGGTGCGGTTATTCTGCGTGGCCACCTGCGTATTGCGGGTCA
CCACCTGGGCCGTTGCGACATCAAGGATCTGCCGAAAGAAATTACCGTGGCGACCAGCCGTACCCTG
AGCTACTATAAGCTGGGTGCGAGCCAGCGTGTTGCGGGTGACAGCGGTTTTGCGGCGTACAGCCGTT
ATCGTATCGGCAACTACAACTGAACACCGACCACAGCAGCAGCAGCGATAACATTGCGCTGCTGGTT
CAG**GGGCCC**
- B. MADSNGTITVEELKKLLEQWNLVIGFLFTWICLLQFAYANRNRFLYIIKLIFLWLLWPVTLACFVLAAYRIN
WITGGIAIAMACLVGLMWLSYFIASFRLFARTRSMWSFNPETNILLNVPLHGTILTRPLLESELVIGAVILRG
HLRIAGHHLGRCDIKDLPKEITVATSRTLSTYKLGASQRVAGDSGFAAYSRYRIGNYKLNTDHSSSDNIA
LLVQ
- C. nCoV-M NotI-F: 5'-ATATGCGGCCGCTCATGGCGGACAGCAACGGT-3'
nCoV-M ApaI-R: 5'-ATATGGGCCCTGAACCAGCAGCGCAAT-3'

Supplemental FigureS1. Nucleotide sequence and cloning of codon optimized synthetic gene of SARS-CoV-2 membrane (M) protein. The codon optimized synthetic membrane (M) protein gene of SARS-CoV-2 Wuhan-Hu-1 strain (NCBI Reference Sequence: NC_045512.2) was chemically synthesized and cloned in the plasmid pET-32a(+)-M protein (kindly provided by Dr. Shih-Wein Li at Institut Pasteur of Shanghai, Chinese Academy of Sciences). Nucleotide and deduced amino acid sequences of SARS-CoV-2 M protein gene were listed (A, B), and used as the template for PCR amplification with a specific primer pair (C). Finally, the codon optimized synthetic gene of SARS-CoV-2 M protein was cloned into the restriction enzyme sites NotI/ApaI of pcDNA3.1/HisC-T2A-F2A.

- A. **ATCGAT**ATGTACTCATTTCGTTTCGGAAGAGACAGGTACGTTAATAGTTAATAGCGTACTTCTTTTCTTGCTTCGTGGTATTCTTGCTAGTTACACTAGCCATCCTTACTGCGCTTCGATTGTGTGCGTACTGCTGCAATATTGTTAACGTGAGTCTCGTAAACCTTCTTTTACGTTTACTCTCGTGTTAAAAATCTGAATTCTTCTAGAGTTCCTGATCTTCTGGTC**ATCGAT**
- B. MYSFVSEETGLIVNSVLLFLAFVVFLVTLAILTALRLCAYCCNIVNVSLVKPSFYVYSRVKNLNSSRPDL
LV
- C. nCoV-E ClaI-F: 5'-ATATATCGATATGTACTCATTTCGTTTCGG-3'
nCoV-E ClaI-R: 5'-ATATATCGATGACCAGAAGATCAGGAAC-3'

Supplemental FigureS2. Nucleotide/amino acid sequences and cloning of SARS-CoV-2 envelope (E) protein gene. The envelope (E) protein gene of SARS-CoV-2 Wuhan-Hu-1 strain (NCBI Reference Sequence: NC_045512.2) was chemically synthesized and cloned in the plasmid pCAG.2-HA-SARS-CoV-2-E-HA (kindly provided by Professor Che Alex Ma at Genomics Research Center, Academia Sinica, Taiwan). Nucleotide and deduced amino acid sequences of SARS-CoV-2 E protein gene were listed (A, B), and used as the template for PCR amplification with a specific primer pair (C). Finally, the codon optimized synthetic gene of SARS-CoV-2 E protein was cloned into the restriction enzyme site ClaI of pcDNA3.1/HisC-T2A-F2A-nCoV M.

B. MFVFLVLLPLVSSQCVNLTTTRTQLPPAYTNSFTRGVYPDKVFRSSVLHSTQDLFLPFFSNVTWFHAIHVS GTNGTKRFDNPVLPFNDGVYFASTEKSNIIRGWIFGTTLD SKT
QSSLIVNNATNVV/VIKVCE/FQFCNDPFLGVYYHKNNKSWMESEFRVYSSANNCTFEYVSQPFMDLEGKQGNFKNREFVFKNIDGYFKIYSKHTPINLVRDLPQGFSALEPLV
DLPIGINITRFQTLALLHRSYLT PGDSSSGW TAGAAAYYVG YLQPRTFLLKY NENG TITDAVDCALDPLSETKCTLKSFTVEKGIYQTSNFRVQPTESIVRFPNITNLCPFGEVFN
ATRFASVYAWNRKRISNCVADY SVLYNSASFSTFKCYGVSP TKLNDLCFTNVYADSFVIRGDEVRQIAPGQTGKIADYNYKLPDDFTGCVIAWNSNNLDSKVGGNYNYLYRLF
RKSNLKPFERDISTEIIYQAGSTPCNGVEGFNCYFPLQSYGFGQPTNGVGYPYRVVLSFELLHAPATVCGPKKSTNLVKNCVNFNFNGLTGQVLTESNKKYLPFQQFGGRDI
ADTTDAVRDPQTLEILDITPCSF GGVSVITPGTNTSNQVAVLYQDVNCTEYVPVAIHADQLPTWRVYSTGSNVFQTRAGCLIGA EHVNSYECDIPGAGIKASQQTQTSNPRR
ARSVASQSIIAYTMSLGAENSVAYSNN SIAIPTNFTISV TTEILPVSMKTCTSV DCTMYICGDSTEC SNLLQLQYGSFCTQLNRALT GIAVEQDKNTQEVFAIGKQKIYKTPPIKDFGGF
NFSQILPDPSKPSKRSFIEDLLFNKVT LADAGFIKQYGDCLGDIAARDLICAQKFNGLT VLPPLLTDEMAIAYTSALLAGTITSGWTFGAGAALQIPFAMQMAYRFNGIGVTQNVL
YENQKLIANQFN SIAIGKIQDSL S STASALGKLQDVVNQNAQAALNTLVKQLSSNFGAISSVLNDILSRLDKVEAEVQIDRLITGR LQSLQTYVTQQLIRAAEIRASANLAATKMSEC
VLGQSKRVDFCKGKYHLSMFPQSAPHGVVFLHVTYVPAQEKNFTTAPAICH DGKAHFPREGV FVSNGTHWFVTQRNFYEPQIITDNTFVSGNCDVVIGIVNNTVYDPLQPE
LDSFPEELDKYFKNHTSPD VDLGDISGINASVVNIQKEIDRLNEVAKNLN ES LIDLQELGKYEQYIKWPWYIWLGFIAGLIAIVMVTIMLCCMTSCC SCLKGCCSCGSCCKFDED
DSEPKVLKGVKLHYHT

Supplemental Figure S3. Nucleotide/amino acid sequences and cloning of SARS-CoV-2 Wuhan-Hu-1 spike (S) protein gene. Spike (S) protein gene of SARS-CoV-2 Wuhan-Hu-1 strain (NCBI Reference Sequence: NC_045512.2) was chemically synthesized and cloned in the plasmid pCMV3-2019-nCoV-Spike (S1+S2)-long (kindly provided by Professor Che Alex Ma at Genomics Research Center, Academia Sinica, Taiwan). Nucleotide and deduced amino acid sequences of SARS-CoV-2 S protein gene were listed (A, B), and used as the template for PCR amplification with a specific primer pair (C). Finally, the SARS-CoV-2 Wuhan-Hu-1 S protein gene was cloned into the restriction enzyme sites KpnI/AscI of pcDNA3.1/HisC-T2A-F2A-nCoV E-nCoV M.