

Pseudovirus-SARS-CoV-2-S

Cat# PV025

Store at -20°C for 6 months

INFORMATION

DESCRIPTION:	The partial S gene coding sequence were cloned into retroviral vector by chemical synthesis to obtain SARS-CoV-2-S pseudovirus. The pseudovirus are prepared with 293T and concentrate and purify by the ultracentrifuge. Viral envelope of the SARS-CoV-2-S pseudovirus include the partial S gene and coding sequence, using in experiments which are related to RNA extraction and become the positive control of qPCR.
APPLICATIONS:	Research Recommended amount: 50-100 µl / reaction. According to experimental conditions, it can be adjusted.
Main ingredient:	glucose 、KH ₂ PO ₄ 、Na ₂ HPO ₄ 、NaCl 、KCl 、SARS-CoV-2-S pseudovirus
FORMULATION:	Liquid
STORAGE & STABILITY:	The product can be stored at -20°C or below for 6 months. Avoid repeated freezing and thawing cycles.
PRODUCT:	>1 x 10 ⁷ copy/ml in 1 ml

PROTOCOL

1. The pseudovirus are took from -20°C and melted in 4°C or ice bath. You can execute the related experiments when the pseudovirus completely melted.
2. Pseudovirus Inactivate (optional operation): Extract the sufficient pseudovirus in eppendroft at 56°C for 30 min that need to operation in Biological safety cabinet (BSC).
3. Pseudovirus extraction and qPCR detection (materials prepare by yourself): Perform relevant experimental operations in accordance with the instructions of the RNA extraction kit and qPCR detection kit.

4. qPCR Detection (materials prepare by yourself): The qPCR detection was performed after pseudovirus RNA was synthesized cDNA by RT-PCR.
5. Supplementary: The produce maybe a small amount of plasmid DNA remain. If the experiments need high purification, it can use DNase-DEPC water by RNA extraction. You can add EDTA (final con. 5 mM) for 10 min at 75°C to inactivate DNase (optional operation).

NOTE

1. Freezing and thawing will reduce the stability of the pseudovirus, which will affect the effect of RNA extraction and results of qPCR detection. Avoid repeated freezing and thawing when using.
2. Virus inactivation treatment may lead to RNA explanation, please choose according to the needs of the experiment.
3. If you need to dilute the pseudovirus, you can choose the 1X PBS or physiological saline (0.9% NaCl).
4. Please rinse immediately with plenty of water when the pseudovirus accidentally splashed on eyes, skin or other body parts.
5. According to the medical waste disposal specifications, the experimental waste generated by using the pseudovirus needs to perform high pressure thermal sterilization process.

SEQUENCE INFORMATION

S Gene 1.3kb

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GTTCTTGTTAACTAAACGAACAATGTTTGTTCCTTGTTT  
TATTGCCACTAGTCTCTAGTCAGTGTGTTAATCTTACAACCAGA  
ACTCAATTACCCCTGCATACACTAATTCTTTCACACGTGGTGT  
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CATGCTATACATGTCTCTGGGACCAATGGTACTAAGAGGTTTGA  
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CTGAGAAGTCTAACATAATAAGAGGCTGGATTTTTGGTACTACT  
TTAGATTGAAGACCCAGTCCCTACTTATTGTTAATAACGCTAC  
TAATGTTGTTATTAAGTCTGTGAATTTCAATTTGTAATGATC  
CATTTTTGGGTGTTTATTACCACAAAAACAACAAAAGTTGGATG  
GAAAGTGAGTTCAGAGTTTATTCTAGTGCGAATAATTGCACTTT  
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AGTGCGTGATCTCCCTCAGGGTTTTTCGGCTTTAGAACCATTGG
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AACCTAGGACTTTTCTATTAATAATAATGAAAATGGAACCATT
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CCTAATATTACAACTTGTGCCCTTTTGGTGAAGTTTTAACGC
CACCAGATTTGCATCTGTTTATGCTTGGAACAGGAAGAGAATCA
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GAGGTGATGAAGTCAGACAAATCGCTCCAGGGCAAACCTGAAAAG
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CGTTATAGCTTGGAAATCTAACAATCTTGATTCTAAGGTTGGTG
GTAATTATAA

PRODUCT USE LIMITATION

These products are intended for research use only.