



VERSION 2

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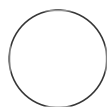
**MANUSCRIPT CITATION:**

Generic novel system for genomic characterization of Respiratory Syncytial Virus obtaining whole genome sequencing and a full-length G and F sequences.

## RSVAB WGS and GF protocols V.2

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### ABSTRACT

This SOP describes the procedure for generating cDNA from RSV viral nucleic acid extracts and subsequently producing amplicons tiling the viral genome using. We propose two systems for genomic characterization of RSV. First, a novel RSV amplicon-based system for WGS, and second, a method focused on obtaining the specific sequences of the main antigens, G and F.

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**Protocol status:** Working  
We use this protocol and it's working

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**PROTOCOL integer ID:**  
91949

## dsCDNA generation:

1

During this step three master mixes will be prepared: MMI, MMII and MMIII.

Materials: **Kit Superscript III First Strand (Invitrogen)**  
**100% DMSO**  
**RNAseH (Invitrogen)**  
**Klenow fragment 3' ->5' exo (New England Biolabs)**  
**Primer FR26RV-N : 5'GCC GGA GCT CTG CAG ATA TCNNNNNN 3'**

### Note

This step must be performed in a RNase free, pre-PCR environment in which post PCR RSV amplicons are not present, to minimise risk of sample contamination.

### CITATION

Díez-Fuertes F, Iglesias-Caballero M, García-Pérez J, Monzón S, Jiménez P, Varona S, Cuesta I, Zaballos Á, Jiménez M, Checa L, Pozo F, Pérez-Olmeda M, Thomson MM, Alcamí J, Casas I (2021). A Founder Effect Led Early SARS-CoV-2 Transmission in Spain..

LINK

<https://doi.org/10.1128/JVI.01583-20>

## 2 MMI Preparation:

A	B
FR26RV-N (10uM)	2
DMSO	0,5
Total	2,5 ul

Mix thoroughly by vortexing.

## 3 MMII Preparation:

A	B
10x First Strand Buffer	2
DTT 100 mM	2
MgCl <sub>2</sub> 25mM	4
dNTPs	1
RNaseOUT	0,5
SSIII RT	0,5
Total	10 ul

Kit Superscript III First Strand (Invitrogen)

## 4 MMIII Preparation:

A	B
Klenow 5'-3'	1
RNaseH	0,5
Total	1,5 ul

## 5 Defrost extracted RNA.

Maintain **on ice** the MMI,MMII and MMIII mixes.

## 6 MMI Amplification:


Add  5  $\mu\text{L}$   Sample in MMI mix

Place the tube on a thermocycler and run the following program:

A	B
65°C	5 min
4°C	2 min

Briefly tube centrifugation

## 7 MMII Amplification:


Addition of  10  $\mu\text{L}$  from MMII in the tube with the MMI and the viral extraction.

Place the tube on a thermocycler and run the following program:

A	B
25°C	10 min
50°C	50 min
85°C	10 min
4°C	$\infty$

Briefly tube centrifugation

## 8 MMIII Amplification:

Addition of  1.5  $\mu\text{L}$  of MMIII into the tube with the previous mixes and the viral extraction

Place the tube on a thermocycler and run the following program:

A	B
37°C	60 min
75°C	15 min

Briefly tube centrifugation

9 **STOP POINT:** cDNA can be stored at 4°C (same day) or -20°C (up to a week).

## RSVAB WGS protocol

10

Materials:

**2x MyTaqRed mix (Bioline)**

**Primers:**

A	B
Primer ID	Sequence (5'-3')
<b>Mix 1</b>	
RSVCombinitial	ACGCGAAAAAATGCGTACWACA
RSVWGS4R	CATGWTGWYTTATTTGCCCC
RSVWGS2F	CACTWACAATATGGGTGCC
RSVWGS1R	TCCATKGTTATTTGCCCC
RSVWGS3.2F	ACATGGAAAGAYATYAGCC
RSVWGS2R.2	CRTTYCTTAARGTRGGCC
RSVWGS3.2R	TTGCATCTGTAGCAGGAATGG
OG1-21	GGGGCAAATGCAACCATGTCC
RSVGF-R	TTCGYGACATATTTGCCCC
RSVCombending	ACGAGAAAAAAGTGTCAAAAATAA
<b>Mix 2</b>	
RSVCombinitial	ACGCGAAAAAATGCGTACWACA
RSVWGS1R	TCCATKGTTATTTGCCCC
RSVWGS8R.2	TCMAWYTCWGCAGCTCC
RSVWGS5R	CAAACATTTAATCTRCTAAGGC

A	B
RSVWGS6F	TTATAYAGATATCAYATGGGTGG
RSVWGS6R	CCCTCTCCCAATCTTTTTC
RSVWGS9F	GARCAACTCAAAGAAAATGG
RSVWGS9R	AYTGRAACATRGGCACCC
RSVCombending	ACGAGAAAAAAGTGTCAAAAATAA

#### Note


The protocol is based in the RSV genome amplification in two separate mixes with an unique amplification program. The mixes that will be mixed at the end of cycling.

## 11 Preparation of RSV Amplification Mix 1:

A	B
MyTaq Red 2x	15
H2O	8,4
RSV Combinitial	0,2
RSVWGS1R (5uM)	0,2
RSVWGS2F (5 uM)	0,2
RSVWGSW2R.2 (5 uM)	0,2
RSVWGS4R (5 uM)	0,2
RSVWGS3.F (5 uM)	0,2
RSVWGS3.2R (5 uM)	0,2
OG1-21	0,2
RSVGF-R	0,2
RSV Combending	0,2
Total	25

## 12 Preparation of RSV Amplification Mix 2:

A	B
2x My Taq Red	15
H2O	8,6
RSV Combinatorial (10uM)	0,2
RSVWGS1R (10 uM)	0,2
RSVWGS5R (10 uM)	0,2
RSVWGS8R.2 (10 uM)	0,2
RSVWGS6F (10 uM)	0,2
RSVWGS6R (10 uM)	0,2
RSVWGS9F	0,2
RSVWGS9R	0,2
RSV Combending	0,2
Total	25 ul

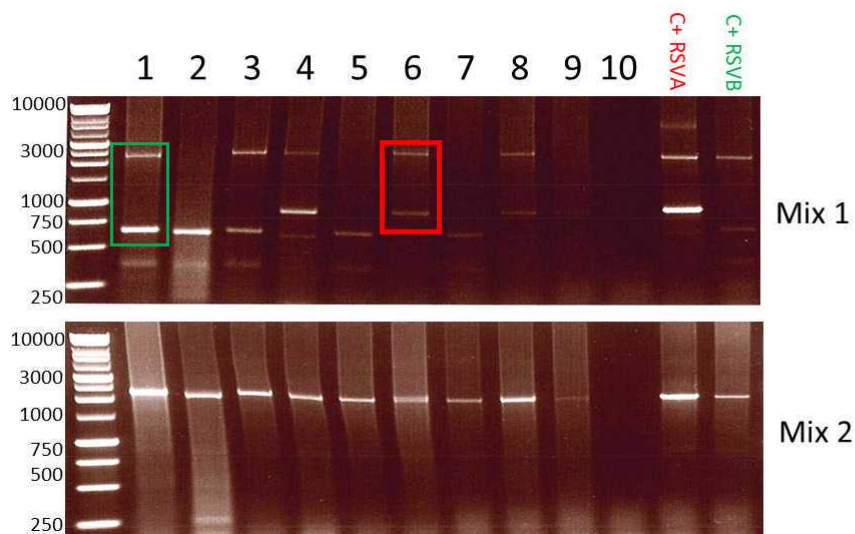
**13** Addition of  5 µL of the previous prepared double stranded cDNA on each mix.

**14 Amplification protocol:**

A	B	C
95°C	1 min	
95°C	30 seg	x45
55°C	8 min	
72°C	2 min	
72°C	5 min	
12°C	∞	

**15** To assess PCR performance, the amplicons can be loaded onto a 1% agarose gel for electrophoresis.

## Expected result



- 16 Finally, mix in one single tube both mixes and proceed to purification and library preparation.

## RSVAB GF protocol starting from ds cDNA

- 17 Due to the significance of achieving accurate RSV genomic characterization, it was developed the RSVAB-GF PCR to complement the genomic coverage of both antigenic major proteins in cases where WGS encounters difficulties, and to provide a simpler and more cost-effective method of obtaining the sequences of both antigens.


- 18 Materials:  
**2x MyTaqRed mix (Bioline)**  
**Primers:**

A	B
OG1-21	GGGGCAAATGCAACCATGTCC
RSVGF-R	TTCGYGACATATTTGCCCC



## 19 Preparation of cDNA GF amplification mix:

A	B
H2O	5,5
2X MyTaqRed	12,5
OG1-21 (10 uM)	1
RSVGF-R (10 uM)	1
Total	20 ul

20 Addition of  5 µL of the previous prepared double stranded cDNA on the mix.

## 21 Amplification protocol cDNA GF:

A	B	C
95°C	1 min	
95°C	30 seg	x35
60°C	3 min	
72°C	2 min	
72°C	5 min	
12 °C	∞	

## RSVAB GF protocol starting from viral extraction

22 Materials:  
**Qiagen OneStep RT-PCR kit.**  
**Glycerolised 1% H2O**

## 23 Preparation of GF amplification mix:

A	B
H2Ogly	10
5xQ PCR MM	6
dNTPs	1
OG1-21 (10uM)	1
RSVGF-R (10 uM)	1
RT-PCR mix	1
Total	20 ul

24 Addition of  10 µL of the viral extraction

25 **Amplification protocol GF:**

A	B	C
48°C	60 min	
95°C	15 min	
95°C	30 seg	x 35
60°C	3 min	
72°C	2 min	
72°C	5 min	
12°C	∞	