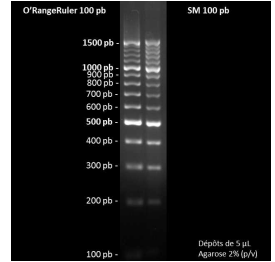


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🌐 Home made SM 100bp DNA ladder for agarose gel

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Protocol status: Working

We use this protocol and it's working

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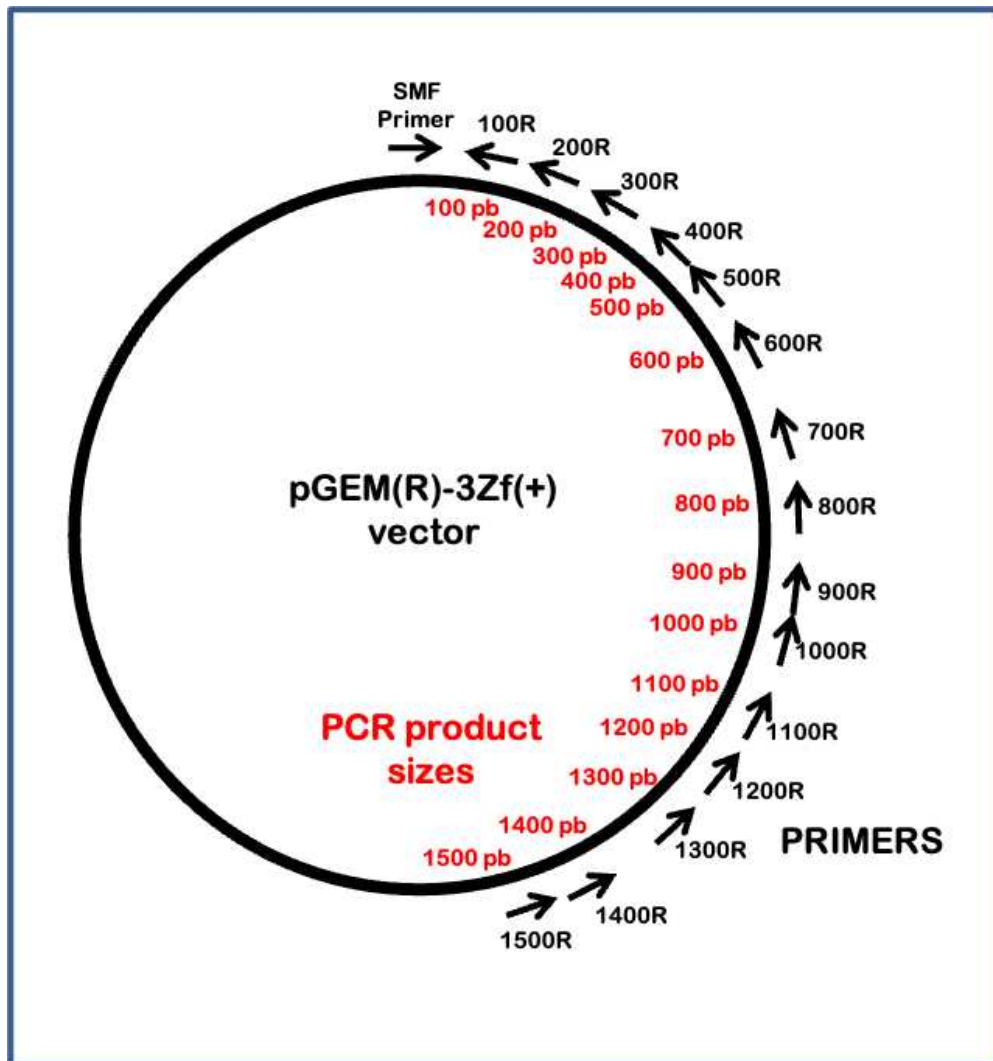
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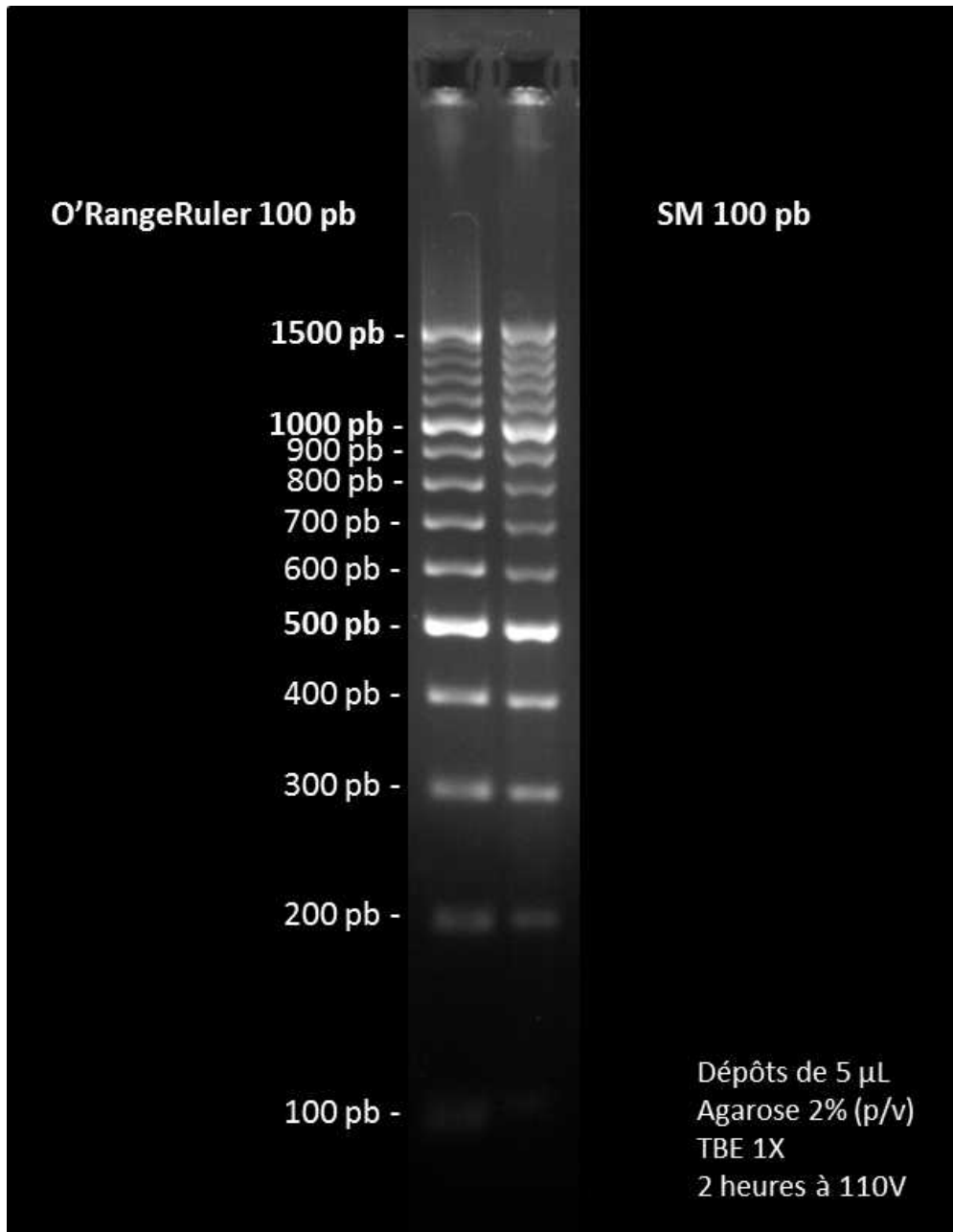
Abstract

The SM 100pb DNA Ladder is a standard size marker equivalent to the Fisher's O'RangeRuler® 100 pb DNA Ladder (#SM0623). The SM 100pb Ladder allows to determine the size of double-stranded DNA fragments between 100 bp and 1500 bp and it is composed of 15 double-stranded DNA fragments of 100, 200, 300, 400, 500, 600, 700, 800, 900, 1000, 1100, 1200, 1300, 1400 and 1500 bp. Like the O'RangeRuler 100 bp, the SM 100 bp gives more intense bands at 500, 1000 and 1500 bp.

At the end, the SM 100 pb DNA Ladder is 25 times cheaper than the O'RangeRuler 100 pb.



Map of the universal cloning vector pGEM®-3Zf(+) showing the position of the 16 primers



Comparaison between O'rangeRuler 100pb (Thermofisher) and SM100bp ladder on 2% agarose gel

Protocol materials

 GoTaq® Flexi DNA Polymerase **Promega Catalog #M8291**

 pGEM®-3Zf() Vectors **Promega Catalog #P2271**



Primers and pGEM®-3Zf(+) Vectors preparation

1 Oligo primers dilution

in 16  1.5 mL microtubes, dilute 1:10 each oligo primer at 100µM to final concentration 10µM, **see the SM100pb_primers.xlsx file below.**

 50 µL oligo primer (100µM)

 450 µL nuclease free water

Homogenize and store at  4 °C (or at  -20 °C for a long-term storage).

SM100pb oligo primers sequences (salt free purification):



SM100pb_primers.xlsx 12.8KB

2 pGEM®-3Zf(+) Vectors dilution

In  1.5 mL microtube, dilute 1:100 pGEM®-3Zf(+) vector at 1µg/µL to final concentration 10ng/µL.

 5 µL pGEM®-3Zf(+) (1µg/µL)

 495 µL nuclease free water

Homogenize and store at  4 °C (or at  -20 °C for a long-term storage).



pGEM®-3Zf() Vectors **Promega Catalog #P2271**

PCR amplification to generate 100pb-1500pb double-stranded DNA fragments

10s

3 PCR mix preparation

10s

SM100pb is produced using **28 PCRs amplifications** in a total of  50 µL reaction volume:

1 x  50 µL for 600pb, 700pb, 800pb, 900pb, 1100pb, 1200pb, 1300pb and 1400pb fragments

2 x  50 µL for 200pb, 300pb and 400pb fragments

3 x  50 µL for 1000pb and 1500pb fragments

4 x  50 µL for 100pb and 500pb fragments

Defreeze and vortex all reagents, except enzymes (stored at -20°C), for approximately 00:00:05

Spin down all reagents for approximately 00:00:05 and place On ice.

In 1.5 μL microtube, prepare the PCR mix according to the following table :

A	B	C	D	E
	Initial concentration	Final concentration	n=1	n=28
SMF forward primer	10 μM	400mM	2 μL	56μL
Green GoTaq buffer	5X	1X	10 μL	280μL
MgCl ₂	25mM	1mM	2 μL	56μL
pGEM®-3Zf(+) vecto	10ng/ μL	20ng	2 μL	56μL
dNTP mix	2.5mM	150 μM each	3 μL	84μL
GoTaq polymerase	5 u/ μL	1.75U	0.35 μL	9.8μL
nuclease free water			28.65 μL	802.2μL
TOTAL			48 μL	1344μL

PCR mix composition

 GoTaq® Flexi DNA Polymerase **Promega Catalog #M8291**

4 Reverse primers and mix combinaison

Defreeze and vortex all the 15 reverse oligo primers at 10 μM

In a 96-well plate PCR, transfer 2 μL of each reverse oligo primers (10 μM) according to the following map :

A	B	C	D	E	F	G	H
600R	200R	1000R	500R				
700R	200R	1500R	500R				
800R	300R	1500R	500R				

	A	B	C	D	E	F	G	H
	900R	300R	1500R	500R				
	1100R	400R	100R					
	1200R	400R	100R					
	1300R	1000R	100R					
	1400R	1000R	100R					

Map of the PCR plate

Vortex and spin down the PCR mix tube, transfer  48 µL in each of the 28 wells.
Seal the PCR plate.

In thermocycler, run PCR amplification with cycles follows:

	A	B	C	D
	Cycles step	Temperature	Time	Cycles
	Initial denaturation	94°C	5 min	1
	Denaturation	94°C	30 sec	40
	Annealing	60°C	30 sec	40
	Extension	72°C	30 sec	40
	Final extension	72°C	60 min	1
	Hold	4°C		

PCR program

After PCR, pool and dilute 1:2 the PCR amplification

In a  5 mL tube:

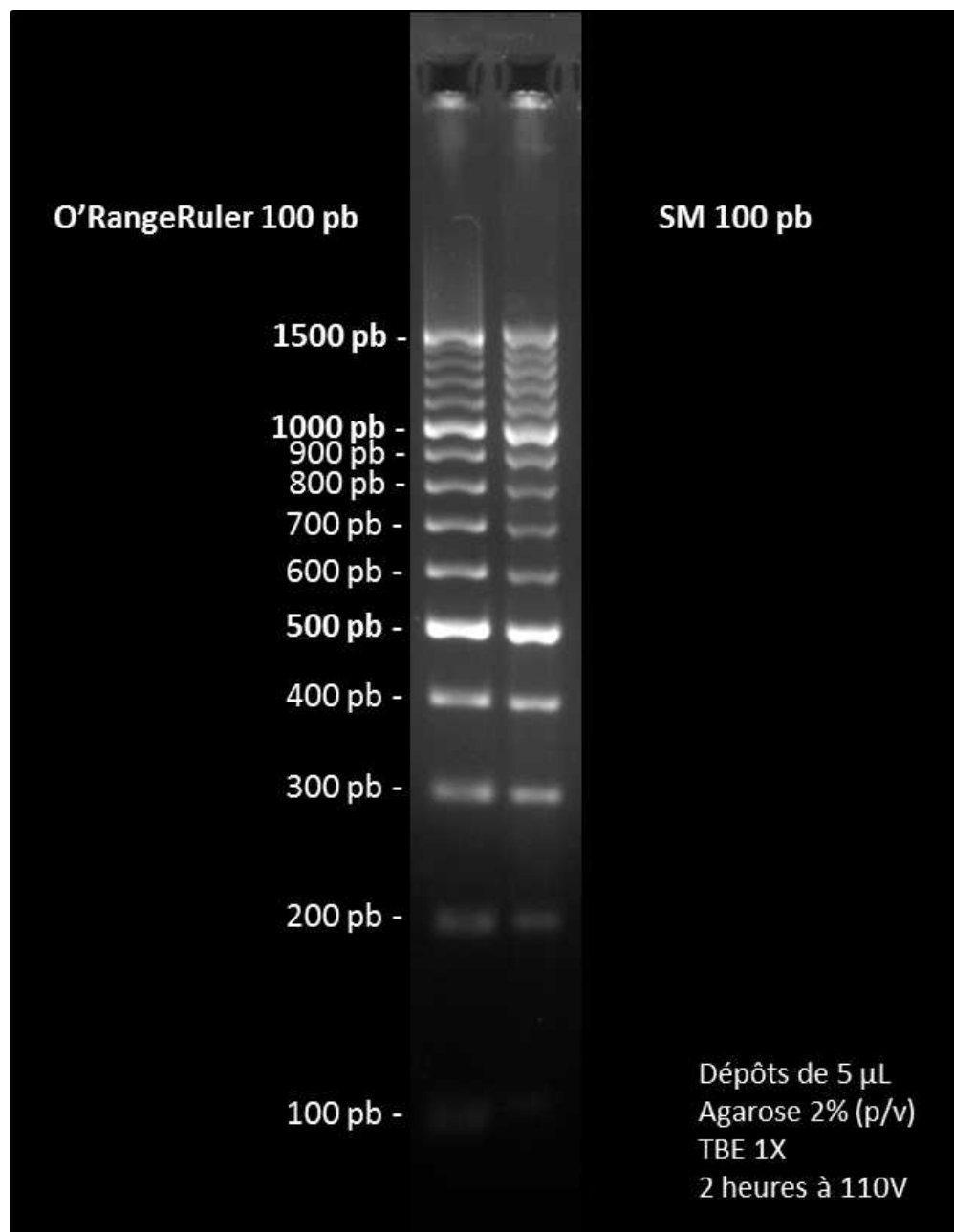
 1400 µL of PCR amplification (28 x  50 µL)

 420 µL 5X green GoTag buffer

 980 µL nuclease free water

Homogenize and store at  4 °C (or at  -20 °C for a long-term storage).

Load 5 μL to 10 μL per line in a agarose gel.



Comparaison between O'rangeRuler 100pb (Thermofisher)
and SM100bp ladder on 2% agarose gel