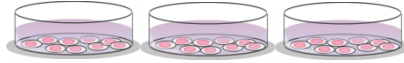


To find the ideal conditions, **ViroMag** must be tested at ratio **1.5 to 6  $\mu\text{L}$  /  $10^5$  infectious viral particles**. Adapt your MOI depending of your viral vector and the type of cells used.

**Seed cells to be at 70% confluent the day of transfection**

1



**Prepare 3 identical tubes of viral particles (usually MOI 1)**

2



**96 well plate**

For  $1 \times 10^4$  cells per well,  
dilute  $1 \times 10^4$  infectious units in  
 $50 \mu\text{L} \times 3$

**24 well plate**

For  $1 \times 10^5$  cells per well,  
dilute  $1 \times 10^5$  infectious units in  
 $100 \mu\text{L} \times 3$

**6 well plate**

For  $5 \times 10^5$  cells per well,  
dilute  $5 \times 10^5$  infectious units in  
 $200 \mu\text{L} \times 3$

**Prepare 3 tubes of ViroMag (with different amounts of magnetic beads)**

3



**96 well plate**

$0.15 \mu\text{L} / 0.3 \mu\text{L} / 0.6 \mu\text{L}$   
in an empty microtube

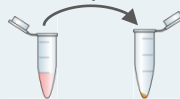
**24 well plate**

$1.5 \mu\text{L} / 3 \mu\text{L} / 6 \mu\text{L}$   
in an empty microtube

**6 well plate**

$7.5 \mu\text{L} / 15 \mu\text{L} / 30 \mu\text{L}$   
in an empty microtube

**Mix each tube of viral particles (step 2) to each tube of ViroMag (step 3)**



4

**96 well plate**

**24 well plate**

**6 well plate**

96 well plate			24 well plate			6 well plate		
Infectious units		ViroMag	Infectious units		ViroMag	Infectious units		ViroMag
$1 \times 10^4$	+	$0.15 \mu\text{L}$	$1 \times 10^5$	+	$1.5 \mu\text{L}$	$5 \times 10^5$	+	$7.5 \mu\text{L}$
$1 \times 10^4$	+	$0.3 \mu\text{L}$	$1 \times 10^5$	+	$3 \mu\text{L}$	$5 \times 10^5$	+	$15 \mu\text{L}$
$1 \times 10^4$	+	$0.6 \mu\text{L}$	$1 \times 10^5$	+	$6 \mu\text{L}$	$5 \times 10^5$	+	$30 \mu\text{L}$

**Incubate 20 min at room temperature**

5



**Distribute each mix onto the cells and Incubate the cells 20 min on the magnetic plate**

6



**Remove the cells from the magnetic plate and incubate cells for 24 to 72h at  $37^\circ\text{C}$  until evaluation of transgene expression**

7



**Choose the best ratio DNA:ViroMag**

8



**NOTES:** (1) 24H before transfection seed the cells in 6 wells of a 96-well plate, 3 wells of a 24-well plate or 3 wells of a 6-well plate in respectively 150  $\mu$ L, 400  $\mu$ L and 2 mL of complete culture medium. (2) Allow reagents to reach RT and gently vortex them before forming complexes . (3) DMEM without serum & supplement is recommended for complexes preparation. (4) For doses less than 1 $\mu$ L, dilute your reagent with deionized water.

The complete instruction manual is accessible on OZ Biosciences website:  
[www.ozbiosciences.com](http://www.ozbiosciences.com)

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