Tinzyme Co., Limited

Email: sales@tinzyme.com Website: www.tinzyme.com
Tel: +86-755-86134126 WhatsApp/Facebook/Twitter: +86-189-22896756

EZ-AV Transduction kit

Product Number: TR3011

Shipping and Storage

This product is transported at room temperature and stored for a long time at 4℃, with a validity period of 12 months.

This product is safe to use and no biological or chemical toxicity has been found. If accidentally contaminated, rinse with clean water.

Description

EZ-AV Transmission Kit is an enhancer developed and synthesized by our company for adenovirus infection, which is synthesized using nanotechnology. The EZ-AV Transmission kit enriches the virus on the cell surface through physical interactions to enhance infection efficiency. Due to the application of nanotechnology, the EZ-AV Transmission kit significantly improves the efficiency of adenovirus infection while minimizing cell toxicity and not interfering with cellular physiological functions.

Note

1. After using the EZ-AV Transmission kit for enhancement, the degree of cell fusion during infection has an impact on the infection effect. It is recommended to conduct infection experiments when the degree of cell fusion is around 30-50%. If it is a suspended cell, the usual density of cells infected with adenovirus can be used.
2. Before using the EZ-AV Transmission kit enhancer, the optimal MOI (Multiplicity of Infection) of infected cells should be determined, which refers to the average number of viruses infected per cell. Generally, the higher the MOI, the more viruses integrate into chromosomes and the higher the surface expression of the target protein. Different cells have their optimal MOI values. We need to conduct a MOI gradient experiment first to determine the optimal MOI value for infected cells. The application of enhancers on the basis of the optimal MOI value has a better effect.
3. The dilution solution for EZ-AV Transmission kit and virus is a serum-free liquid that is consistent with the cell base culture medium, such as serum-free DMEM or 1640.
4. The relationship between EZ-AV Transmission kit and infection efficiency. Under other fixed conditions, the larger the dosage of enhancer, the higher the infection efficiency. Generally, the dosage of enhancer can be used 1-3 times the recommended dosage. However, excessive use of enhancers may lead to cytotoxicity.

Protocol

1. Cell laying one day in advance

   It is advisable to plant cells in advance, with a cell fusion degree of around 30-50% during infection (adjust suspension cells as needed, do not use high cell density).

2. Infect

   2.1. Dilute the EZ-AV Transmission kit with serum-free diluent (see table below for dosage), mix thoroughly, and prepare an enhancer diluent.

   2.2. Dilute the virus concentrate with serum-free diluent (see table below for dosage), mix thoroughly, and prepare the virus diluent.

   Note: Due to different virus concentrations, the specific amount of virus concentrate may be based on conventional dosage. It is recommended to conduct gradient experiments in combination with enhancers.

   2.3. Mix the diluent of the enhancer and virus thoroughly, and let it stand at 4 ℃ for 15 minutes.

   2.4. Add the above mixture to cells containing complete culture medium, culture at 37 ℃ for 8 hours, and observe the cell state. If there is no significant change, do not change the culture medium and continue to culture. Due to the rapid infection of adenovirus, the expression can be observed at 12, 24, and 48 hours respectively.

Table 1 Recommended dosages for different cell culture containers

Cell culture container	Surface area (cm^2)	Ratio of surface area to 24 well	Quantity of EZ-AV Transduction per Hole	Dosage of diluent per well	Total amount of culture medium per well
96-well	0.3	0.2	0.5-1.5μl	10μl	100μl
48-well	0.7	0.4	1-3μl	15μl	200μl
24-well	1.9	1	2.5-7.5μl	25μl	500μl
12-well	3.8	2	5-15μl	25μl	1ml
6-well/35-mm	10	5	10-30μl	50μl	2ml
60mm/T25 flask	21	10	25-75μl	125μl	5ml
100 mm	58	30	75-225μl	250μl	15ml

Taking one well in a 6-well plate as an example, explain the amount of each liquid component used:

Enhanced reagents	Serum-free diluent
10μl*	50μl

Mix

Concentrated virus	Serum-free diluent
10μl**	50μl

Complete culture medium per well
2000μl-60μl-60μl=1880μl

***Can be increased or decreased separately based on factors such as toxicity and efficiency**

****The amount used here is for illustration only, and the specific amount can be determined based on the regular amount at your discretion**

Common problems and solutions

Problem	Reason	Solution
The effect is not significant after using enhancers	Low virus concentration, insufficient dosage	Increase virus usage
	Excessive cell density	Reduce cell density
	The enhancement is not significant	Increase the dosage of enhancer by 1-3 times
Cytotoxicity	Excessive dosage of enhancer	Reduce the dosage of enhancers appropriately
	Excessive virus volume	Reduce virus usage appropriately
	Less cell culture media used	Increase the amount of culture medium or replace the culture medium
Sedimentation occurs in the mixed solution	Analysis of virus preservation solution	It can still be used, but pay attention to observing cell toxicity. The same serum-free dilution can also be added immediately to help precipitate and dissolve

Other related reagents

TR3012: Chronic virus infection enhancement reagent.

TR3003: Adenovirus infection enhancing reagent.

TR3041: Retroviral infection enhancing reagent.

TR3015: Virus infection enhancing reagent in animal bodies.