



Code GD1004-RV (GeneDetect rAVETM gene delivery reagent).

Vector AAV-SAR-CAG-eGFP-WPRE-BGH (Green fluorescent protein)

Vector description The CAG promoter consists of the chicken B-actin promoter hybridised

with the CMV immediate early enhancer sequence and is highly efficient in most tissue types. Addition of a scaffold attachment region (SAR) stabilises the WPRE regulatory element and the presence of a BGH polyadenylation sequence ensures transcription of the vector transgene

following transduction.

Lot Number 88765

Quantity $0.3\text{mL} (300\mu\text{l})$

Purity Affinity purified against immobilised heparan sulfate proteoglycan.

Titer/concentration 3.6 x 10¹⁰ genomic particles/ml

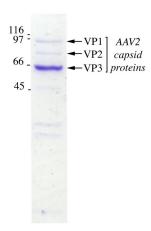
Presentation Liquid in phosphate buffered saline (PBS) containing 1mM MgSO₄

Storage and stability Upon receipt spin contents of vial to collect sample, aliquot as necessary

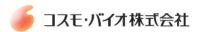
and store: 4°C for short term (<1 month), -20 °C or -80°C for long term.

Avoid repeated freeze-thaw cyles.

Quality control 10µl was analysed by SDS-PAGE to verify purity.



Note: GeneDetectTM and rAVETM are trademarks of GeneDetect.com Limited.



Handling

Always wear laboratory gloves, protective glasses and a suitable protective laboratory coat when using rAVE reagents. Recent NIH guidelines state that "adeno-associated virus (AAV) types 1 through 4, and recombinant AAV constructs, in which the transgene does not encode either a potentially tumorigenic gene product or a toxin molecule and are produced in the absence of a helper virus" can in most cases be handled at biosafety level 1 (BL1). You should follow the guidelines set by your Institutional biosafety committee for the handling of adeno-associated virus.

Disposal

rAVE reagents are susceptible to 5% phenol, 10% bleach, 10% Wescodyne or Virkon. We recommend using a fresh solution of 10% bleach for 30 minutes for decontamination.

Applications

For *in vitro* applications, mix 2µl rAVE sample with 200µl pre-warmed culture media and apply per well to cells of 60 - 80% confluency (24 well plate). Allow at least three days for viral integration and gene expression before analysis. For *in vivo* applications, dose should be determined by end user.

Refer to www.GeneDetect.com for a selection of protocols.

References

For a comprehensive list of references refer to www.GeneDetect.com