

Code MJFF-GD1007 (GeneDetect ® rAVE[™] Gene Delivery Reagent)

Vector scAAV2/5-CBh-Empty/Null-WPRE3-enSV40pA

Vector description scAAV2/5 Empty/Null Control Vector. The CBh promoter consists of a

hybrid form of the chicken β -actin promoter which drives robust expression. The Woodchuck post-transcriptional regulatory element (WPRE3) and the presence of a polyadenylation sequence (enSV40pA)

ensures high transcription following transduction.

Lot Number 56815

Quantity 0.2 ml

Purity Purified and concentrated by modified Iodixanol/cation exchange/Q-

Sepharose.

Titer/concentration 0.3 x 10¹³ genomic particles/ml (vg/ml)

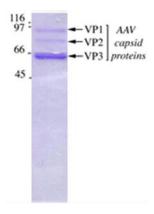
Presentation Liquid in phosphate buffered saline (PBS) containing 1mM MgCl₂

Storage & stability Upon receipt, briefly spin contents of vial to collect sample, aliquot on ice

under sterile conditions and store: 4°C for short term (<1 month), -20 °C

or -80°C for long term. Avoid repeated freeze-thaw cycles.

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



Note: GeneDetect $^{\otimes}$ 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.

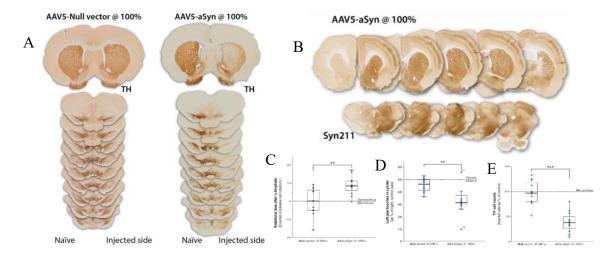
Disposal

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 $2\mu l$ rAVETM sample with $200\mu l$ pre-warmed culture media and apply per well to cells of 60 - 80% confluency (24well 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.

MJFF In Vivo Testing



Injection paradigm for wild type Sprague rats injected with 3 μ L doses confirmed by ddPCR and qPCR. Rats were sacrificed 6 weeks post injection. Representative images of TH **(A)** and aSyn **(B)** stained sections. 100% is the undiluted virus, ddPCR-matched at 2.4E12. **(C)** TH positive cell counts in the SNpc. **(D)** Left paw touches in cylinder. **(E)** Rotational bias after D- Amphetamine.

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