

Code MJFF-GD1006 (GeneDetect[®] rAVE[™] Gene Delivery Reagent)

Vector scAAV2/5-CBh-hu WT aSyn-WPRE3-enSV40pA

Vector description scAAV2/5 Expression 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 43411

Quantity 0.2 ml

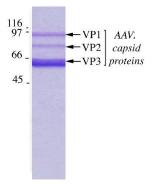
 Purity
 Purified and concentrated by modified Iodixanol/cation exchange/Q-Sepharose.

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

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

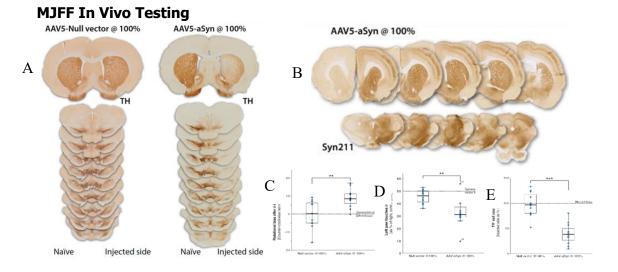
Storage & stability Upon receipt, briefly spin contents of vial to collect sample, aliquot on ice under sterile conditions and store: 4°C for short term (<1 month), -20 °C or -80°C for long term. <u>Avoid repeated freeze-thaw cycles</u>.

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



Note: $\mathsf{GeneDetect}^{\, \circledast}$ and $\mathsf{rAVE}^{\mathsf{TM}}$ are trademarks of $\mathsf{GeneDetect.com}$ 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
- 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 (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.



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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