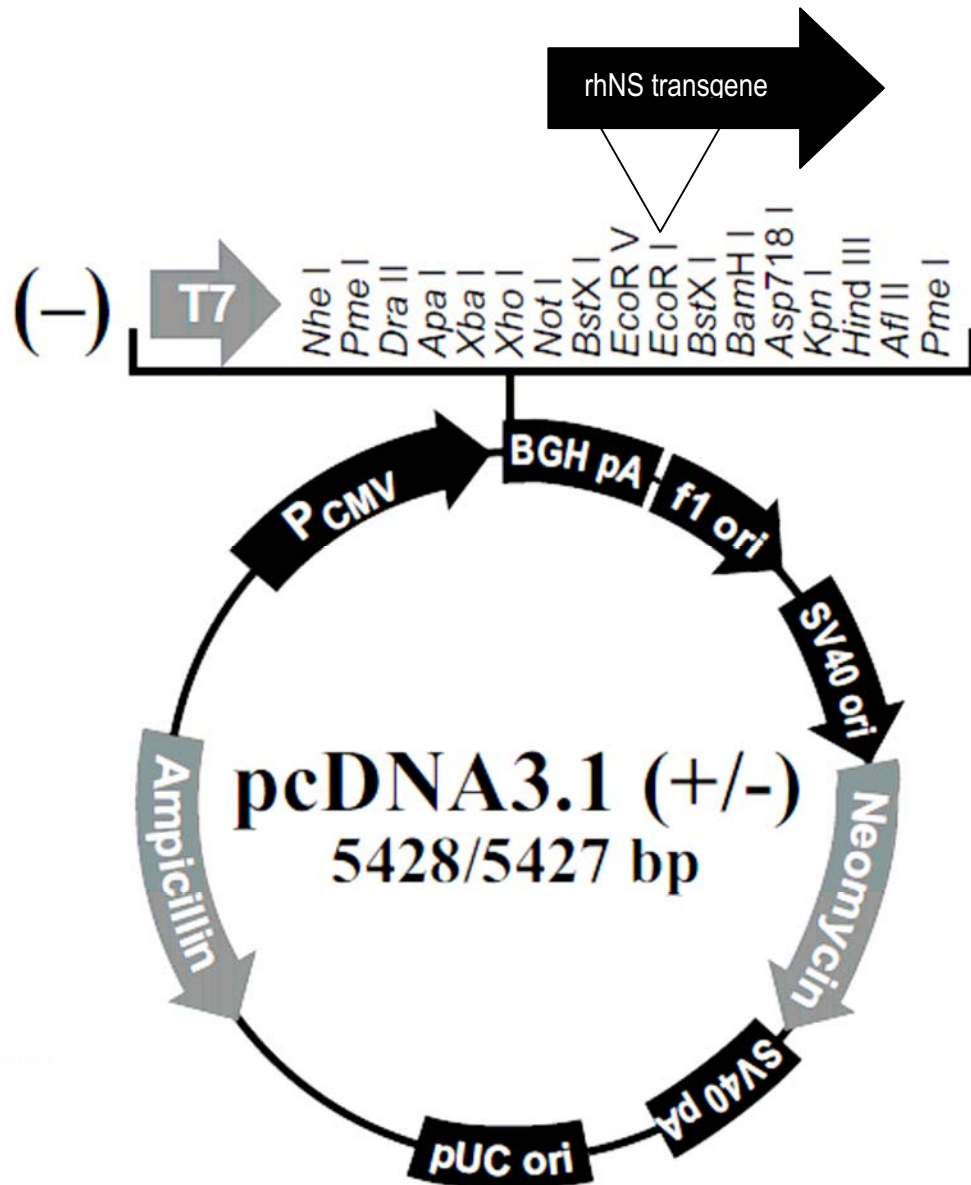


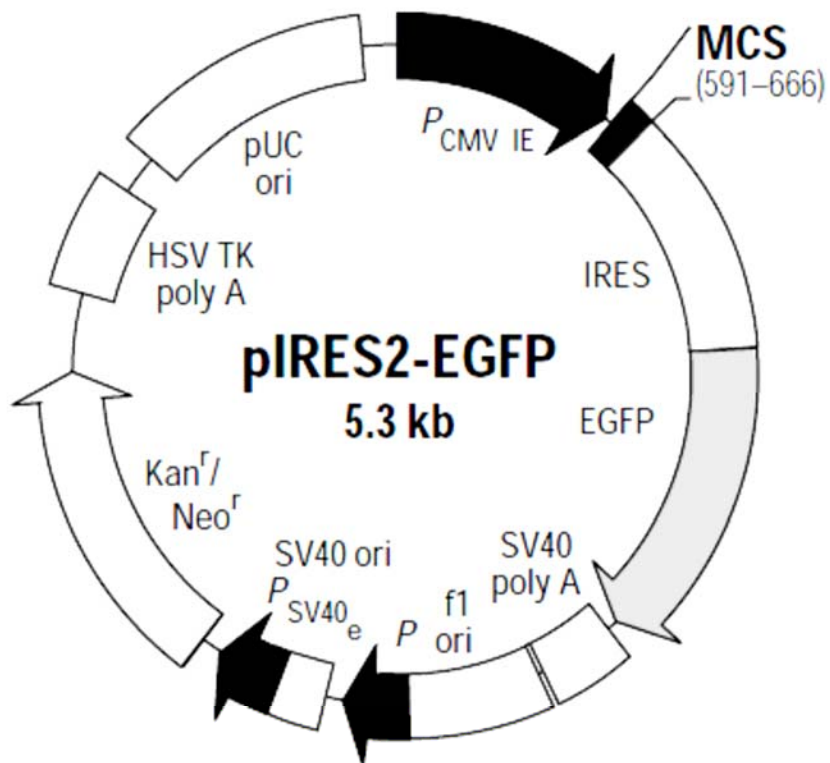
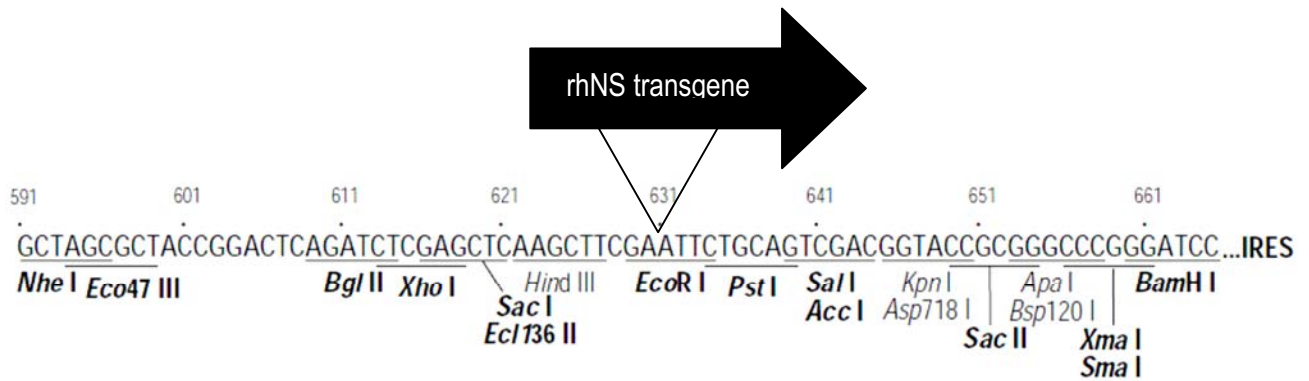
APPENDIX:
Plasmid DNA Maps



PLASMID NAME: rhNS-pcDNA3.1

PLASMID SIZE: 7233 bp

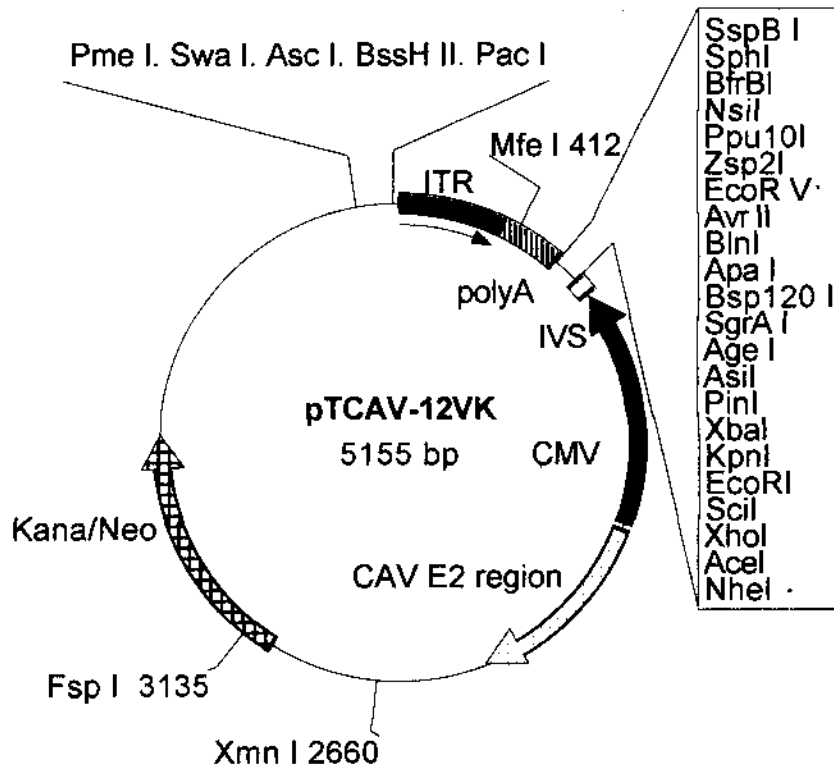
VECTOR DETAILS: The rhNS sequence was derived from human liver as described in Scott *et al* (1995). After the addition of *EcoRI* linkers, the rhNS transgene (bases 1 to 1806 of GenBank accession no. U30894) was cloned into the *EcoRI* site of the multiple cloning site of the pcDNA3.1 (-) vector (Invitrogen, cat. no. V95-20).



PLASMID NAME: pNS-IRES

PLASMID SIZE: 7114 bp

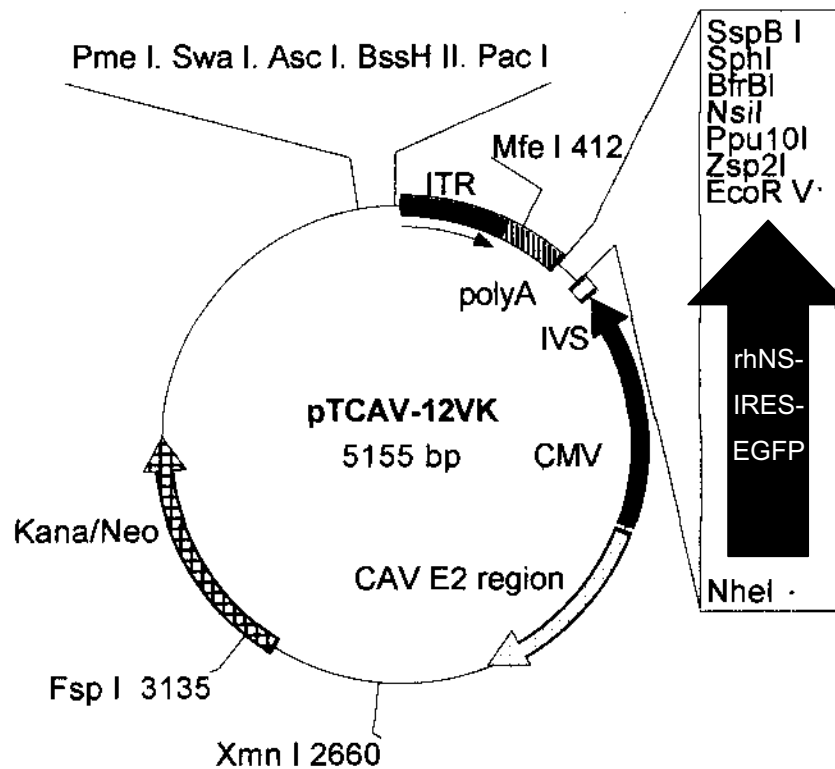
VECTOR DETAILS: The rhNS sequence (bases 1 to 1806 of GenBank accession no. U30894) was removed by *Eco*RI digestion from the rhNS-pcDNA3.1 vector and sub-cloned into the *Eco*RI site of the pIRES-2EGFP vector (Clontech, cat. no. 6029-1).



PLASMID NAME: pTCAV-12VK

PLASMID SIZE: 5155 bp

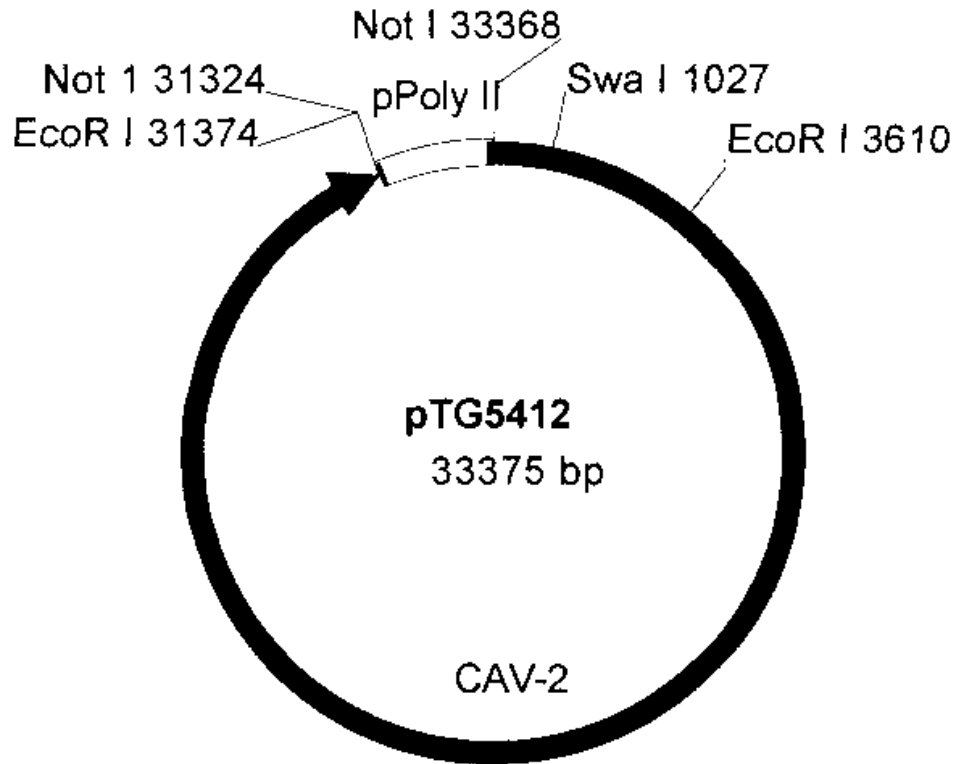
VECTOR DETAILS: This plasmid is derived from ptCAV-12a and has been modified by the insertion of the kanamycin/neomycin resistance gene in the *FspI* site of the ampicillin gene. The CAV-2 inverted terminal repeat (ITR) and the CAV-2 E2 region are homologous to the sequences in the pTG5412 plasmid. The desired transgene is subcloned into the multiple cloning site following an intervening sequence (IVS) and is driven by the CMV promoter.



PLASMID NAME: pTCAV-NS

PLASMID SIZE: 8275 bp

VECTOR DETAILS: The 5' rhNS-IRES-EGFP 3' sequence from the plasmid pNS-IRES was removed by digesting with *NotI*, blunting with Klenow fragment and then by digestion with *NheI* (i.e. sticky/blunt fragment). The pTCAV-12VK vector was digested with *NheI* and *EcoRV* (a blunt-cutter) and ligated with the rhNS-IRES-EGFP fragment to form the plasmid pTCAV-NS.



PLASMID NAME: pTG5412

PLASMID SIZE: 33375 bp

VECTOR DETAILS: This vector was constructed by TransGene SA (France) and contains the entire CAV-2 genome (Genbank accession no. J04368), including the inverted terminal repeats and E2B regions. The CAV-2 genome is cloned into a pPolyII backbone and is flanked by *NotI* linkers.

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NOTE: This image is included in the print copy of the thesis held in the University of Adelaide Library.

Addenda

Page 14, line 30: change to “benzodiazepines, chloral hydrate trimeprazine tartrate have had some success”.

Page 15, line 4: delete “antipsychotic”.

Page 60, line 15: delete “complete” and replace with “completely”.

Page 69, line 1: change “25 mL” to “25 μ L”.

Page 136, line 11: replace with “... was able to have...”

Page 137, lines 1-4: change to “Injection of 10^8 or 10^9 particles/ μ L CAV-NS at birth did not significantly impact on the concentration of HNS-UA detected in the olfactory bulb at both 1-wk and 6-wks post-injection, given the variation in the PBS-treated MPS IIIA group at these time-points ($p=0.80$, $p=0.30$).

Page 145, line 30: delete final sentence and replace with “The operator was placed at a point equidistant from an open and closed arm”.

Page 153, line 17: change “marginally significant” to “approached significance”.

Page 160, paragraph 2: after “1- 7-days after the initial exposure” insert the sentence “This discrepancy may relate to the time difference between the first trial and subsequent re-exposure to the EPM apparatus in these studies (less than 1 week) compared to the 3 weeks interval utilised in our study”.

Page 162, line 5: delete “Neither antibodies towards rhNS nor”.

Page 179, line 5: insert after the end of sentence “CAV-NS treatment did not increase the amount of NS activity measured in unaffected or MPS IIIA mice”.

Page 203, line 18: change “Compared to PBS-treated animals, CAV-NS treatment had no effect on the functional changes associated with MPS IIIA.” to read: “No functional changes were able to be demonstrated between PBS- and CAV-NS-treated MPS IIIA animals.”