

[Mouse Models](#)[MSA/OPCA](#)[Publication](#)

TgM83+/? Snca+/?

IDENTIFICATION

| | |
|-------------------------|--|
| Causal gene(s) | Sporadic |
| Repeat size or mutation | human A53T ?-synuclein (heterozygous) and mouse ?-synuclein (homozygous) |
| Animal model | Mouse |

MODEL DETAILS

| | |
|---------------------------|-------------------------|
| Mouse strain / background | 129/SvEvTac x C57Bl/C3H |
| Type of model | Transgenic |

TRANSGENIC CONSTRUCT

| | |
|-------------------------------------|---------------------------------------|
| Transgenic construct: sequence type | cDNA for human ?-synuclein expression |
| Promoter: gene | PrP |
| Promoter: species | Mouse |

PHENOTYPE

| | |
|-------------------|---|
| Hallmark features | No development of spontaneous illness, neurodegeneration only occurred after addition of MSA brain extracts |
|-------------------|---|

SOURCE & PUBLICATIONS

| | |
|---|---|
| Originating lab / institution | University of California |
| Links to publications or public resources | Evidence for u03b1-synuclein prions causing multiple system atrophy in humans with parkinsonism - PubMed https://pubmed.ncbi.nlm.nih.gov/26324905/ |